

Clinical Trials in Myeloma and Related Disorders at PMH
(Version: May 2018)

NEWLY DIAGNOSED MULTIPLE MYELOMA

Phase 3, Randomized, Placebo-Controlled, Double-Blind Study of Oral Ixazomib Maintenance Therapy After Initial Therapy in Patients With Newly Diagnosed Multiple Myeloma Not Treated With Stem Cell Transplantation (Protocol Number: C16021)

Inclusion Criteria

Each patient must meet all the following inclusion criteria to be randomized to treatment:

1. Adult male or female patients aged 18 years or older with a confirmed diagnosis of symptomatic NDMM according to standard criteria.
 2. Completed 6 to 12 months (+/- 2 weeks) of initial therapy, during which the patient was treated to best response, defined as the best response maintained for 2 cycles after the M-protein nadir is reached.
 3. Documented major response (PR, VGPR, and CR) according to the IMWG uniform response criteria, version 2011, after this initial therapy.
 4. Female patients who:
 - Are postmenopausal for at least 1 year before the screening visit, OR
 - Are surgically sterile, OR
 - If they are of childbearing potential, agree to practice 2 effective methods of contraception, at the same time, from the time of signing the informed consent through 90 days after the last dose of study drug, or
 - Agree to practice true abstinence, when this is in line with the preferred and usual lifestyle of the subject. (Periodic abstinence (eg, calendar, ovulation, symptothermal, post ovulation methods] and withdrawal are not acceptable methods of contraception.)
- Male patients, even if surgically sterilized (ie, status postvasectomy), who:
- Agree to practice effective barrier contraception during the entire study
 - Treatment period and through 90 days after the last dose of study drug, or agree to practice true abstinence, when this is in line with the preferred and usual lifestyle of the subject. (Periodic abstinence [eg, calendar, ovulation, symptothermal, post ovulation methods for the female partner] and withdrawal are not acceptable methods of contraception.)
5. Voluntary written consent must be given before performance of any study-related procedure not part of standard medical care, with the understanding that consent may be withdrawn by the patient at any time without prejudice to future medical care.
 6. Complete documentation of the details of the initial therapy before randomization including cytogenetics and ISS is available.
 7. Eastern Cooperative Oncology Group Performance Status of 0 to 2.

8. Suitable venous access for the study-required blood sampling and consent for the specific amounts that will be taken.
9. Patient is willing and able to adhere to the study visit schedule and other protocol requirements including blood sampling and bone marrow aspiration.
10. Patients must meet the following clinical laboratory criteria at study entry:
 - Absolute neutrophil count (ANC) $\geq 1,000/\text{mm}^3$ without growth factor support and platelet count $\geq 75,000/\text{mm}^3$. Platelet transfusions to help patients meet eligibility criteria are not allowed within 3 days before randomization.
 - Total bilirubin $\leq 1.5 \times$ the upper limit of the normal range (ULN).
 - Alanine aminotransferase and aspartate amino transferase $\leq 3 \times$ ULN.
 - Calculated creatinine clearance $\geq 30 \text{ mL/min}$ (using the Cockcroft-Gault equation)

Exclusion Criteria

Patients meeting any of the following exclusion criteria are not to be randomized to treatment:

1. Multiple myeloma that has relapsed after, or was not responsive to, initial therapy.
2. Prior SCT.
3. Radiotherapy within 14 days before randomization.
4. Diagnosed or treated for another malignancy within 5 years before randomization or previous diagnosis with another malignancy. Patients with nonmelanoma skin cancer or carcinoma in situ of any type are not excluded if they have undergone complete resection.
5. Female patients who are lactating and breastfeeding or have a positive serum pregnancy test during the Screening period.
6. Major surgery within 14 days before randomization.
7. Central nervous system involvement.
8. Infection requiring IV antibiotic therapy or other serious infection within 14 days before randomization.
9. Diagnosis of Waldenstrom's Macroglobulinemia, POEMS (polyneuropathy, organomegaly, endocrinopathy, monoclonal gammopathy, and skin changes) syndrome, plasma cell leukemia, primary amyloidosis, myelodysplastic syndrome, or myeloproliferative syndrome.
10. Evidence of current uncontrolled cardiovascular conditions, including uncontrolled hypertension, uncontrolled cardiac arrhythmias, uncontrolled congestive heart failure, unstable angina, or myocardial infarction within the past 6 months.
11. Systemic treatment with strong inhibitors of CYP1A2 (fluvoxamine, enoxacin, ciprofloxacin), strong inhibitors of CYP3A (clarithromycin, telithromycin, itraconazole, Voriconazole, ketoconazole, nefazodone, posaconazole), or strong CYP3A inducers (rifampin, rifapentine, rifabutin, carbamazepine, phenytoin, phenobarbital) or use of Ginkgo biloba or St. John's wort within 14 days before randomization.
12. Ongoing or active infection, known human immunodeficiency virus positive, active hepatitis B or C infection.

Clinical-grade Molecular Profiling of Patients with Multiple Myeloma and Related Plasma Cell Malignancies (Protocol MMRF-002; Non-Interventional)

Inclusion Criteria:

1. Patients must have a diagnosis of multiple myeloma or related plasma cells malignancies
2. Patients are undergoing standard of care bone marrow aspirates.
3. Patients (male or female) from any race or ethnicity must be 18 years of age at the time of registration
4. Procedure-specific signed informed consent form (ICF) prior to initiation of any study-related procedures.

Exclusion Criteria:

1. It is the enrolling study physician's discretion to decide if a patient is not fit enough to undergo tissue biopsy.
2. Patients who are incarcerated are not eligible to participate.
3. Women who are pregnant.
4. Patients who have had another malignancy within the last five (5) years (except for basal or squamous cell carcinoma, or in situ cancer of the cervix)

Health-related quality of life and caregiver burden assessment in multiple myeloma and lymphoma patients and their caregivers undergoing outpatient autologous stem cell transplantation as compared to inpatient transplantations: a needs assessment (Non-Interventional)

Inclusion criteria:

1. Males or females aged 18 years or older undergoing an autologous stem cell transplant for multiple myeloma, or Hodgkin or Non-Hodgkin Lymphoma
2. Able to provide consent
3. Able to read, write and speak English
4. Available primary caregiver for the caregiver QOL and burden component of study who is able to provide consent and read, write and speak English

Exclusion criteria:

1. Geographically inaccessible/will not be followed at Princess Margaret Cancer Centre for the 100d period post-transplant.
2. Unable to provide consent.

The Terry Fox Pan-Canadian Multiple Myeloma Molecular Monitoring Cohort Study (The M4 Study; Non-Interventional)

Inclusion criteria:

1. Age \geq 19 years
2. Ability to give informed consent
3. Diagnosed with active multiple myeloma (refer to Appendix I for IMWG definition)
4. Also enrolling in the CMM-DB project; and
5. Previously untreated and eligible for autologous stem-cell transplantation (ASCT)
6. Patients who are going to be treated on a clinical trial are also eligible to participate in this study if they meet the other eligibility criteria.

Detection of AL amyloid fibrils and oligomers in blood plasma of Multiple Myeloma and Related Plasma Cell Dyscrasias using Immuno-Gold Electron Microscopy (Non-Interventional)

Inclusion criteria:

1. Patients must have or be suspected of a diagnosis of AL amyloidosis, MM, or related clonal plasma cell disorder (PCD) such as smoldering myeloma or MGUS.
2. Patient must be \geq 18 years old.
3. Patients are undergoing standard of care blood draw.
4. All patients must have signed and dated an informed consent form.

Healthy Subject Inclusion Criteria

1. 18-60 years old
2. 110 lbs. and above
3. Not pregnant
4. Not known to be anemic

Phase 1/2 Multicenter, Open-Label Study to Determine the Recommended Dose and Regimen of Durvalumab (Medi4736) In Combination with Lenalidomide (Len) With and Without Dexamethasone (Dex) In Subjects with Newly Diagnosed Multiple Myeloma (Protocol: MEDI4736-MM-002)

Inclusion criteria

1. Subjects must satisfy the following criteria to be enrolled into the study:
 - Subject is ≥ 18 years of age at the time of signing the informed consent form (ICF)
 - Subject must understand and voluntarily sign an ICF prior to any study-related assessments/procedures being conducted
 - Subject is willing and able to adhere to the study visit schedule and other protocol requirements
 - Subject must have documented diagnosis with previously untreated (for cohort C, the induction and consolidation treatment along with the first ASCT are allowed), symptomatic MM as defined by the criteria below (Rajkumar, 2014; NCCN-MM, 2015):
2. MM diagnostic criteria (all 3 required);
 - Monoclonal protein present in the serum and/or urine
 - Clonal bone marrow plasma cells $\geq 10\%$ or biopsy-proven bony or extramedullary plasmacytoma*
 - Any one or more of the following myeloma defining events:
 - i. One or more of the following Myeloma-related organ dysfunction (at least one)
 - [C] Calcium elevation (serum calcium >11.5 mg/dl) [> 2.65 mmol/L]
 - [R] Renal insufficiency (serum creatinine >2 mg/dl) [$177 \mu\text{mol/L}$ or more] or creatinine clearance < 40 ml/min
 - [A] Anemia (hemoglobin <10 g/dl or >2 g/dL below the lower limit of laboratory normal)
 - [B] Bone lesions (lytic or osteopenic) one or more bone lesions on skeletal radiography, CT, or PET-CT
 - ii. One or more of the following biomarkers of malignancy:
 - Clonal bone marrow plasma cell percentage* $\geq 60\%$
 - Abnormal serum free light-chain ratio ≥ 100 (involved kappa) or < 0.01 (involved lambda)
 - >1 focal lesions detected by functional imaging including PET/CT and/or whole body magnetic resonance imaging (MRI)
- AND have measurable disease by protein electrophoresis analyses as defined by the following:**
 - IgG MM: Serum monoclonal paraprotein (M-protein) level ≥ 1.0 g/dl or urine M protein level ≥ 200 mg/24 hours
 - IgA MM: Serum M-protein level ≥ 0.5 g/dl or urine M-protein level ≥ 200 mg/24 hours
 - IgM MM (IgM M-protein plus lytic bone disease documented by skeletal survey plain films): Serum M-protein level ≥ 1.0 g/dl or urine M-protein level ≥ 200 mg/24 hours
 - IgD MM: Serum M-protein level ≥ 0.05 g/dl or urine M-protein level ≥ 200 mg/24 hours
 - Light chain MM: Serum M-protein level ≥ 1.0 g/dl or urine M-protein level ≥ 200 mg/24 hours
3. Eastern Cooperative Oncology Group (ECOG) performance status of 0, 1, or 2
4. Females of childbearing potential (FCBP1) must:
 - Have two negative pregnancy tests as verified by the investigator prior to starting study treatment. She must agree to ongoing pregnancy testing during the course of the study, and after end of study treatment. This applies even if the subject practices true abstinence² from heterosexual contact.

