MagnetisMM-6: A Phase 3 Study of Elranatamab (PF-06863135) + Daratumumab + Lenalidomide Versus Daratumumab + Bortezomib + Lenalidomide + Dexamethasone in Transplant-Ineligible Participants With Newly Diagnosed Multiple Myeloma

Estado	**Tipo de Participantes**	**Rangos de Edad**
Interrumpido	No aportado	Mayores de 64 , Adultos (18-64 años)
Género	**Fases**	**Participantes esperados**
Ambos	Fase III	616
Resultados	**Bajo nivel intervención**	**Enfermedad rara**
Sin resultados	No	No
Cobertura geográfica	**Ámbitos del ensayo**	**Terapia avanzada**
Sin determinar	tratamiento, seguridad, eficacia, farmacocinética, farmacodinamica, farmacoeconómica	NO

Información

Identificador
2024-514139-50-00 (CTIS)

Cod. Protocolo
C1071006

Transicionado 2021-000803-20

Área terapéutica
Enfermedades [C] - Cáncer [C04]|

Enfermedad investigada
Multiple myeloma

Título Científico
C1071006 AN OPEN-LABEL, 2-ARM, MULTICENTER, RANDOMIZED PHASE 3 STUDY TO EVALUATE THE

EFFICACY AND SAFETY OF ELRANATAMAB (PF-06863135) + DARATUMUMAB + LENALIDOMIDE VERSUS DARATUMUMAB + BORTEZOMIB + LENALIDOMIDE + DEXAMETHASONE IN TRANSPLANT-INELIGIBLE PARTICIPANTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

Justificación

No aportado

Objetivo Principal

Part 1
To assess dose limiting toxicities DLTs of EDR to select an RP3D for the combination to be used in Part 2 of this study.
Part 2
To compare the efficacy of Arm A (EDR) vs Arm B (D-VRd) as measured by MRD status and PFS

Variables de Evaluación Primaria

Part 1: DLTs during the DLT observation period:For all dose levels (DLs) except DL F: From the first priming dose of elranatamab in the 2 Step-up Priming Dose Period until 28 days (± visit windows) from the first administration of the EDR combination.
For DL F: 28 days (± visit windows) from the day of the first full dose of elranatamab (76 mg) in combination with daratumumab (D) and lenalidomide (R.).
Part 2: MRD negative CR rate (central lab) at 12 months after randomization per IMWG as assessed via next generation sequencing (NGS)
PFS per IMWG

Momentos temporales de evaluación primaria

No aportado

Objetivo Secundario

Part 1 To evaluate the overall safety profile of EDR or ER to select an RP3D for the combination to be used in Part 2 of this study.
To evaluate the efficacy of EDR or ER to select an RP3D for the combination to be used in Part 2 of this study.
To evaluate the PK of elranatamab when used in the EDR or ER combinations
To evaluate the immunogenicity of elranatamab when used in the EDR or ER combinations (Part 1 only).
To evaluate the PK of daratumumab and lenalidomide when used in EDR or ER combinations
Part 2. To compare the efficacy of Arm A (EDR) vs Arm B (D-VRd) as measured by OS