- She must either commit to true abstinence from heterosexual contact (which must be reviewed on a monthly basis and be source documented) or agree to use, and be able to comply with, effective contraception without interruption, 28 days prior to starting study treatment, during the study therapy (including dose interruptions), and for 90 days after discontinuation of study treatment.
 - Refrain from egg cell and blood donation for 90 days after the final dose of Durvalumab.
5. Male subjects must:
 - Practice true abstinence (which must be reviewed on a monthly basis) or agree to use a condom during sexual contact with a pregnant female or a FCBP while participating in the study, during dose interruptions and for at least 90 days following study treatment discontinuation, even if he has undergone a successful vasectomy.
 - Refrain from sperm and blood donation for at least 90 days after the final dose of Durvalumab
 6. For Cohort A subject must be transplant non-eligible (TNE) and meet at least one of the following high risk factors:
 - Cytogenetic abnormalities finding in malignant myeloma clone with t(4; 14); and / or del(17p); and / or 1q rearrangement; and / or t(14;16); or
 - ISS Stage III; or
 - Serum LDH > 2 x ULN;
 7. For Cohort B subject must be ≥ 65 years of age at the time of signing the informed consent form (ICF) and transplant non-eligible (TNE); excluding the subjects who meet the Cohort A criteria
 8. For Cohort C subject must be after first autologous stem cell transplantation (ASCT) for NDMM and meet the following criteria:
 - Have a post-transplant response as PR or better at the time of enrollment to this study;
 - Have one of the following high risk factors at the time of NDMM diagnosis;
 - i. Cytogenetic abnormalities finding in malignant myeloma clone with t(4; 14); and / or del(17p); and / or 1q rearrangement; and / or t(14; 16); or
 - ii. ISS stage III; or
 - iii. Serum LDH > 2 x ULN;
 9. MRD positive (defined as more than 1 malignant cell in 105 cells) measured by ClonoSIGHT™ NGS assay of a BMA sample) at the time of enrollment to this study; BMA sample collected at the time of multiple myeloma diagnosis, prior to induction therapy available for central MRD assessment by ClonoSIGHT™ NGS assay

Exclusion criteria

The presence of any of the following will exclude a subject from enrollment:

1. Previous treatment with anti-myeloma therapy (does not include radiotherapy, bisphosphonates, or a single short course of steroid [ie, less than or equal to the equivalent of dexamethasone 40 mg/day for 4 days; such a short course of steroid treatment must not have been given within 14 days of Cycle 1 Day 1], for Cohort C, the induction and consolidation treatment along with the first ASCT are allowed)
2. Any of the following laboratory abnormalities:
 - Absolute neutrophil count (ANC) < 1,000/μL
 - Untransfused platelet count < 75,000 cells/μL
 - Serum aspartate aminotransferase/serum glutamic oxaloacetic transaminase (SGOT/AST) or alanine aminotransferase (SGPT/ALT) > 2.5 × upper limit of normal (ULN)
 - Serum total bilirubin > 1.5 × ULN or > 3.0 mg/dL for subjects with documented Gilbert's syndrome
 - Corrected serum calcium >13.5 mg/dL (> 3.4 mmol/L)
3. Renal failure requiring hemodialysis or peritoneal dialysis

4. Any serious medical condition that places the subject at an unacceptable risk if he or she participates in this study. Examples of such a medical condition are, but are not limited to, subject with unstable cardiac disease as defined by: cardiac events such as myocardial infarction (MI) within the past 6 months, NYHA (New York Heart Association) heart failure class III-IV, uncontrolled atrial fibrillation or hypertension; subjects with conditions requiring chronic steroid or immunosuppressive treatment, such as rheumatoid arthritis, multiple sclerosis and lupus, that likely need additional steroid or immunosuppressive treatments in addition to the study treatment
5. Peripheral neuropathy \geq Grade 2
6. Primary AL (immunoglobulin light-chain) amyloidosis and myeloma complicated by amyloidosis
7. Prior history of malignancies, other than MM, unless the subject has been free of the disease for \geq 5 years with the exception of the following non-invasive malignancies:
 - Basal cell carcinoma of the skin
 - Squamous cell carcinoma of the skin
 - Carcinoma in situ of the cervix
 - Carcinoma in situ of the breast
 - Incidental histologic finding of prostate cancer (T1a or T1b using the TNM [tumor, nodes, metastasis] clinical staging system) or prostate cancer that is curative
8. Subject is positive for human immunodeficiency virus (HIV); chronic or active hepatitis B or active hepatitis A, or C
9. Subject had prior exposure to immunotherapy, including, but not limited to, other anti-CTLA-4, anti-PD-1, anti-PD-L1 monoclonal antibody or inhibitor, cell-based therapies, or cancer vaccines
10. Subject has history of organ or allogeneic stem cell transplantation
11. Subject who has had clinical evidence of central nervous system (CNS) or pulmonary leukostasis, disseminated intravascular coagulation, or CNS multiple myeloma, or plasma cell leukemia
12. Known or suspected hypersensitivity to the excipients contained in the formulation of
13. Durvalumab, lenalidomide, or dexamethasone
14. Major surgery (as defined by the investigator) within the 28 days prior to the first dose of study treatment
15. Received prior treatment (for any reason) with a monoclonal antibody within 5 half-lives of initiating study treatment
16. Use of any investigational agents within 28 days or 5 half-lives (whichever is longer) of initiating study treatment.

RELAPSED OR REFRACTORY MULTIPLE MYELOMA

A Phase 2, Multicenter Study to Determine the Efficacy and Safety of bb2121 in Subjects with Relapsed and Refractory Multiple Myeloma (Protocol Number: BB2121-MM-001)

Inclusion Criteria

1. Documented diagnosis of multiple myeloma
2. Must have received at least 3 prior MM treatment regimens.
3. Must have undergone at least 2 consecutive cycles of treatment for each regimen, unless PD was the best response to the regimen.
4. Must have received a proteasome inhibitor, an immunomodulatory agent and an anti-CD38 antibody.
5. Must be refractory to the last treatment regimen. Refractory is defined as documented progressive disease during or within 60 days (measured from the last dose) of completing treatment with the last anti-myeloma drug regimen before study entry.
6. Subjects must have measurable disease, including at least one of the criteria below:
 - Serum M-protein greater or equal to 1.0 g/dL
 - Urine M-protein greater or equal to 200 mg/24 h
 - Serum free light chain (FLC) assay: involved FLC level greater or equal to 10 mg/dL (100 mg/L) provided serum FLC ratio is abnormal (Clarification by Medical Monitor: Measurable FLC would count for eligibility. However, if patient has disease that is followed by M-protein in serum for response, they need to progress by serum M protein. They cannot progress by FLC alone if the disease has always been followed by serum m protein)

Exclusion Criteria:

1. Subjects with solitary plasmacytomas or non-secretory myeloma without other evidence of measurable disease
2. Inadequate bone marrow function defined by:
 - ANC < 1000 cells/mm³ in the absence of growth factor support (Filgrastim within 7 days or pegfilgrastim within 14 days of screening)
 - Platelet count < 50,000 mm³ in the absence of transfusion support (platelet transfusion within 7 days of screening).
3. Inadequate renal function defined by CrCl ≤ 45 ml/min using Cockcroft-Gault equation.
4. International ratio (INR) or partial thromboplastin time (PTT) > 1.5 × ULN,
5. Echocardiogram or MUGA with LVEF < 45%.
6. Inadequate pulmonary function (oxygen saturation (SaO₂) < 92 % on room air).
7. Ongoing treatment with chronic immunosuppressants (e.g., cyclosporine or systemic steroids at any dose). Intermittent topical, inhaled or intranasal corticosteroids are allowed.
8. Following transmissible disease tests will be done as part of the screening for this study:
 - HIV 1 and 2 antibody/p24 antigen combination, Hepatitis B Surface Antibody (HBsAb), Hepatitis B Surface Antigen (HBsAg), Hepatitis B Core Antibody (HBcAb), Hepatitis C antibody (HCV antibody), syphilis (by CMIA), HTLV 1 & 2 antibody, HBV DNA and HCV RNA testing. West Nile Virus; CMV, EBV (Dr. Reece's agreement to Bio-Safety Committee)
9. Patients with positive tests for following diseases will be excluded from this study (as per protocol): HIV; hepatitis B virus (HBV); hepatitis C virus.
10. Patient will be verbally screened for the history of travelling in the Zika Virus affected area. If an assay for antibody titer is positive for Zika virus. Patient will be excluded from the study (Dr. Reece's agreement to Bio-Safety Committee)

11. Dr. Reece has agreed to follow the BMT program protocol for patient's eligibility in addition to protocol (patients with West Nile virus and Syphilis will be excluded).
12. Subject requires ongoing treatment with chronic, therapeutic dosing of anti-coagulants (eg, warfarin, low molecular weight heparin, or Factor Xa inhibitors).
13. Washout-page 53:
 - Any prior systemic therapy for MM within 7 days prior to scheduled protocol required leukapheresis.
 - Experimental agents within 4 weeks prior to leukapheresis unless no response or PD is documented on the experimental therapy (Clarification: If patient's best response to an Investigational Agent (e.g. anti-PDL1) is progressive disease, then 7 days. If not, then 28 days for this experimental agent.
 - Therapeutic doses of corticosteroids (defined as > 20 mg/day prednisone or equivalent) within 7 days prior to leukapheresis.

AN OPEN-LABEL, MULTICENTER, PHASE I TRIAL EVALUATING THE SAFETY AND PHARMACOKINETICS OF ESCALATING DOSES OF BFCR4350A IN PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA (Protocol Number: GO39775)

Key Inclusion Criteria

1. Patients must have R/R MM for which no established therapy for MM is appropriate and available or be intolerant to those established therapies
2. Agreement to provide bone marrow biopsy and aspirate samples as per protocol
3. Adverse events from prior anti-cancer therapy resolved to Grade \leq 1, with the following exceptions:
 - Any grade alopecia, peripheral sensory or motor neuropathy must have resolved to Grade \leq 2
 - Measurable disease defined as at least one of the following:
 - Serum monoclonal protein (M-protein) \geq 0.5 g/dL (\geq 5 g/L)
 - Urine M-protein \geq 200 mg/24 hr.
 - Serum free light chain (SFLC) assay: Involved SFLCs \geq 10 mg/dL (\geq 100 mg/L) and an abnormal SFLC ratio ($<$ 0.26 or $>$ 1.65)
4. Laboratory values:
 - Hepatic function: AST and ALT \leq 3 \times ULN; Total bilirubin \leq 1.5 \times ULN; patients with a documented history of Gilbert syndrome and in whom total bilirubin elevations are accompanied by elevated indirect bilirubin are eligible.
 - Hematologic function: Platelet count \geq 75,000/mm³ without transfusion within 14 days prior to first dose of BFCR4350A, ANC \geq 1000/mm³, Total hemoglobin \geq 8 g/dL
 - Creatinine \leq 2.0 mL/dL and creatinine clearance (CrCl) \geq 30 mL/min (either calculated or per 24-hr urine collection)
 - Serum calcium (corrected for albumin) level at or below the ULN
5. For women of childbearing potential: agreement to remain abstinent (refrain from heterosexual intercourse) or use contraceptive methods with a failure rate of $<$ 1% per year during the treatment period and for at least 3 months after the last dose of BFCR4350A and tocilizumab (if applicable)
6. For men: agreement to remain abstinent (refrain from heterosexual intercourse) or use a condom, and agreement to refrain from donating sperm

Key Exclusion Criteria

1. Prior use of any monoclonal antibody, radioimmunoconjugate, or antibody-drug conjugate within 4 weeks before first BFCR4350A infusion
2. Prior treatment with systemic immunotherapeutic agents, including, but not limited to, cytokine therapy and anti-CTLA4, anti-PD-1, and anti-PD-L1 therapeutic antibodies, within 12 weeks or 5 half-lives of the drug, whichever is shorter, before first BFCR4350A infusion
3. Treatment-related, immune-mediated adverse events associated with prior immunotherapeutic agents as follows:
 - Grade \geq 3 adverse events with the exception of Grade 3 endocrinopathy managed with replacement therapy