Variables de Evaluación Secundaria

Part 1: Adverse events (AEs) as characterized by type, frequency, severity (as graded by National Cancer Institute Common Terminology Criteria for Adverse Events [NCI CTCAE] v5.0), timing, seriousness, and relationship to study treatment. Severity of cytokine release syndrome (CRS) and immune effector cell-associated neurotoxicity syndrome (ICANS) will be graded according to American Society for Transplantation and Cellular Therapy (ASTCT) criteria.
Laboratory abnormalities as characterized by type, frequency, severity (as graded by NCI CTCAE v5.0), and timing.
Objective response rate (ORR) and complete response rate (CRR,), per International Myeloma Working Group (IMWG) response criteria as determined by investigator.
Time to event endpoints: time to response (TTR,), duration of response (DOR,), duration of complete response (DOCR) and progression-free survival (PFS) per IMWG response criteria as determined by investigator, and overall survival (OS;);
Minimal residual disease (MRD) negative CR rate (central laboratory) per IMWG sequencing criteria.
Predose and post dose concentrations of elranatamab against elranatamab
Anti-drug antibodies (ADAs) and neutralizing antibodies (NAbs) against elranatamab
Predose concentrations of daratumumab and lenalidomide
Part 2 Key Secondary: OS
Secondary: Overall MRD negative CR rate per IMWG
Duration of MRD negative CR per IMWG
PFS and progression-free survival on next line of therapy (PFS2) by investigator per IMWG
ORR, CRR, TTR, DOR, and DOCR per IMWG
AEs as characterized by type, frequency, severity (as graded by NCI CTCAE v5.0), timing, seriousness, and relationship to study treatment. The severity of CRS and ICANS will be graded according to ASTCT criteria (Lee, 2019).
Laboratory abnormalities as characterized by type, frequency, severity (as graded by NCI CTCAE v5.0), and timing.
Predose and post dose concentrations of elranatamab.
ADAs and NAbs against elranatamab.
EORTC QLQ-C30 and MY20European Organisation for Research and Treatment of Cancer (EORTC) Core Quality of Life Questionnaire (QLQC30) and Myeloma Module 20 (MY20).

Momentos temporales de evaluación secundaria

No aportado

Criterios de Inclusión

Participants must meet the following inclusion criteria to be eligible for enrollment into the study. Criteria are for both Part 1 and Part 2 unless otherwise specified: Age and Sex:Participant's age 18 years (or the minimum country specific age of consent if >18) at Visit 1 (Screening). Part 2 only: Participant has NDMM and is transplant-ineligible. In Part 2, transplant-ineligible is defined as: Participants not considered candidates for high-dose chemotherapy and ASCT due to age or Participants with important comorbidities likely to have a negative impact on tolerability of high dose chemotherapy and ASCT. Eastern Cooperative Oncology Group (ECOG) performance status 2. Adequate hepatic, renal, and bone marrow (BM) function (absolute neutrophil count [ANC], platelet count, hemoglobin). Corrected serum calcium 14 mg/dL (3.5 mmol/L), or free ionized calcium 6.5 mg/dL (1.6 mmol/L). Resolved acute effects of any prior therapy to baseline severity or CTCAE Grade 1. Type of Participant and Disease Characteristics: Diagnosis of multiple myeloma (MM) as defined according to IMWG criteria. Measurable disease based on IMWG criteria as defined by at least 1 of the following (as assessed by the central laboratory for Part 2) Serum M-protein 0.5 g/dL (Part 1) and 1g/dL (Part 2) Involved free light chain (FLC) 10 mg/dL (100 mg/L) AND abnormal serum immunoglobulin kappa to lambda FLC ratio (<0.26 or >1.65). Part 1 only: Participant with NDMM or RRMM. NDMM participant must be transplant-ineligible. In Part 1 transplant-ineligible is defined by age 65 years or by age <65 years with comorbidities impacting the possibility of transplant. Participants with RRMM must have received 1-2 prior lines of MM therapy including at least one immunomodulatory drug (IMiD) and one proteasome inhibitor (PI).

Criterios de Exclusión

Medical Conditions: Smoldering MM. Any other active malignancy within 3 years prior to enrollment, except for adequately treated basal cell or squamous cell skin cancer, or carcinoma in situ, or Stage 0/1 with minimal risk of recurrence per investigator. Participants with known or suspected hypersensitivity to the study interventions or any of their excipients. Participants with known or suspected central nervous system (CNS) or clinical signs of myelomatous meningeal involvement. Other surgical (including major surgery within 14 days prior to enrollment), medical or psychiatric conditions including recent (within the past year) or active suicidal ideation/behaviour or laboratory abnormality that may increase the risk of study participation or, in the investigator's judgment, make the participant inappropriate for the study. Active inflammatory gastrointestinal disease, chronic diarrhea, known diverticular disease or previous gastric resection or lap band surgery that may significantly alter the absorption of oral drugs. Gastroesophageal reflux disease under treatment with proton pump inhibitors is allowed (assuming no drug interaction potential). Monoclonal gammopathy of undetermined significance (MGUS). Plasma cell leukemia. Waldenström`s Macroglobulinemia. Systemic light chain amyloidosis. Polyneuropathy, Organomegaly, Endocrinopathy, Monoclonal gammopathy, and Skin abnormalities (POEMS) Syndrome. Impaired cardiovascular function or clinically significant cardiovascular diseases within 6 months prior to enrollment. Ongoing Grade 3 (Part 1)/Grade 2 (Part 2) or higher peripheral sensory or motor neuropathy, history of Guillain-Barré syndrome (GBS) or GBS variants, or history of any Grade >3 (Part 1)/Grade >2 (Part 2) peripheral motor polyneuropathy. Active, uncontrolled bacterial, fungal, or viral infection, including (but not limited to) coronavirus disease 2019 (COVID-19)/severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), hepatitis B virus (HBV), hepatitis C virus (HCV), and known human immunodeficiency virus (HIV) or acquired immunodeficiency syndrome (AIDS)-related illness. Active infections must be resolved at least 21 days prior to enrollment. Participants treated with systemic therapeutic anti-infective agents within 28 days prior to enrollment are not eligible. Prophylactic use of systemic anti-infective agents is permitted.