- Grade 1–2 adverse events that did not resolve to baseline after treatment discontinuation
4. Treatment with radiotherapy, any chemotherapeutic agent, or treatment with any other anti-cancer agent (investigational or otherwise) within 4 weeks or 5 half-lives of the drug, whichever is shorter, prior to first BFCR4350A infusion
 5. Autologous stem cell transplantation (SCT) within 100 days prior to first BFCR4350A infusion
 6. Prior allogeneic SCT
 7. Primary or secondary plasma cell leukemia as defined by an absolute plasma cell count exceeding 2000/ μ L or 20% of the peripheral blood white cells
 8. Prior solid organ transplantation
 9. History of autoimmune disease, including, but not limited to, myasthenia gravis, myositis, autoimmune hepatitis, systemic lupus erythematosus, rheumatoid arthritis, inflammatory bowel disease, vascular thrombosis associated with antiphospholipid syndrome, Wegener’s granulomatosis, Sjögren’s syndrome, Guillain-Barré syndrome, multiple sclerosis, vasculitis, or glomerulonephritis. Patients with a history of autoimmune-related hypothyroidism on a stable dose of thyroid replacement hormone may be eligible for this study.
 10. Patients with history of confirmed progressive multifocal leukoencephalopathy
 11. History of severe allergic or anaphylactic reactions to monoclonal antibody therapy (or recombinant antibody-related fusion proteins)
 12. History of other malignancy that could affect compliance with the protocol or interpretation of results. - Patients with a history of curatively treated basal or squamous cell carcinoma of the skin or in situ carcinoma of the cervix are allowed.
 - Patients with a malignancy that has been treated with curative intent will also be allowed if the malignancy has been in remission without treatment for \geq 2 years prior to first BFCR4350A infusion.
 13. Current or past history of CNS disease, such as stroke, epilepsy, CNS vasculitis, neurodegenerative disease, or CNS involvement by MM
 - Patients with a history of stroke who have not experienced a stroke or transient ischemic attack in the past 2 years and have no residual neurologic deficits as judged by the investigator are allowed.
 - Patients with a history of epilepsy who have had no seizures in the past 2 years while not receiving any anti-epileptic medications are allowed.
 14. Significant cardiovascular disease (such as New York Heart Association Class III or IV cardiac disease, myocardial infarction within the last 6 months, unstable arrhythmias, or unstable angina)
 15. Significant active pulmonary disease (e.g., bronchospasm and/or obstructive pulmonary disease)
 16. Known active bacterial, viral, fungal, mycobacterial, parasitic, or other infection (excluding fungal infections of nail beds) at study enrollment, or any major episode of infection requiring treatment with IV antibiotics within 4 weeks prior to first BFCR4350A infusion
 17. Known or suspected chronic active EBV infection.
 18. Recent major surgery within 4 weeks prior to first BFCR4350A infusion
 19. Positive serologic or PCR test results for acute or chronic HBV infection: Patients whose HBV infection status cannot be determined by serologic test results
 20. Acute or chronic HCV infection
 21. Known history of HIV seropositivity
 22. Administration of a live, attenuated vaccine within 4 weeks before first BFCR4350A infusion or anticipation that such a live attenuated vaccine will be required during the study.
 23. Received systemic immunosuppressive medications (including, but not limited to, cyclophosphamide, azathioprine, methotrexate, thalidomide, and anti-tumor necrosis factor agents) with the exception of corticosteroid treatment \leq 10 mg/day prednisone or equivalent within 2 weeks prior to first dose of BFCR4350A

- Patients who received acute, low-dose, systemic immunosuppressant medications (e.g., single dose of dexamethasone for nausea) may be enrolled in the study after discussion with and approval of the Medical Monitor
 - The use of inhaled corticosteroids, mineralocorticoids for management of orthostatic hypotension, physiologic doses of corticosteroids for management of adrenal insufficiency is permitted.
24. History of illicit drug or alcohol abuse within 12 months prior to screening,

MCRN 004 – The LAURENTIANS Trial: A Randomized Phase II, Open Label, Study of Daratumumab, Weekly Low-Dose Oral Dexamethasone and Cyclophosphamide with or Without Pomalidomide in Patients with Relapsed and Refractory Multiple Myeloma

Inclusion Criteria

1. Males or females, age 18 years or older.
2. ECOG performance status score of 0, 1 or 2.
3. Life expectancy of at least 3 months
4. Measurable disease according to the IMWG criteria defined as
 - Serum monoclonal paraprotein (M-protein) ≥ 10 g/L (if IgG) or ≥ 5 g/L (if IgA, D, E or M).
 - Urine M-protein ≥ 200 mg/24 h.
 - Serum free light chains (FLC) assay: Involved FLC level ≥ 100 mg/L and an abnormal serum free light chain ratio (< 0.26 or > 1.65) if disease otherwise unmeasurable by a and b.
5. Relapsed or relapsed and refractory disease defined as documented disease progression during or after completing their last treatment line and it must have contained either bortezomib and/or lenalidomide. The only exception for non-refractory patients is when re-treatment with these agents is medically contra-indicated.
6. Have undergone at least 1 prior line of therapy. Induction therapy followed by ASCT and consolidation/maintenance will be considered as one line.
7. Have achieved at least a Minimal Response (MR) or better to at least one previous line of therapy.
8. Have received at least 2 consecutive cycles of prior treatment that have included lenalidomide or bortezomib, either alone or in combination regimens, unless intolerant to these agents.
9. Subjects must be eligible for pomalidomide reimbursement by their provincial jurisdictions or by the criteria of their private insurance companies.
10. Have the following laboratory values:
 - ANC $\geq 1.0 \times 10^9$ /L
 - Hemoglobin ≥ 80 g/L
 - Platelets $\geq 70 \times 10^9$ /L (or $\geq 50 \times 10^9$ /L if $\geq 50\%$ plasmacytosis in bone marrow).
 - Calculated CrCl ≥ 45 mL/min
 - AST and ALT $\leq 3.0 \times$ ULN
 - Total bilirubin $\leq 2 \times$ ULN unless known to have Gilbert's disease
 - Corrected serum calcium ≤ 3.5 mmol/L
11. Have signed the informed consent documents indicating that the subject understands the purpose of the procedures required for the study and is willing to participate and adhere to the study protocol.
12. Females with child-bearing potential (FCBP⁺) must agree to use 2 reliable forms of contraception* simultaneously or practice complete abstinence from heterosexual contact for at least 28 days before starting study drug, while participating in the study (including during dose interruptions), and for at least 90 days after study treatment discontinuation.
13. Females must agree to abstain from breastfeeding during study participation and 90 days after study drug discontinuation.

14. Males must agree to use a latex condom during any sexual contact with FCBP while participating in the study and for 90 days following discontinuation from this study, even if he has undergone a successful vasectomy.
15. Males must also agree to refrain from donating semen or sperm during the treatment phase and for 90 days after discontinuation from this study treatment.
16. All subjects must agree to refrain from donating blood while on study therapy and for 28 days after discontinuation from this study treatment.
17. All subjects must be eligible for enrollment and enrolled in the RevAid® program (refer to <https://www.revaid.ca/revaid/> for RevAid® eligibility).

Exclusion Criteria

1. Prior exposure to daratumumab (or other anti-CD38 monoclonal antibody) or pomalidomide
2. History of prior allogeneic stem cell transplantation and showing evidence of active graft-versus-host disease or graft-versus-host disease that requires immunosuppressive therapy.
3. Chemotherapy or other anti-myeloma therapy within 14 days prior to the first dose of study drug.
4. Treatment-related toxicity that has not recovered \leq Grade 1 unless deemed to be irreversible (an example of an irreversible toxicity would include steroid induced cataracts). Peripheral neuropathy $>$ Grade 2 or Grade 2 with pain will be excluded.
5. Subjects who have received steroids within 2 weeks prior to starting study treatment or who have not recovered from side effects of such therapy. Concomitant therapy medications that include corticosteroids are allowed if subject receive \leq 10 mg of prednisone per day, or equivalent, as indicated for other medical conditions, or up to 100 mg of hydrocortisone as pre-medication for administration of certain medications or blood products prior to enrolment in this study.
6. Subjects who have received any investigational agents within 28 days or 5 half-lives (whichever is longer) of the first dose (Cycle 1, Day 1).
7. Prior history of malignancies, other than MM, unless the subject has been free of the disease for 3 years or longer. Exceptions include the following:
 - Basal or squamous cell carcinoma of the skin
 - Carcinoma in situ of the cervix or breast
 - Adenocarcinoma of the prostate (TNM stage of T1a or T1b)
8. Other concurrent severe and/or uncontrolled medical conditions (i.e. uncontrolled diabetes, active or uncontrolled infection, acute diffuse pulmonary disease, pericardial disease, uncontrolled thyroid dysfunction) including abnormal laboratory values, that could cause unacceptable safety risks or compromise compliance with the protocol.
9. Known chronic obstructive pulmonary disease (COPD), defined as a FEV1 $<$ 50% predicted value.
10. Known moderate or severe persistent asthma within the last 2 years, or currently has uncontrolled asthma of any classification.
11. History of or current uncontrolled cardiovascular disease including:
 - Unstable angina, myocardial infarction, or known congestive heart failure Class III/IV (Appendix 5) within the preceding 12 months
 - Transient ischemic attack within the preceding 3 months, pulmonary embolism within the preceding 2 months.
 - Any of the following: sustained ventricular tachycardia, ventricular fibrillation, Torsades de Pointes, cardiac arrest, Mobitz II second degree heart block or third degree heart block; known presence of dilated, hypertrophic, or restrictive cardiomyopathy.
 - QTc prolongation as confirmed by ECG assessment at screening (QTc $>$ 470 milliseconds).
12. Women who are pregnant, breastfeeding or planning to become pregnant while enrolled in this study, or within 90 days after the last dose of study medications. Male subject who plans to father a child while enrolled in this study, within 90 days after the last dose of study medications.
13. Known HIV positivity or active infectious hepatitis B or C
14. Known allergies, hypersensitivity to mannitol, corticosteroids, monoclonal antibodies or human proteins, or their excipients (refer to the Daratumumab IB), or known sensitivity to mammalian-derived products

15. Known CNS involvement, plasma cell leukemia or amyloidosis.
16. Subjects who are receiving any other investigational agent
17. Autologous, peripheral stem cell transplant within 12 weeks of the first dose of study drug
18. Any other condition that, in the Investigator's opinion, would contraindicate the patient's participation in the clinical study due to safety concerns or compliance with clinical study procedures.