Calendario

(Última actualización: 04/08/2026)

Autorización	Inicio de Ensayo	Inclusión Primer Paciente	Interrumpido	Reiniciado
22/12/2022	01/03/2023	17/05/2023	29/10/2025	No aportado

Fin de reclutamiento	Fin prematuro (España)	Fin prematuro (Global)	Fin del ensayo en España	Fin del ensayo global
No aportado	No aportado	No aportado	No aportado	No aportado

Promotor

Pfizer Inc. United States

66 Hudson Boulevard East

Contact Person

Clinical Medical Lead

+18007181021

ClinicalTrials.gov_Inquiries@pfizer.com

Monetary support: N/A

Tipo de promotor: Comercial

Ceim

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ceic.germanstrias@gencat.cat

934978956

Centros

Activo (09/03/2020)

Clinica Universidad De Navarra

Pamplona

NAVARRA

N/A

Activo (21/11/2025)

Complexo Hospitalario Universitario De Santiago

Santiago De Compostela

CORUÑA

N/A

Activo (03/03/2020)

Hospital Clinic De Barcelona

Barcelona

BARCELONA

N/A

Activo (03/05/2020)

Hospital Germans Trias I Pujol

Badalona

BARCELONA

Instituto Catalán de Oncología de Badalona

Activo (09/02/2024)

Hospital San Pedro De Alcantara

Caceres

CÁCERES

N/A

Activo (21/12/2023)

Hospital Universitari Mutua Terrassa

Terrassa

BARCELONA

N/A

Activo (01/03/2020)

Hospital Universitario De La Princesa

Madrid

MADRID

N/A

Activo (23/03/2020)

Hospital Universitario Dr Peset Aleixandre

Valencia

VALENCIA

N/A

Activo (10/06/2024)

Hospital Universitario Fundacion Jimenez Diaz

Madrid

MADRID

N/A

Activo (23/06/2026)

Hospital Universitario Marques De Valdecilla

Santander

CANTABRIA

N/A

Activo (05/06/2026)

Hospital Universitario Regional De Malaga

Malaga

MÁLAGA

N/A

Activo (15/06/2026)

Hospital Universitario Y Politecnico La Fe

Valencia

VALENCIA

Hematología y Hemoterapia

Activo (15/11/2023)

Institut Catala D´oncologia

L'hospitalet De Llobregat

BARCELONA

Haematology Department

No iniciado (--/--/----)

Institut Catala D´oncologia

Girona

GERONA

Hematología Clínica

Medicamentos

Zelvina 15 mg hard capsules
CAPSULE, HARD
ORAL USE

Código ATC: L04AX04 - LENALIDOMIDA
Principios Activos: N/A

Experimental

Privigen 100 mg/ml solution for infusion
SOLUTION FOR INFUSION
IV INFUSION

Código ATC: J06BA02 - INMUNOGLOBULINAS HUMANAS NORMALES PARA ADM. INTRAVASCULAR
Principios Activos: N/A

Experimental

ELRANATAMAB
SOLUTION FOR INJECTION
SUBCUTANEOUS USE

-
Principios Activos: N/A

Experimental

Zelvina 2.5 mg hard capsules
CAPSULE, HARD
ORAL USE

Código ATC: L04AX04 - LENALIDOMIDA
Principios Activos: N/A

Experimental

Zelvina 5 mg hard capsules
CAPSULE, HARD
ORAL USE

Código ATC: L04AX04 - LENALIDOMIDA
Principios Activos: N/A

Experimental

Dexamethasone 0.5mg Tablets
TABLET
ORAL USE

Código ATC: H02AB02 - DEXAMETASONA
Principios Activos: N/A

Experimental

Zelvina 25 mg hard capsules
CAPSULE, HARD
ORAL USE

Código ATC: L04AX04 - LENALIDOMIDA
Principios Activos: N/A

Experimental

Nodexon 20 mg tabletid
TABLET
ORAL USE

Código ATC: H02AB02 - DEXAMETASONA
Principios Activos: N/A

Experimental

Nodexon 20 mg tabletes
TABLET
ORAL USE

Código ATC: H02AB02 - DEXAMETASONA
Principios Activos: N/A

Experimental

Nodexon 20 mg tablets
TABLET
ORAL USE

Código ATC: H02AB02 - DEXAMETASONA
Principios Activos: N/A

Experimental

Nodexon 20 mg tablets
TABLET
ORAL USE

Código ATC: H02AB02 - DEXAMETASONA
Principios Activos: N/A

Experimental

DARZALEX 1800 mg solution for injection
SOLUTION FOR INJECTION
SUBCUTANEOUS USE

Código ATC: L01FC01 - DARATUMUMAB
Principios Activos: N/A

Huérfano
Experimental

Dexamethasone 4 mg tablets
TABLET
ORAL USE

Código ATC: H02AB02 - DEXAMETASONA
Principios Activos: N/A

Experimental

VELCADE 3.5 mg powder for solution for injection
SOLUTION FOR INJECTION
SUBCUTANEOUS

Código ATC: L01XG01 - BORTEZOMIB
Principios Activos: N/A

Experimental

Zelvina 10 mg hard capsules
CAPSULE, HARD
ORAL USE

Código ATC: L04AX04 - LENALIDOMIDA
Principios Activos: N/A

Experimental

Sin resultados

MagnetisMM-6: A Phase 3 Study of Elranatamab (PF-06863135) + Daratumumab + Lenalidomide Versus Daratumumab + Bortezomib + Lenalidomide + Dexamethasone in Transplant-Ineligible Participants With Newly Diagnosed Multiple Myeloma

State	Type of participants	Age Ranges
Halted	Not provided	Older than 64 , Adults (18-64 years)
Gender	**Phases**	**Expected Participants**
Both	Phase III	616
Results	**Low level of intervention**	**Rare disease**
No results	No	No
Geographic coverage	**Areas of the study**	**Advanced therapy**
Undetermined	treatment, safety, effectiveness, pharmacokinetics, pharmacodynamics, pharmacoeconomic	NO

Information

Identifier
2024-514139-50-00 (CTIS)

Protocol Code
C1071006

Transitioned 2021-000803-20

Therapeutic area
Diseases [C] - Cancer [C04]|

Investigated Disease
Multiple myeloma

Scientific Title
C1071006 AN OPEN-LABEL, 2-ARM, MULTICENTER, RANDOMIZED PHASE 3 STUDY TO EVALUATE THE

EFFICACY AND SAFETY OF ELRANATAMAB (PF-06863135) + DARATUMUMAB + LENALIDOMIDE VERSUS DARATUMUMAB + BORTEZOMIB + LENALIDOMIDE + DEXAMETHASONE IN TRANSPLANT-INELIGIBLE PARTICIPANTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

Rationale

Not provided

Main Objective

Part 1
To assess dose limiting toxicities DLTs of EDR to select an RP3D for the combination to be used in Part 2 of this study.
Part 2
To compare the efficacy of Arm A (EDR) vs Arm B (D-VRd) as measured by MRD status and PFS