Phase 3, Randomized Multicenter Study of Subcutaneous vs. Intravenous Administration of Daratumumab in Subjects with Relapsed of Refractory Multiple Myeloma (Study Number: 54767414MMY3012)

Inclusion Criteria

1. At least 18 years of age.
2. Documented multiple myeloma as defined by the criteria below:
 - Multiple myeloma diagnosis according to the IMWG diagnostic criteria (refer to Attachment 1).
 - Measurable disease at Screening as defined by any of the following:
 - IgG multiple myeloma: Serum M-protein level ≥ 1.0 g/dL or urine M-protein level ≥ 200 mg/24 hours; or
 - IgA, IgD, IgE, IgM multiple myeloma: serum M-protein level ≥ 0.5 g/dL or urine M-protein level ≥ 200 mg/24 hours; or
 - Light chain multiple myeloma without measurable disease in the serum or the urine: Serum immunoglobulin FLC ≥ 10 mg/dL and abnormal serum immunoglobulin Kappa lambda FLC ratio.
3. Evidence of a response (PR or better based on investigator's determination of response by IMWG criteria) to at least 1 prior treatment regimen.
4. Relapsed or refractory disease as defined below:
 - Relapsed disease is defined as an initial response to previous treatment, followed by Confirmed PD by IMWG criteria >60 days after cessation of treatment.
 - Refractory disease is defined as $<25\%$ reduction in M-protein or confirmed PD by IMWG criteria during previous treatment or ≥ 60 days after cessation of treatment.
5. Received at least 3 prior lines of therapy (refer to Attachment 2) including a PI (≥ 2 cycles or 2 months of treatment) and an IMiD (≥ 2 cycles or 2 months of treatment) in any order during the course of treatment (except for subjects who discontinued either of these treatments due to a severe allergic reaction within the first 2 cycles/months).
 - A single line of therapy may consist of 1 or more agents, and may include induction, hematopoietic stem cell transplantation, and maintenance therapy. Radiotherapy, bisphosphonate, or a single short course of corticosteroids (no more than the equivalent of dexamethasone 40 mg/day for 4 days) would not be considered prior lines of therapy.
 - or
 - Refractory to both a PI and an IMiD. For subjects who have received more than 1 type of PI, their disease must be refractory to the most recent one. Similarly, for those who have received more than 1 type of IMiD, their disease must be refractory to the most recent one.
6. ECOG Performance Status score of 0, 1, or 2 (refer to Attachment 3).
7. Pretreatment clinical laboratory values meeting the following criteria during the Screening Phase:
 - hemoglobin ≥ 7.5 g/dL (≥ 7 mmol/L) (without prior red blood cells [RBC] transfusion within 7 days before the laboratory test; recombinant human erythropoietin use is permitted);
 - absolute neutrophil count $\geq 1.0 \times 10^9$ /L (prior growth factor support is permitted);
 - platelet count $\geq 50 \times 10^9$ /L (transfusions are not permitted within 7 days of testing to achieve this minimum platelet count);
 - aspartate aminotransferase (AST) $\leq 2.5 \times$ upper limit of normal (ULN);
 - alanine aminotransferase (ALT) $\leq 2.5 \times$ ULN;

- total bilirubin $\leq 2.0 \times \text{ULN}$; except in subjects with congenital bilirubinemia, such as Gilbert syndrome (in which case direct bilirubin $\leq 2.0 \times \text{ULN}$ is required);
 - estimated creatinine clearance ≥ 30 mL/min per 1.73m^2 (refer to Attachment 4);
 - albumin-corrected serum calcium ≥ 14 mg/dL (≥ 3.5 mmol/L) or free ionized calcium ≥ 6.5 mg/dL (≥ 1.6 mmol/L) (refer to Attachment 5).
8. Women of childbearing potential must commit to either abstain continuously from heterosexual sexual intercourse or to use 2 methods of reliable birth control simultaneously. This includes one highly effective form of contraception (tubal ligation, intrauterine device, hormonal [birth control pills, injections, hormonal patches, vaginal rings or implants] or partner's vasectomy) and one additional effective contraceptive method (male latex or synthetic condom, diaphragm, or cervical cap). Contraception must begin 4 weeks prior to dosing. Reliable contraception is indicated even where there has been a history of infertility, unless due to hysterectomy.
 9. Women of childbearing potential must have a negative urine or serum pregnancy test at screening within 14 days prior to randomization.
 10. Each subject (or their legally acceptable representative) must sign an Informed Consent Form (ICF) indicating that he or she understands the purpose of and procedures required for the study and are willing to participate in the study. Subjects must be willing and able to adhere to the prohibitions and restrictions specified in this protocol, as referenced in the ICF.

Exclusion Criteria

1. Received daratumumab or other anti-CD38 therapies previously.
2. Received anti-myeloma treatment within 2 weeks or 5 pharmacokinetic half-lives of the treatment, whichever is longer, before the date of randomization. The only exception is emergency use of a short course of corticosteroids (equivalent of dexamethasone 40 mg/day for a maximum of 4 days [refer to Attachment 6]) before treatment. A list of anti-myeloma treatments with the corresponding pharmacokinetic half-lives is provided in the Site Investigational Product Procedures Manual.
3. Received autologous stem cell transplant within 12 weeks before the date of randomization, or the subject has previously received allogeneic stem cell transplant (regardless of timing).
4. Plans to undergo a stem cell transplant prior to progression of disease on this study (these subjects should not be enrolled to reduce disease burden prior to transplant).
5. History of malignancy (other than multiple myeloma) within 3 years before the date of randomization (exceptions are squamous and basal cell carcinomas of the skin and carcinoma in situ of the cervix, or breast, or other non-invasive lesion, that in the opinion of the investigator, with concurrence with the sponsor's medical monitor, is considered cured with minimal risk of recurrence within 3 years).
6. Clinical signs of meningeal involvement of multiple myeloma.
7. Either of the following:
 - Known chronic obstructive pulmonary disease (COPD) with a forced expiratory volume in 1 second (FEV1) is $< 50\%$ of predicted normal. Note that FEV1 testing also is required for subjects suspected of having COPD and subjects must be excluded if FEV1 is $< 50\%$ of predicted normal.
 - Known moderate or severe persistent asthma, or a history of asthma within the last 2 years, or currently has uncontrolled asthma of any classification (refer to Attachment 7). (Subjects who currently have controlled intermittent asthma or controlled mild persistent asthma are allowed to participate in the study.)
8. Any of the following:
 - Known to be seropositive for human immunodeficiency virus (HIV)
 - Known to be seropositive for hepatitis B (defined by a positive test for hepatitis B surface antigen [HBsAg]) or with known prior hepatitis B infection without evidence of immunity (ie, patients who are positive for antibodies to hepatitis B core antigen [anti-HBc] but negative for antibodies to hepatitis B surface antigen [anti-Hbs])
 - Known to be seropositive for hepatitis C (except in the setting of a sustained virologic response [SVR], defined as aviremia at least 12 weeks after completion of antiviral therapy).

9. Concurrent medical or psychiatric condition or disease (eg, active systemic infection, uncontrolled diabetes, acute diffuse infiltrative pulmonary disease) that is likely to interfere with study procedures or results, or that in the opinion of the investigator would constitute a hazard for participating in this study.
10. Clinically significant cardiac disease, including:
 - Myocardial infarction within 6 months before date of randomization, or unstable or uncontrolled disease/condition related to or affecting cardiac function (eg, unstable angina, congestive heart failure, New York Heart Association Class III-IV [refer to Attachment 8]).
 - Uncontrolled cardiac arrhythmia (Grade 2 or higher by NCI-CTCAE Version 4.03) or clinically significant ECG abnormalities.
 - Screening 12-lead ECG showing a baseline QT interval as corrected by Fridericia's formula >470 msec.
11. Known allergies, hypersensitivity, or intolerance to any of the study drugs, hyaluronidase, mAbs, human proteins, or their excipients (refer to daratumumab IB10), or known sensitivity to mammalian-derived products.
12. Plasma cell leukemia (>2.0 × 10⁹/L circulating plasma cells by standard differential) or Waldenstrom's macroglobulinemia or POEMS syndrome (polyneuropathy, organomegaly, endocrinopathy, M-protein, and skin changes) or amyloidosis.
13. Known or suspected of not being able to comply with the study protocol (eg, because of alcoholism, drug dependency, or psychological disorder) or the subject has any condition for which, in the opinion of the investigator, participation would not be in the best interest of the subject (eg, compromise their well-being) or that could prevent, limit, or confound the protocol-specified assessments.
14. Pregnant, breast-feeding, or planning to become pregnant while enrolled in this study or within 3 months after the last dose of study drug.
15. Plans to father a child while enrolled in this study or within 3 months after the last dose of study drug.
16. Received an investigational drug (including investigational vaccines) or used an invasive investigational medical device within 4 weeks before the planned first dose of study drug (except for investigational anti-myeloma treatments, which cannot be taken within 2 weeks before Cycle 1 Day 1).
17. Major surgery within 2 weeks before randomization, or has not fully recovered from an earlier surgery, or has major surgery planned during the time the subject is expected to participate in the study or within 2 weeks after the last dose of study drug administration. Kyphoplasty or vertebroplasty are not considered major surgery. Note: subjects with planned surgical procedures to be conducted under local anesthesia may participate. If there is a question whether a procedure is considered a major surgery, the investigator must consult with the appropriate sponsor representative and resolve any issues before enrolling a subject in the study.
18. Plasmapheresis within 28 days before randomization.

Phase 1/2 Trial of Idasanutlin in combination with Ixazomib and dexamethasone in patients with 17p deleted, relapsed multiple myeloma (MC1582/MMRC-061; Protocol#: MC1582/MMRC-061)

Inclusion Criteria

1. Diagnosis of MM with deletion 17p (del17p) or monosomy 17 by FISH who have received at least one line of therapy.
2. The following laboratory values obtained ≤14 days prior to registration.
 - Calculated creatinine clearance ≥30 mL/min
 - AST (SGOT) and ALT (SGPT) ≤3.0 x upper limit of normal (ULN)
 - Total bilirubin ≤1.5 × the upper limit of the normal range (ULN)
 - Absolute neutrophil count (ANC) ≥1500/mm³
 - Platelet count ≥75,000/mm³
 - Hemoglobin ≥8.0 g/dL

NOTE: White blood count and platelet count criteria must be met without any transfusion or growth factor support.

3. Patients with measurable disease defined as at least one of the following:
 - Serum monoclonal protein ≥ 1.0 g/dL by protein electrophoresis
 - >200 mg of monoclonal protein in the urine on 24-hour electrophoresis
 - Serum immunoglobulin free light chain ≥ 10 mg/dL AND abnormal serum immunoglobulin kappa to lambda free light chain ratio.
4. ECOG performance status 0, 1 or 2

Exclusion Criteria

1. Other malignancy requiring active therapy.
 - EXCEPTIONS: Non-melanoma skin cancer, DCIS or carcinoma-in-situ of the cervix.
 - NOTE: If there is a history of prior malignancy, they must not be receiving other specific treatment for their cancer
2. Other concurrent chemotherapy, radiotherapy, or any ancillary therapy considered investigational.
 - NOTE: Bisphosphonates are considered to be supportive care rather than therapy, and are thus allowed while on protocol treatment.
3. Patient has $>$ Grade 2 peripheral neuropathy, or Grade 1 with pain on clinical examination during the screening period.
4. All CYP2C8 inhibitors, inducers, and substrates should be discontinued ≥ 7 days prior to registration. Systemic treatment with CYP2C8 inhibitors (anastrozole, montelukast, quercetin, trimethoprim, gemfibrozil, rosiglitazone, pioglitazone), inducers (carbamazepine, phenytoin, rifabutin, rifampin), or substrates (amiodarone, repaglinide, rosiglitazone, sorafenib, toremide) should be discontinued ≥ 7 days prior to registration.
5. Systemic treatment with strong inhibitors of CYP3A4 (clarithromycin, telithromycin, itraconazole, Voriconazole, ketoconazole, nefazodone, posaconazole) or strong CYP3A4 inducers (rifampin, rifapentine, rifabutin, carbamazepine, phenytoin, phenobarbital, Gingko biloba, St. John's wort) are not allowed ≤ 14 days before registration.
6. Evidence of current uncontrolled cardiovascular conditions, including cardiac arrhythmias, congestive heart failure, angina, or myocardial infarction within the past 6 months. Note: Prior to study entry, any ECG abnormality at screening must be documented by the investigator as not medically relevant.
7. QTc >470 milliseconds (msec) on a 12-lead ECG obtained during the Screening period.
8. Known human immunodeficiency virus (HIV) positive.
9. Known hepatitis B surface antigen-positive status, or known or suspected active hepatitis C infection
10. Known GI disease or GI procedure that could interfere with the oral absorption or tolerance of ixazomib or idasanutlin including difficulty swallowing
11. Diarrhea $>$ Grade 1, based on the NCI CTCAE grading, or currently taking antidiarrheals
12. Need for ongoing therapeutic anticoagulation.
13. Patients that have previously been treated with ixazomib, or who participated in a blinded study with ixazomib (whether treated with ixazomib or not).

A Phase 1b Study Evaluating the Safety, Tolerability, Pharmacokinetics and Efficacy of Oprozomib in Combination with Pomalidomide and Dexamethasone in Subjects with Relapsed or Refractory Multiple Myeloma (Protocol: 20160104)

Inclusion Criteria

1. Subject must have a pathologically documented, definitively diagnosed, multiple myeloma relapse, or refractory progressive disease after at least 2 lines of therapy for multiple myeloma. Prior therapeutic treatment or regimens must include a proteasome inhibitor and lenalidomide
2. Measurable disease (assessed within 28 days prior to day 1), as indicated by one or more of the following:
 - Serum M-protein ≥ 0.5 g/dL

- Urine M-protein ≥ 200 mg/24 hours
 - In subjects without detectable serum or urine M-protein: serum Free Light
 - Chain (sFLC) ≥ 10 mg/dL (≥ 100 mg/L) and an abnormal sFLC ratio
3. Eastern Cooperative Oncology Group (ECOG) performance status of ≤ 2
 4. Hematological function, as follows, without transfusion support:
 - Absolute neutrophil count $\geq 1.0 \times 10^9/L$
 - Platelet count $\geq 75 \times 10^9/L$ (in patients with $< 50\%$ of bone marrow nucleated cells were plasma cells) or $\geq 50 \times 10^9/L$ (in patients with $\geq 50\%$ of bone marrow nucleated cells were plasma cells) without transfusion or growth factor support
 - Hemoglobin > 8 g/dL (> 80 g/L) Use of erythropoietic stimulating factors and red blood cell (RBC) transfusions per institutional guidelines is allowed, however most recent RBC transfusion must not be within 7 days prior to obtaining screening hemoglobin
 5. Coagulation function as follows: PT/INR and PTT < 1.5 x Institutional Upper Limit of Normal (ULN)
 6. Renal function as follows: estimated glomerular filtration rate based on Modification of Diet in Renal Disease calculation (MDRD) > 30 mL/min/1.73 m²
 7. Hepatic function, as follows: AST and ALT < 3 x ULN, Total bilirubin < 1.5 x ULN (except subjects with Gilbert's syndrome)

Exclusion Criteria

1. Currently receiving treatment in another investigational device or drug study, or less than 28 days since ending treatment on another investigational device or drug study(s)
2. Previously received an allogeneic stem cell transplant and the occurrence of one or more of the following:
 - Received the transplant within 6 months prior to study day 1
 - Received immunosuppressive therapy within the last 3 months prior to study day
 - Having signs or symptoms of acute or chronic graft-versus-host disease
3. Autologous stem cell transplant < 90 days prior to study day 1
4. Multiple myeloma with IgM subtype
5. POEM syndrome (polyneuropathy, organomegaly, endocrinopathy, monoclonal protein, and skin changes)
6. Plasma cell leukemia ($> 2.0 \times 10^9/L$ circulating plasma cells by standard differential)
7. Waldenstrom's macroglobulinemia, Amyloidosis
8. Requirement for plasmapheresis during the screening period
9. Dexamethasone at cumulative doses of greater than 160 mg or equivalent within 21 days prior to study day 1 is not allowed. Use of topical or inhaled steroids is Acceptable
10. History of other malignancy, except:
 - Current use of therapeutic doses of anticoagulation unless agreed upon by the investigator and the Amgen Medical Monitor. Please note: thromboprophylaxis is recommended with pomalidomide treatment
11. History of clinically significant GI hemorrhage (Grade ≥ 2) in the 6 months prior to study day 1, unless agreed upon by the investigator and the Amgen Medical Monitor
12. Known positive results for Human Immunodeficiency Virus (HIV)
13. Prior systemic radiation therapy must have been completed at least 28 days before study day 1. Prior focal radiotherapy completed at least 14 days before study day 1
14. Prior use of pomalidomide if subjects required pomalidomide dose reduction or pomalidomide discontinuation due to toxicity

A Phase II Open Label, Multicenter, Trial of JNJ-42756493 in Combination with Dexamethasone for the Treatment of FGFR3 Wild-type or Mutation Positive Relapsed and/or Refractory Multiple Myeloma (Protocol: PM-MM003)

Inclusion Criteria

Patients must meet all of the following inclusion criteria to be eligible for participation in this study.

1. A diagnosis of MM and documentation of at least 1 prior line of therapy including proteasome and immunomodulatory agents (in separate regimens or in combination).
2. Documented lab results confirming FGFR3 expression and mutational status determined by a clinical grade, next generation sequencing platform (e.g. Michigan Center for Translational Pathology, Foundation Medicine, Inc.) approved by the Sponsor-Investigator, the results of which must be obtained prior to registration.
3. Patients with measurable disease defined as at least one of the following (these baseline laboratory studies for determining eligibility must be obtained within 28 days prior to start of study drug):
 - Serum M-protein ≥ 0.5 g/dl (≥ 5 g/l)
 - Urine M-protein ≥ 200 mg/24 h
 - Serum free light chains (FLC) assay: Involved FLC level ≥ 10 mg/dl (≥ 100 mg/l) and an abnormal serum free light chain ratio (< 0.26 or > 1.65)
4. The following laboratory results must be met within 7 days of first study drug administration:
 - Absolute neutrophil count (ANC) $\geq 1,000$ cells/dL ($1.0 \times 10^9/L$). Growth factors cannot be given within 7 days of study drug administration.
 - Serum AST and ALT ≤ 2.5 x upper limit of normal (ULN).
 - Creatinine clearance ≥ 40 mL/min either directly measured via 24-hour urine collection or calculated using Cockcroft-Gault.
 - Platelet count $\geq 50,000$ cells/dL ($50 \times 10^9/L$). Platelet transfusions to help patients meet eligibility criteria are not allowed within 7 days before study enrollment.
 - Hemoglobin ≥ 8.0 g/dL.
 - Total bilirubin ≤ 1.5 x ULN, unless known to have Gilbert's disease.
 - Albumin ≥ 2.0 g/dL (20 g/L).
 - Electrolytes: Magnesium within 0.85 to 1.25 x institutional ULN; Sodium ≥ 130 mEq/L; and potassium within institutional normal limits (correction with supplementation and re-testing is permitted).

Exclusion Criteria

Patients who meet any of the following exclusion criteria are not eligible for enrollment.

1. Patients in whom FGFR3 expression or mutational status cannot be determined.
2. Chemotherapy, limited palliative radiotherapy or other anti-myeloma therapy within 14 days prior to the first dose of study drug. In addition, any treatment-related toxicity should have recovered $< \text{Grade } 1$ unless deemed to be irreversible (an example of an irreversible toxicity would include steroid induced cataracts or peripheral neuropathy).
3. Patients who are receiving any other investigational agent.
4. Patients with known CNS involvement, plasma cell leukemia or amyloidosis.
5. Use of an investigational drug within 21 days or five-half-lives, whichever is shorter but not less than 14 days, preceding the first dose of study drug.
6. History of allogeneic stem cell transplant.
7. Autologous, peripheral stem cell transplant within 12 weeks of the first dose of study drug.

MCRN 005 – The BOSTON Trial: A PHASE 3 RANDOMIZED, CONTROLLED, OPEN-LABEL STUDY OF SELINEXOR, BORTEZOMIB, AND DEXAMETHASONE (SVD) VERSUS BORTEZOMIB AND DEXAMETHASONE (VD) IN PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA (RRMM) - Protocol number: KCP-330-023

Key Inclusion Criteria

1. Documented evidence of progressive MM (based on the Investigator's determination according to the modified IMWG response criteria) on or after their most recent regimen.
2. Prior treatment with bortezomib or other PI is allowed. Must have had at least a 6-month PI-treatment-free interval prior to C1D1 of study treatment.
3. Resolution of any clinically significant non-hematological toxicities (if any) from previous treatments to \leq Grade 1 by C1D1.
4. Adequate hepatic function within 28 days prior to C1D1:
 - Total bilirubin $< 1.5 \times$ upper limit of normal (ULN) (except patients with Gilbert's syndrome who must have a total bilirubin of $< 3 \times$ ULN), and
 - Aspartate aminotransferase (AST) and alanine aminotransferase (ALT) normal to $< 2 \times$ ULN.
5. Adequate renal function within 28 days prior to C1D1 (estimated creatinine clearance [CrCl] of ≥ 20 mL/min)
6. Adequate hematopoietic function within 7 days prior to C1D1: total white blood cell (WBC) count $\geq 1500/\text{mm}^3$, absolute neutrophil count $\geq 1000/\text{mm}^3$, hemoglobin ≥ 8.5 g/dL and platelet count $\geq 75,000/\text{mm}^3$ (patients for whom $< 50\%$ of bone marrow nucleated cells are plasma cells) or $\geq 50,000/\text{mm}^3$ (patients for whom $\geq 50\%$ of bone marrow nucleated cells are plasma cells).
 - Patients receiving hematopoietic growth factor support must have a 2-week interval between growth factor support and the Screening assessments, but they may receive growth factor support during the study.
 - Patients must have at least a 2-week interval from the last red blood cell (RBC) transfusion and 1-week interval prior to the Screening

Key Exclusion Criteria

1. Has received Selinexor or another XPO1 inhibitor previously.
2. Prior malignancy that required treatment, or has shown evidence of recurrence
3. Has any concurrent medical condition or disease (e.g., uncontrolled active hypertension, uncontrolled active diabetes, active systemic infection, active, unstable cardiovascular function).
4. Active plasma cell leukemia, systemic light chain amyloidosis, Polyneuropathy, organomegaly, endocrinopathy, monoclonal gammopathy, and skin changes (POEMS) syndrome, MM involving the central nervous system or Spinal cord compression.
5. Greater than Grade 2 neuropathy or \geq Grade 2 neuropathy with pain at baseline, regardless of whether or not the patient is currently receiving medication.
6. Intolerance, hypersensitivity, or contraindication to glucocorticoids.
7. Radiation, chemotherapy, or immunotherapy or any other anticancer therapy ≤ 2 weeks prior to C1D1.
8. Prior autologous stem cell transplantation < 1 month or allogeneic stem cell transplantation < 4 months prior to C1D1.
9. Active graft versus host disease (after allogeneic stem cell transplantation) at C1D1.
10. BSA < 1.4 m² at baseline.
11. Life expectancy of < 4 months.