Primary Endpoints

Part 1: DLTs during the DLT observation period:For all dose levels (DLs) except DL F: From the first priming dose of elranatamab in the 2 Step-up Priming Dose Period until 28 days (± visit windows) from the first administration of the EDR combination.
For DL F: 28 days (± visit windows) from the day of the first full dose of elranatamab (76 mg) in combination with daratumumab (D) and lenalidomide (R.).
Part 2: MRD negative CR rate (central lab) at 12 months after randomization per IMWG as assessed via next generation sequencing (NGS)
PFS per IMWG

Temporary moments of secondary assessment

Not provided

Secondary Objective

Part 1 To evaluate the overall safety profile of EDR or ER to select an RP3D for the combination to be used in Part 2 of this study.
To evaluate the efficacy of EDR or ER to select an RP3D for the combination to be used in Part 2 of this study.
To evaluate the PK of elranatamab when used in the EDR or ER combinations
To evaluate the immunogenicity of elranatamab when used in the EDR or ER combinations (Part 1 only).
To evaluate the PK of daratumumab and lenalidomide when used in EDR or ER combinations
Part 2. To compare the efficacy of Arm A (EDR) vs Arm B (D-VRd) as measured by OS

Secondary Endpoints

Part 1: Adverse events (AEs) as characterized by type, frequency, severity (as graded by National Cancer Institute Common Terminology Criteria for Adverse Events [NCI CTCAE] v5.0), timing, seriousness, and relationship to study treatment. Severity of cytokine release syndrome (CRS) and immune effector cell-associated neurotoxicity syndrome (ICANS) will be graded according to American Society for Transplantation and Cellular Therapy (ASTCT) criteria.
Laboratory abnormalities as characterized by type, frequency, severity (as graded by NCI CTCAE v5.0), and timing.
Objective response rate (ORR) and complete response rate (CRR,), per International Myeloma Working Group (IMWG) response criteria as determined by investigator.
Time to event endpoints: time to response (TTR,), duration of response (DOR,), duration of complete response (DOCR) and progression-free survival (PFS) per IMWG response criteria as determined by investigator, and overall survival (OS;);
Minimal residual disease (MRD) negative CR rate (central laboratory) per IMWG sequencing criteria.
Predose and post dose concentrations of elranatamab against elranatamab
Anti-drug antibodies (ADAs) and neutralizing antibodies (NAbs) against elranatamab
Predose concentrations of daratumumab and lenalidomide
Part 2 Key Secondary: OS
Secondary: Overall MRD negative CR rate per IMWG
Duration of MRD negative CR per IMWG
PFS and progression-free survival on next line of therapy (PFS2) by investigator per IMWG
ORR, CRR, TTR, DOR, and DOCR per IMWG
AEs as characterized by type, frequency, severity (as graded by NCI CTCAE v5.0), timing, seriousness, and relationship to study treatment. The severity of CRS and ICANS will be graded according to ASTCT criteria (Lee, 2019).
Laboratory abnormalities as characterized by type, frequency, severity (as graded by NCI CTCAE v5.0), and timing.
Predose and post dose concentrations of elranatamab.
ADAs and NAbs against elranatamab.
EORTC QLQ-C30 and MY20European Organisation for Research and Treatment of Cancer (EORTC) Core Quality of Life Questionnaire (QLQC30) and Myeloma Module 20 (MY20).