An Open-Label, Randomized Phase 3 Study of Combinations of Nivolumab, Elotuzumab, Pomalidomide and Dexamethasone in Relapsed and Refractory Multiple Myeloma (Protocol number: CA209-602) - Enrollment on Hold

Inclusion Criteria

1. Signed Written Informed Consent
2. Target Population
 - Must have received 2 prior lines of therapy which must have included at least 2 consecutive cycles of each immune modulatory drug (IMiD) and a proteasome inhibitor alone or in combination
 - Documented refractory or relapsed and refractory (R/R) multiple myeloma
 - Refractory (progressed on or within 60 days of treatment) to their last treatment
 - Subjects must have failed treatment with a proteasome inhibitor and an IMiD in one of the following ways
 - i. “Double Refractory” = Refractory to a proteasome inhibitor and an IMiD, and to their last treatment
 - ii. “Relapsed and refractory” = patients had achieved at least a partial response to previous treatment with proteasome inhibitor or IMiD, or both, but progressed within 6 months, and were refractory to their last treatment
 - Measurable disease at screening, based on central lab results within 28 days of randomization.
 - Eastern Cooperative Oncology Group (ECOG) performance status ≤ 2
3. Age and Reproductive Status
 - Males and Females at least 18 years or legal age of consent per local regulations
 - Women of childbearing potential (WOCBP) must have two negative serum or urine pregnancy tests.
 - i. Men who are sexually active with WOCBP must agree for method(s) of contraception for the duration of treatment with study drug plus 5 half-lives of study drug plus 90 days (duration of sperm turnover) for a total of 31 weeks (except the Pd arm -where only 4 weeks is required) post-treatment completion
 - Women must not be breastfeeding
 - Male patients must not donate sperm, for up to 180 days (4 weeks only for Pd arm) post treatment completion
 - Azoospermic males and WOCBP who are not heterosexually active are exempt from contraceptive requirements. However, they must still undergo pregnancy testing as described in this section
 - All subjects must not donate blood for 90 days post treatment completion
 - All subjects must be willing and able to comply with Pomalyst® REMS program, where applicable
 - All subjects must agree not to share study medication

Exclusion Criteria

1. Target Disease Exceptions
 - Subjects with solitary bone or extramedullary plasmacytoma as the only evidence of plasma cell dyscrasia
 - Subjects with monoclonal gammopathy of undetermined significance (MGUS), smoldering multiple myeloma (SMM), amyloidosis, Waldenström’s Macroglobulinemia, or POEMS syndrome (plasma cell dyscrasia with poly neuropathy, organomegaly, endocrinopathy, monoclonal protein, and skin changes)
 - Subjects with active plasma cell leukemia (defined as either 20% of peripheral blood white blood cell count comprised of plasma/CD138+ cells or an absolute plasma cell count of $2 \times 10^9/L$)
2. Medical History and Concurrent Diseases
 - Women who are of childbearing potential not complying to the above described contraceptive measures or are breastfeeding, and sexually active fertile men whose partners are WOCBP if they are not complying to the above described contraceptive measures
 - Any uncontrolled or severe cardiovascular or pulmonary disease determined by the investigator, including:
 - i. NYHA functional classification III or IV, congestive heart failure, unstable or poorly controlled angina, uncontrolled hypertension, arrhythmia, or myocardial infarction in the past 12 months

ii. Subjects with interstitial lung disease that is symptomatic or may interfere with the detection or management of suspected drug-related pulmonary toxicity

- Active infection
- Subjects with an active, known or suspected autoimmune disease. Subjects with type I diabetes mellitus, hypothyroidism only requiring hormone replacement, skin disorders (such as vitiligo, psoriasis, or alopecia) not requiring systemic treatment, or conditions not expected to recur in the absence of an external trigger are permitted to enroll
- Subjects with a condition requiring systemic treatment with either corticosteroids (> 10 mg daily prednisone equivalent) or other immunosuppressive medications within 14 days of initiation of study drug. Inhaled or topical steroids, and adrenal replacement steroid doses > 10 mg daily prednisone equivalent, are permitted in the absence of active autoimmune disease
- Unable to tolerate thromboembolic prophylaxis while on the study
- Hypersensitivity reaction to prior IMiD (thalidomide or lenalidomide)
- Grade ≥ 2 peripheral neuropathy (per NCI CTCAE v4.0)
- Any positive test for hepatitis B virus or hepatitis C virus indicating acute or chronic infection
- Known history of positive test for human immunodeficiency virus (HIV) or known acquired immunodeficiency syndrome (AIDS). NOTE: Testing for HIV must be performed at sites where mandated by local regulation
- Gastrointestinal disease that may significantly alter the absorption of pomalidomide
- Prior or concurrent invasive malignancy, except for the following:
 - i. Adequately treated basal cell or squamous cell skin cancer
 - ii. Adequately treated in-situ cancer
 - iii. Any cancer (other than those noted above) from which the subject has been disease free for > 3 years prior to study entry

3. Prior Therapy or Surgery

- Prior treatment with pomalidomide, Nivolumab (or any PD-1 or PD-L1 inhibitor) or Elotuzumab
- Use of any anti-myeloma drug therapy, within 14 days of the initiation of study drug treatment or use of any experimental drug therapy or plasmapheresis within 28 days (or 5 half-lives of the experimental drug; whichever is longer) of the initiation of study drug treatment (includes dexamethasone). Bisphosphonate use permitted if initiated prior to first dose of study medication
- Treatment with melphalan or monoclonal antibodies within 4 weeks (or 5 half-lives of the monoclonal antibody; whichever is longer) of the first dose of study drug
- Prior autologous stem cell transplant within 12 weeks of the first dose of study drug
- Prior allogeneic stem cell transplant except subjects who have completed the stem cell transplant > 12 months prior to first dose of study drug, have no current or history of graft versus host disease, and are not on topical or systemic immunosuppressive therapy
- Treatment with corticosteroids within 2 weeks of the first dose of study drug, except for the equivalent of 10 mg prednisone per day or corticosteroids with minimal to no systemic absorption (ie, topical or inhaled steroids) or for short course (4 days) of 40 mg dexamethasone or equivalent for emergency use (baseline M proteins must be drawn after this short course and prior to randomization). Adrenal replacement steroid doses > 10 mg daily prednisone equivalent, are permitted in the absence of active autoimmune disease
- Major cardiac surgery within 8 weeks prior to the first dose of study drug; all other major surgery within 4 weeks prior to the first dose of study drug. (Kyphoplasty is not considered major surgery); subjects should have been fully recovered from any surgical related toxicities

4. Physical and Laboratory Test Findings

Screening Laboratory evaluations within the following parameters:

- Absolute neutrophil count (ANC) < 1,000 cells/ μ L (1.0×10^9 /L) (Growth factors cannot be used within 1 week of first drug administration. No pegylated growth factors within 3 weeks of first drug administration)
- Platelet count < 75,000 cells/ μ L (75×10^9 /L) (< 30×10^9 /L if $\geq 50\%$ of bone marrow nucleated cells were plasma cells). Qualifying laboratory value must occur at most recent measurement prior to study entry. No transfusions are allowed within 72 hours prior to qualifying laboratory value

- Hemoglobin < 8 g/dl (No transfusions are allowed within 72 hours prior to qualifying laboratory value)
 - Total Bilirubin > 1.5 X upper limit of normal (ULN) (except subjects with Gilbert Syndrome, who can have total bilirubin up to 3.0 X ULN)
 - AST (SGOT) and ALT (SGPT) > 3.0x ULN
 - Renal function: Estimated creatinine clearance by Cockcroft-Gault formula < 30 mL/min
 - Corrected serum calcium \geq 11.5 mg/dl within 2 weeks of initiation of study drug (despite appropriate measures such a short course of steroids, bisphosphonates, hydration, calcitonin)
5. Allergies and Adverse Drug Reaction a) History of allergy or hypersensitivity to study drug components

A Phase 2, Multicenter, Open-label, Study to Determine the Safety and Efficacy for the Combination of Durvalumab (DURVA) and Daratumumab (DARA) (D2) in Subjects with Relapsed and Refractory Multiple Myeloma (Protocol Number: MEDI4736-MM-003) – Enrollment on Hold

Inclusion Criteria

Subjects must satisfy the following criteria to be enrolled in the study:

1. Subject received at least 3 prior anti-myeloma regimen including a PI and an immunomodulatory agent or is double-refractory to a PI and an immunomodulatory agent.
 - Induction, bone marrow transplant with or without maintenance therapy is considered one regimen
 - Refractory is defined as disease that is nonresponsive on therapy, or progresses within 60 days of last therapy. Nonresponsive disease is defined as either failure to achieve minimal response or development of progressive disease while on therapy.
 - For subjects who received more than 1 regimen containing a PI their disease must be refractory to the most recent PI containing regimen.
 - For subjects who received more than 1 regimen containing an immunomodulatory agent their disease must be refractory to the most recent immunomodulatory agent containing regimen
2. Subject has measurable disease defined as:
 - M-protein (serum protein electrophoresis (sPEP) or urine protein electrophoresis (uPEP): sPEP \geq 0.5 g/dL or uPEP \geq 200 mg/24 hours) and/or
 - Light chain MM without measurable disease in the serum or the urine: serum immunoglobulin free light chain \geq 10 mg/dL and abnormal serum immunoglobulin kappa lambda free light chain ratio
3. Subject achieved a response (MR or better) to at least 1 prior treatment regimen
4. Subject has evidence of PD on or within 60 days of the most recent prior treatment regimen
5. Subject received an alkylating agent alone or in combination with other myeloma treatment
6. Subject has an Eastern Cooperative Oncology Group performance-status score of 2 or less
7. Subject's toxicities resulting from previous therapy (including peripheral neuropathy) have resolved or stabilized to \leq Grade 1.

Exclusion Criteria:

The presence of any of the following will exclude a subject from enrollment:

1. Subject has had prior exposure to anti-CTLA-4, anti-PD-1, anti-PD-L1 mAbs, cell-based therapies (eg, CAR-T cells), or cancer vaccines
2. Subject received DARA or other anti-CD38 therapies previously
3. Subject received prior treatment with a monoclonal antibody within 5 half-lives of initiating study treatment

4. Subject used any investigational agents within 28 days or 5 half-lives (whichever is longer) of initiating study treatment
5. History of organ or allogeneic stem cell transplantation
6. Subject has any of the following laboratory abnormalities:
 - Absolute neutrophil count (ANC) < 1,000/ μ L
 - Platelet count: < 75,000/ μ L
 - Hemoglobin < 8 g/dL (< 4.9 mmol/L)
 - Creatinine Clearance (CrCl) < 45 mL/min
 - Corrected serum calcium > 13.5 mg/dL (> 3.4 mmol/L)
 - Serum aspartate aminotransferase (AST) or alanine aminotransferase (ALT) > 2.5 \times upper limit of normal (ULN)
 - Serum total bilirubin > 1.5 \times upper limit of normal (ULN) or > 3.0 mg/dL for subjects with documented Gilbert's syndrome
7. Subject has clinical evidence of central nervous system (CNS) or pulmonary leukostasis, disseminated intravascular coagulation, or CNS MM
8. Subject has known chronic obstructive pulmonary disease (COPD) with a forced expiratory volume in 1 second (FEV1) 50% of predicted normal.
9. Subject has known moderate or severe persistent asthma within the past 2 years or uncontrolled asthma of any classification. Note that subjects who currently have controlled intermittent asthma or controlled mild persistent asthma are allowed to participate in the study.
10. Subject has plasma cell leukemia, Waldenstrom's macroglobulinemia, POEMS syndrome, or amyloidosis
11. Subject has nonsecretory MM
12. Subject has active or prior documented autoimmune or inflammatory disorders (including inflammatory bowel disease [eg, colitis, Crohn's disease], diverticulitis, celiac disease, irritable bowel disease, or other serious gastrointestinal chronic conditions associated with diarrhea; systemic lupus erythematosus; Wegener syndrome; myasthenia gravis; Graves' disease; rheumatoid arthritis, hypophysitis, uveitis, etc.) within the past 3 years prior to the start of treatment. The following are exceptions to this criterion:
 - Subjects with vitiligo or alopecia.
 - Subjects with hypothyroidism (eg, following Hashimoto syndrome) stable on hormone replacement.
 - Psoriasis not requiring systemic treatment.
13. Subject has history of primary immunodeficiency
14. Subject is positive for human immunodeficiency virus (HIV), chronic or active hepatitis B or active hepatitis A or C.
15. Clinically significant abnormal electrocardiogram (ECG) finding at screening
16. Subject has prior history of malignancies, other than MM, unless the subject has been free of the disease for \geq 5 years with the exception of the following noninvasive malignancies:
 - Basal cell carcinoma of the skin
 - Squamous cell carcinoma of the skin
 - Carcinoma in situ of the cervix
 - Carcinoma in situ of the breast
 - Incidental histologic finding of prostate cancer (T1a or T1b using the TNM [tumor, nodes, metastasis] clinical staging system) or prostate cancer that is curative

For subjects who will have POM + dex added to the D2 and subjects who will be enrolled into the PD3 cohort, the following exclusions will also apply:

17. Subject has history of anaphylaxis or hypersensitivity to thalidomide, LEN, POM, or dex
18. Subject has history of rash \geq Grade 3 during prior thalidomide, LEN, or POM therapy
19. Subject has known or suspected hypersensitivity to the excipients contained in the formulation of POM or dex

20. Subject is a current smoker

A Phase I Trial of MK-3475 in Combination with Lenalidomide and Dexamethasone in Subjects with Multiple Myeloma (Protocol: Merck 023)

Inclusion Criteria

1. Be willing and able to provide written informed consent/assent for the trial. The subject may also provide consent/assent for Future Biomedical Research. However, the subject may participate in the main trial without participating in Future Biomedical research.
2. Be 18 years of age on day of signing informed consent.
3. Has a confirmed diagnosis of multiple myeloma based on standard criteria (see Durie 1986 for criteria [55]).
4. Currently has MM with measurable disease, defined as:
 - A monoclonal immunoglobulin spike on serum electrophoresis of at least 0.5 g/dL and/or
 - Urine monoclonal protein levels of at least 200 mg/24 hours
 - For subjects without measurable serum and urine M-protein levels, an abnormal free light chain ratio (normal value: 0.26 - 1.65) with involved FLC level ≥ 10 mg/dL (≥ 100 mg/L).
5. Has relapsed/refractory MM who has failed at least two lines of prior therapy, including bortezomib and an IMiD (thalidomide, pomalidomide, lenalidomide).

Relapsed MM defined as disease progression following stabilization or a response to at least one anti-myeloma regimen.

Refractory MM defined as meeting one or more of the following:

 - Nonresponsive to most recent therapy (e.g., stable disease only, or progressive disease while on treatment)
 - Disease progression within 60 days of discontinuation from most recent Therapy
6. Be able to provide archival (if available) and newly obtained bone marrow aspirate/biopsy material for biomarker analysis and disease assessment.
7. Has a performance status of 0 or 1 on the ECOG Performance Scale.
8. Demonstrate adequate organ function. All screening labs should be performed within 7 days of treatment initiation.
9. All subjects must agree to follow the regional requirements for lenalidomide counseling, pregnancy testing, and birth control; and be willing and able to comply with the regional requirements (for example, periodic pregnancy tests, safety labs, etc.).

Exclusion Criteria

1. Is currently participating in or has participated in a study of an investigational agent or using an investigational device within 4 weeks of the first dose of treatment.
2. Has myeloma and a history of repeated infections, primary amyloidosis, hyperviscosity, plasma cell leukemia, POEMS syndrome, Waldenstrom's Macroglobulinemia or IgM myeloma.
3. Has a diagnosis of immunosuppressive disorder or is on any other form of immunosuppressive therapy within 7 days prior to the first dose of trial treatment.
4. Has had a prior monoclonal antibody within 4 weeks prior to study Day 1 or who has not recovered (i.e. \leq Grade 1 or at baseline) from adverse events due to agents administered more than 4 weeks earlier.
5. Has had prior chemotherapy (including dexamethasone), targeted small molecule therapy, or radiation therapy within 2 weeks prior to study Day 1 or who has not recovered (i.e. \leq Grade 1 or at baseline) from adverse events due to a previously administered agent.
 - Note: Subjects with \leq Grade 2 neuropathy are an exception to this criterion and may qualify for the trial.
 - Note: If subject received major surgery, they must have recovered adequately from the toxicity and/or complications from the intervention prior to starting therapy.

- Note: Toxicity that has not recovered to \leq Grade 1 is allowed if it meets the inclusion requirements for laboratory parameters defined in Table 1.
6. Has been free of additional malignancy for at least 5 years. Exceptions include basal cell carcinoma of the skin, squamous cell carcinoma of the skin, or in situ cervical cancer that has undergone potentially curative therapy.
 7. Has known clinically active CNS involvement.
 8. Has an active autoimmune disease or a documented history of autoimmune disease, or a syndrome that requires systemic steroids or immunosuppressive agents. Subjects with vitiligo or resolved childhood asthma/atopy would be an exception to this rule. Subjects that require intermittent use of bronchodilators or local steroid injections would not be excluded from the study. Subjects with hypothyroidism stable on hormone replacement or Sjogren's syndrome will not be excluded from the trial.
 9. Has evidence of active, non-infectious pneumonitis.
 10. Has an active infection requiring intravenous systemic therapy.
 11. Has known psychiatric or substance abuse disorders that would interfere with cooperation with the requirements of the trial.
 12. Is pregnant or breastfeeding, or expecting to conceive or father children within the projected duration of the trial, starting with the pre-screening or screening visit through 120 days after the last dose of trial treatment.
 13. Has received prior therapy with an anti-PD-1, anti-PD-L1, anti-PD-L2, anti-CD137, or anti-Cytotoxic T-lymphocyte-associated antigen-4 (CTLA-4) antibody (including ipilimumab or any other antibody or drug specifically targeting T-cell co-stimulation or checkpoint pathways).
 14. Has a known Human Immunodeficiency Virus (HIV), Hepatitis B (HBV), or Hepatitis C (HCV) infection.
 15. Has a clinically significant coagulopathy per investigator's assessment.
 16. Has known symptomatic congestive heart failure, unstable angina pectoris, or cardiac arrhythmia.
 17. Has received an allogenic stem cell transplant.
 18. Has received autologous stem cell transplant within 12 weeks before the first infusion.
 19. Has received bortezomib, pomalidomide or thalidomide within 2 weeks before the first infusion.
 20. Subjects with a prior history of Grade 4 rash associated with thalidomide treatment.
 21. Has known hypersensitivity to thalidomide or pomalidomide.
 22. Is planning for or is eligible for allogenic hematopoietic stem cell transplant.
 23. Has known gastrointestinal disease that may significantly alter the absorption of lenalidomide.
 24. Is unable or unwilling to undergo antithrombotic prophylactic treatment.
 25. Has received a live vaccine within 30 days prior to first dose.

A Phase 1b/2 Study of Selinexor (KPT-330) in Combination with Backbone Treatments for Resistant/Refractory Multiple Myeloma (Protocol: KCP-330-017)

Inclusion Criteria

1. Written informed consent
2. Age \geq 18 years.
3. Histologically confirmed diagnosis, measurable disease and evidence of disease Progression of MM, based on IMWG guidelines.
4. Patients must have measurable disease as Defined by at least one of the following:
5. Serum M-protein \geq 0.5 g/dL by serum electrophoresis (SPEP) or for IgA myeloma, by quantitative IgA; or
6. Urinary M-protein excretion at least 200 mg/24 hours; or FLC \geq 100 mg/L, provided that FLC ratio is abnormal

7. If serum protein electrophoresis is felt to be unreliable for routine M-protein measurement (e.g., for IgA MM), then quantitative Ig levels by nephelometry or turbidometry are acceptable.
8. No evidence of prior drug-related toxicities to \geq Grade 2

SdP (Selinexor Dexamethasone Pomalidomide) Only:

1. Relapsed and refractory MM with:
 - Documented evidence of PD after achieving at least SD for \geq 1 cycle during previous MM treatment (i.e., relapsed MM).
 - Disease progression during or within 60 days from the end of the most recent MM treatment (i.e., refractory MM).
 - Previously undergone \geq 2 cycles of lenalidomide and a proteasome inhibitor (in separate regimens or in combination).

SdB (Selinexor Dexamethasone Bortezomib) Only:

1. Relapsed or refractory MM with
 - Documented evidence of relapse after \geq 1 previous line of therapy.
 - Not refractory to bortezomib in their most recent line of therapy.

Both SdP and SdB:

1. Eastern Cooperative Oncology Group (ECOG) Performance Status of \leq 2.
2. Adequate hepatic function within 21 days prior to Cycle 1 Day 1 (i.e., Day -21 to Day -1): Total bilirubin $<$ 2x ULN, AST $<$ 2.5x ULN and ALT $<$ 2.5x ULN.
3. Adequate renal function within 21 days prior to Cycle 1 Day 1: estimated creatinine clearance of \geq 45 mL/min, calculated using the formula of Cockcroft and Gault: $(140 - \text{Age}) \cdot \text{Mass (kg)} / (72 \cdot \text{creatinine mg/dL})$ multiply times 0.85 if the patient is female.
4. Female patients of child-bearing potential must agree to use dual methods of contraception and have a negative serum pregnancy test at screening. Male patients must use an effective barrier method of contraception if sexually active with a female of childbearing potential. Acceptable methods of contraception are condoms with contraceptive foam, oral, implantable or injectable contraceptives, contraceptive patch, intrauterine device, diaphragm with spermicidal gel, or a sexual partner who is surgically sterilized or post-menopausal. For both male and female patients, effective methods of contraception must be used throughout the study and for three months following the last dose.
5. Adequate hematopoietic function within 21 days prior to Cycle 1 Day 1: total WBC count \geq 1,500/mm³, ANC \geq 1000/mm³, hemoglobin (Hb) \geq 8.0 gm/dL, and platelet count \geq 75,000/mm³ for patients with plasma cells of $<$ 50% of bone marrow nucleated cells; or \geq 30,000/mm³ for patients with plasma cells of \geq 50% of bone marrow nucleated cells.
6. Patients receiving hematopoietic growth factor support, including erythropoietin (EPO), darbepoetin, granulocyte-colony stimulating factor (G-CSF), granulocyte macrophagecolony stimulating factor (GM-CSF), and platelet stimulators (e.g., eltrombopag or romiplostim or IL-11) may continue to do so.

Exclusion Criteria

26. Smoldering MM.
27. Multiple myeloma that does not express M-protein or FLC (i.e., non-secretory MM is excluded), and quantitative Ig levels cannot be used instead.
28. Documented systemic amyloid light chain amyloidosis.
29. Active MM involving the central nervous system (CNS).
30. Plasma cell leukemia.
31. Pregnant or breastfeeding.
32. Radiation, chemotherapy, or immunotherapy or any other anticancer therapy \leq 2 weeks prior to Cycle 1 Day 1, and radio-immunotherapy within 6 weeks prior to Cycle 1 Day 1. However, dexamethasone, up to 40 mg per week, is allowed as monotherapy up to the start of study treatment on Cycle 1 Day 1.