Temporary moments of secondary assessment

Not provided

Inclusion criteria

Participants must meet the following inclusion criteria to be eligible for enrollment into the study. Criteria are for both Part 1 and Part 2 unless otherwise specified: Age and Sex:Participant's age 18 years (or the minimum country specific age of consent if >18) at Visit 1 (Screening). Part 2 only: Participant has NDMM and is transplant-ineligible. In Part 2, transplant-ineligible is defined as: Participants not considered candidates for high-dose chemotherapy and ASCT due to age or Participants with important comorbidities likely to have a negative impact on tolerability of high dose chemotherapy and ASCT. Eastern Cooperative Oncology Group (ECOG) performance status 2. Adequate hepatic, renal, and bone marrow (BM) function (absolute neutrophil count [ANC], platelet count, hemoglobin). Corrected serum calcium 14 mg/dL (3.5 mmol/L), or free ionized calcium 6.5 mg/dL (1.6 mmol/L). Resolved acute effects of any prior therapy to baseline severity or CTCAE Grade 1. Type of Participant and Disease Characteristics: Diagnosis of multiple myeloma (MM) as defined according to IMWG criteria. Measurable disease based on IMWG criteria as defined by at least 1 of the following (as assessed by the central laboratory for Part 2) Serum M-protein 0.5 g/dL (Part 1) and 1g/dL (Part 2) Involved free light chain (FLC) 10 mg/dL (100 mg/L) AND abnormal serum immunoglobulin kappa to lambda FLC ratio (<0.26 or >1.65). Part 1 only: Participant with NDMM or RRMM. NDMM participant must be transplant-ineligible. In Part 1 transplant-ineligible is defined by age 65 years or by age <65 years with comorbidities impacting the possibility of transplant. Participants with RRMM must have received 1-2 prior lines of MM therapy including at least one immunomodulatory drug (IMiD) and one proteasome inhibitor (PI).

Exclusion criteria

Medical Conditions: Smoldering MM. Any other active malignancy within 3 years prior to enrollment, except for adequately treated basal cell or squamous cell skin cancer, or carcinoma in situ, or Stage 0/1 with minimal risk of recurrence per investigator. Participants with known or suspected hypersensitivity to the study interventions or any of their excipients. Participants with known or suspected central nervous system (CNS) or clinical signs of myelomatous meningeal involvement. Other surgical (including major surgery within 14 days prior to enrollment), medical or psychiatric conditions including recent (within the past year) or active suicidal ideation/behaviour or laboratory abnormality that may increase the risk of study participation or, in the investigator's judgment, make the participant inappropriate for the study. Active inflammatory gastrointestinal disease, chronic diarrhea, known diverticular disease or previous gastric resection or lap band surgery that may significantly alter the absorption of oral drugs. Gastroesophageal reflux disease under treatment with proton pump inhibitors is allowed (assuming no drug interaction potential). Monoclonal gammopathy of undetermined significance (MGUS). Plasma cell leukemia. Waldenström`s Macroglobulinemia. Systemic light chain amyloidosis. Polyneuropathy, Organomegaly, Endocrinopathy, Monoclonal gammopathy, and Skin abnormalities (POEMS) Syndrome. Impaired cardiovascular function or clinically significant cardiovascular diseases within 6 months prior to enrollment. Ongoing Grade 3 (Part 1)/Grade 2 (Part 2) or higher peripheral sensory or motor neuropathy, history of Guillain-Barré syndrome (GBS) or GBS variants, or history of any Grade >3 (Part 1)/Grade >2 (Part 2) peripheral motor polyneuropathy. Active, uncontrolled bacterial, fungal, or viral infection, including (but not limited to) coronavirus disease 2019 (COVID-19)/severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), hepatitis B virus (HBV), hepatitis C virus (HCV), and known human immunodeficiency virus (HIV) or acquired immunodeficiency syndrome (AIDS)-related illness. Active infections must be resolved at least 21 days prior to enrollment. Participants treated with systemic therapeutic anti-infective agents within 28 days prior to enrollment are not eligible. Prophylactic use of systemic anti-infective agents is permitted.

Calendar

(Last Update: 04/08/2026)

Authorization	Start of Trial	First patient inclusion	Halted	Restarted
22/12/2022	01/03/2023	17/05/2023	29/10/2025	Not aported

End of recruitment	Premature end (Spain)	Premature End (Global)	Trial end (Spain)	Trial end (Global)
Not aported	Not aported	Not aported	Not aported	Not aported

Sponsor

Pfizer Inc. United States

66 Hudson Boulevard East

Contact Person

Clinical Medical Lead

+18007181021

ClinicalTrials.gov_Inquiries@pfizer.com

Monetary support: N/A

Sponsor type: Commercial

Ceim

CEIm Hospital Universitari Germans Trias i Pujol

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ceic.germanstrias@gencat.cat

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Sites

Active (09/03/2020)

Clinica Universidad De Navarra

Pamplona

NAVARRA

N/A

Active (21/11/2025)

Complexo Hospitalario Universitario De Santiago

Santiago De Compostela

CORUÑA

N/A

Active (03/03/2020)