33. Treatment with an investigational anti-cancer therapy within 3 weeks prior to receiving first dose of study drug on Cycle 1 Day 1.

SdB arm only:

1. Prior history of neuropathy Grade > 2, or Grade 2 neuropathy with pain at screening (within 21 days prior to Cycle 1 Day 1).
2. Prior autologous stem cell transplantation < 1 month, or allogeneic stem cell transplantation < 3 months prior to Cycle 1 Day 1.
3. Active graft versus host disease after allogeneic stem cell transplantation. A life expectancy of < 3 months.
4. Major surgery within four weeks prior to Cycle 1 Day 1.
5. Unstable cardiovascular function:
 - Symptomatic ischemia, Or
 - Uncontrolled clinically-significant conduction abnormalities (e.g., patients with ventricular tachycardia on anti arrhythmics are excluded; patients with 1st degree atrioventricular (AV) block or asymptomatic left anterior fascicular block/right bundle branch block (LAFB/RBBB) will not be excluded), or
 - Congestive heart failure (CHF) of New York Heart Association (NYHA) Class \geq 3, or
 - Myocardial infarction (MI) within 3 months prior to Cycle 1 Day 1. Ejection fraction (EF) < 40% at screening.
 - Uncontrolled hypertension.
6. Uncontrolled active infection requiring parenteral antibiotics, antivirals, or antifungals within one week prior to first dose.
7. Known active hepatitis A, B or C.
8. Known HIV infection or HIV seropositivity.
9. Prior malignancies except treated cervical carcinoma in situ. Cancer treated with curative intent > 5 years before study enrollment and without evidence of recurrence will be allowed. Cancer treated with curative intent < 5 years previously will not be allowed unless approved by the medical monitor.
10. Any GI dysfunction that prevents the patient from swallowing tablets, or interferes with absorption of study treatment.
11. A serious psychiatric or medical condition that, in the opinion of the investigator, could interfere with treatment.

An Open Label Continuation Study of the Oral AKT Inhibitor GSK2110183 in Subjects with Hematologic or Solid Tumor Malignancy (PROTOCOL No. PKB115131 [Rollover])

Inclusion Criteria

A subject will be eligible for inclusion in this study only if all of the following criteria apply:

1. **Is currently participating in a GSK2110183 study (monotherapy or in combination with an approved anti-cancer agent) sponsored by GSK or by another research organization working on behalf of GSK.**
2. Currently benefitting from continued treatment and have an acceptable safety profile with GSK2110183 as determined by the investigator following previous treatment with GSK2110183 either as monotherapy or as part of a combination treatment regimen.
3. Continued ability to swallow and retain orally administered study treatment(s) and does not have any clinically significant GI abnormalities that may alter absorption such as malabsorption syndrome or major resection of the stomach or bowels.
4. Has adequate organ function:
 - Absolute neutrophil count (ANC) \geq 1.0 x 10⁹/L
 - Hemoglobin \geq 8.0 g/dL
 - Platelets \geq 50 x 10⁹/L

- PT/INR and PTT $\leq 1.5 \times$ ULN
- Total bilirubin $\leq 1.5 \times$ ULN (isolated bilirubin $> 1.5 \times$ ULN is acceptable if bilirubin is fractionated and direct bilirubin $< 35\%$)
- AST and ALT $\leq 3 \times$ ULN. If liver involvement is present and ALT and AST levels are $> 3 \times$ UNL and $< 5 \times$ ULN, enrollment into PKB115131 can occur as long as there is no concurrent bilirubin or INR elevation
- Serum creatinine OR Calculated creatinine clearance \leq ULN ≥ 30 mL/min
- Ejection Fraction (LVEF) $\geq 50\%$ by TTE or MUGA

Exclusion criteria:

A subject will not be eligible for inclusion in this study if any of the following criteria apply:

1. Permanent discontinuation of GSK2110183 in the parent study due to toxicity or disease progression.
2. Concomitant use of any type of anti-cancer treatment other than studied in the parent protocol.
3. Local access to commercially available GSK2110183.
4. Current use of a prohibitive medication(s) as listed in Section 7.2 of the protocol
5. Current use of anticoagulants is only allowed if PTT/INR values fulfill entry criteria.
6. Any unresolved toxicity $>$ Grade 2, except for alopecia, (National Cancer Institute-Common Toxicity Criteria for Adverse Events [NCI-CTCAE], version 4.0) from parent study treatment at the time of transition to this study.
7. History of HIV infection.
8. Peripheral neuropathy Gr >1
9. History of hepatitis B or C infection (subjects with evidence of cleared hepatitis B are permitted).
10. Evidence of severe or uncontrolled systemic diseases (e.g., unstable, or uncompensated respiratory, hepatic, renal, metabolic or cardiac disease).
11. QTcF interval $>$ 500 msec at the time of transition to this study.
12. Other clinically significant ECG abnormalities including 2nd degree (Type II) or 3rd degree atrioventricular (AV) block.
13. Evidence of current Class II, III, or IV heart failure as defined by the New York Heart Association [NYHA, 1994] functional classification system at the time of transition to this study.
14. Symptomatic or untreated leptomeningeal, CNS or brain metastases or spinal cord compression at the time of transition to this study.

NOTE: Subjects are not permitted to receive enzyme-inducing anti-epileptic drugs (EIAEDs). Continued stability of brain metastases must be confirmed with imaging.

AMYLOIDOSIS TRIALS:

A Phase 3, Randomized, Controlled, Open-label, Multicenter, Safety and Efficacy Study of Dexamethasone Plus MLN9708 or Physician's Choice of Treatment Administered to Patients with Relapsed or Refractory Systemic Light Chain (AL) Amyloidosis (Protocol C16011)

Inclusion Criteria:

1. Male or female patients 18 years or older.
2. Biopsy-proven diagnosis of AL amyloidosis according to the following standard criteria:
 - Histochemical diagnosis of amyloidosis, as based on tissue specimens with Congo red staining with exhibition of an apple-green birefringence
 - If clinical and laboratory parameters insufficient to establish AL amyloidosis or in cases of doubt, amyloid typing may be necessary
3. Measurable disease as defined by serum differential free light chain concentration (dFLC, difference between amyloid forming [involved] and nonamyloid forming [uninvolved] free light chain [FLC]) ≥ 50 mg/L.
4. Objective, measurable major (cardiac or renal) organ amyloid involvement as defined as follows (amyloid involvement of at least 1 required):
 - Cardiac involvement is defined as the presence of a mean left ventricular wall thickness on echocardiogram greater than 12 mm in the absence of a history of hypertension or valvular heart disease, or in the presence of unexplained low voltage (< 0.5 mV) on the electrocardiogram
 - Renal involvement is defined as proteinuria (predominantly albumin) > 0.5 g/day in a 24- hour urine collection
 - Note: Amyloid involvement of other organ systems is allowed, but not required.
5. Must be relapsed or refractory after 1 or 2 prior therapies.

For this protocol, relapsed is defined as PD documented more than 60 days after last dose; refractory is defined as documented absence of hematologic response or hematologic progression on or within 60 days after last dose of prior therapy.

- Patient may not be refractory to proteasome inhibitor therapy
 - Given that the physician may select from an offered list of regimens to treat a specific patient, the patient may be refractory to an agent/s listed within the list of offered treatment choices
 - Must have recovered (ie, \leq Grade 1 toxicity or patient's baseline status) from the reversible effects of prior therapy
 - If a patient has received a transplant as his/her first-line therapy, he/she must be at least 3 months posttransplantation and recovered from the side effects of the stem cell transplant
6. Patient must meet criteria for 1 of the following AL Amyloidosis Risk Stages (as defined by NT-proBNP cut off of < 332 pg/mL and troponin T cut-off of 0.035 ng/mL as thresholds):
 - Stage 1: both NT-proBNP and troponin T under threshold
 - Stage 2: either NT-proBNP or troponin T [but not both] over threshold;
 - Stage 3: both NT-proBNP and troponin T over threshold (but NT-proBNP < 8000 pg/mL)
 7. ECOG Performance Status ≤ 2
 8. Clinical laboratory values:
 - Absolute neutrophil count $\geq 1000/\mu\text{L}$
 - Platelet count $\geq 75,000/\mu\text{L}$
 - Total bilirubin $\leq 1.5 \times \text{ULN}$
 - Alkaline phosphatase $\leq 5 \times \text{ULN}$,
 - ALT or AST $\leq 3 \times \text{ULN}$

- Calculated creatinine clearance ≥ 30 mL/min
9. Female patients who:
- If they are of childbearing potential, agree to practice 2 effective methods of contraception, at the same time, from the time of signing the informed consent through 30 days after the last dose of study treatment, AND
 - Must also adhere to the guidelines of any treatment-specific pregnancy prevention program, if applicable, OR
 - Agree to completely abstain from heterosexual intercourse
- Male patients, even if surgically sterilized (ie, status post vasectomy), who:
- Agree to practice effective barrier contraception during the entire study treatment period and through 4 months after the last dose of study drug, AND
 - Must also adhere to the guidelines of any treatment-specific pregnancy prevention program, if applicable, OR
 - Agree to completely abstain from heterosexual intercourse
10. Voluntary written consent must be given before performance of any study-related procedure not part of standard medical care with the understanding that consent may be withdrawn by the patient at any time without prejudice to future medical care.

Exclusion Criteria

1. Amyloidosis due to mutations of the transthyretin gene or presence of other non-AL amyloidosis.
2. Female patients who are lactating, breastfeeding, or pregnant.
3. Medically documented cardiac syncope, uncompensated NYHA Class 3 or 4 congestive heart failure (Section 15.6), myocardial infarction within the previous 6 months, unstable angina pectoris, clinically significant repetitive ventricular arrhythmias despite antiarrhythmic treatment, or severe orthostatic hypotension or clinically important autonomic disease.
4. Clinically overt multiple myeloma, including monoclonal BM plasma cells $\geq 10\%$ to $\geq 30\%$, and at least 1 of the following:
 - Bone lesions
 - Hypercalcemia, defined as a calcium of > 11 g/dL
5. Inability to swallow oral medication, inability or unwillingness to comply with the drug administration requirements or GI procedure that could interfere with the oral absorption or tolerance of treatment.
6. Requirement for other concomitant chemotherapy, immunotherapy, radiotherapy, or any ancillary therapy considered to be investigational or which would be considered as a treatment of AL amyloidosis. However, patients may be on chronic steroids (maximum dose 20 mg/day prednisone or equivalent [Section 15.7]) if they are being given for disorders other than amyloidosis (eg, adrenal insufficiency, rheumatoid arthritis, etc.).
7. Comorbid systemic illnesses or other severe concurrent disease which, in the judgment of the investigator, would make the patient inappropriate for entry into this study or interfere significantly with the proper assessment of safety and toxicity of the prescribed regimens.
8. Ongoing or active infection, known HIV positive, known to be hepatitis B surface antigen-positive or has known or suspected active hepatitis C infection.
9. Psychiatric illness/social situations that would limit compliance with study requirements.
10. Known allergy to boron, MLN9708, any of the study treatments, their analogues, or excipients.
11. Systemic treatment with strong inhibitors of CYP1A2 (fluvoxamine, enoxacin, ciprofloxacin), strong inhibitors of CYP3A (clarithromycin, telithromycin, itraconazole, Voriconazole, ketoconazole, nefazodone, posaconazole) or strong CYP3A inducers (rifampin, rifapentine, rifabutin, carbamazepine, phenytoin, phenobarbital), or use of Ginkgo biloba or St. John's wort within 14 days before the first dose of study treatment.