Hospital Clinic De Barcelona

Barcelona

BARCELONA

N/A

Active (03/05/2020)

Hospital Germans Trias I Pujol

Badalona

BARCELONA

Instituto Catalán de Oncología de Badalona

Active (09/02/2024)

Hospital San Pedro De Alcantara

Caceres

CÁCERES

N/A

Active (21/12/2023)

Hospital Universitari Mutua Terrassa

Terrassa

BARCELONA

N/A

Active (01/03/2020)

Hospital Universitario De La Princesa

Madrid

MADRID

N/A

Active (23/03/2020)

Hospital Universitario Dr Peset Aleixandre

Valencia

VALENCIA

N/A

Active (10/06/2024)

Hospital Universitario Fundacion Jimenez Diaz

Madrid

MADRID

N/A

Active (23/06/2026)

Hospital Universitario Marques De Valdecilla

Santander

CANTABRIA

N/A

Active (05/06/2026)

Hospital Universitario Regional De Malaga

Malaga

MÁLAGA

N/A

Active (15/06/2026)

Hospital Universitario Y Politecnico La Fe

Valencia

VALENCIA

Hematología y Hemoterapia

Active (15/11/2023)

Institut Catala D´oncologia

L'hospitalet De Llobregat

BARCELONA

Haematology Department

not initialized (--/--/----)

Institut Catala D´oncologia

Girona

GERONA

Hematología Clínica

Medication

Zelvina 15 mg hard capsules
CAPSULE, HARD
ORAL USE

ATC code: L04AX04 - LENALIDOMIDA
Active Principles: N/A

Experimental

Privigen 100 mg/ml solution for infusion
SOLUTION FOR INFUSION
IV INFUSION

ATC code: J06BA02 - INMUNOGLOBULINAS HUMANAS NORMALES PARA ADM. INTRAVASCULAR
Active Principles: N/A

Experimental

ELRANATAMAB
SOLUTION FOR INJECTION
SUBCUTANEOUS USE

-
Active Principles: N/A

Experimental

Zelvina 2.5 mg hard capsules
CAPSULE, HARD
ORAL USE

ATC code: L04AX04 - LENALIDOMIDA
Active Principles: N/A

Experimental

Zelvina 5 mg hard capsules
CAPSULE, HARD
ORAL USE

ATC code: L04AX04 - LENALIDOMIDA
Active Principles: N/A

Experimental

Dexamethasone 0.5mg Tablets
TABLET
ORAL USE

ATC code: H02AB02 - DEXAMETASONA
Active Principles: N/A

Experimental

Zelvina 25 mg hard capsules
CAPSULE, HARD
ORAL USE

ATC code: L04AX04 - LENALIDOMIDA
Active Principles: N/A

Experimental

Nodexon 20 mg tabletid
TABLET
ORAL USE

ATC code: H02AB02 - DEXAMETASONA
Active Principles: N/A

Experimental

Nodexon 20 mg tabletes
TABLET
ORAL USE

ATC code: H02AB02 - DEXAMETASONA
Active Principles: N/A

Experimental

Nodexon 20 mg tablets
TABLET
ORAL USE

ATC code: H02AB02 - DEXAMETASONA
Active Principles: N/A

Experimental

Nodexon 20 mg tablets
TABLET
ORAL USE

ATC code: H02AB02 - DEXAMETASONA
Active Principles: N/A

Experimental

DARZALEX 1800 mg solution for injection
SOLUTION FOR INJECTION
SUBCUTANEOUS USE

ATC code: L01FC01 - DARATUMUMAB
Active Principles: N/A

Orphan
Experimental

Dexamethasone 4 mg tablets
TABLET
ORAL USE

ATC code: H02AB02 - DEXAMETASONA
Active Principles: N/A

Experimental

VELCADE 3.5 mg powder for solution for injection
SOLUTION FOR INJECTION
SUBCUTANEOUS

ATC code: L01XG01 - BORTEZOMIB
Active Principles: N/A

Experimental

Zelvina 10 mg hard capsules
CAPSULE, HARD
ORAL USE

ATC code: L04AX04 - LENALIDOMIDA
Active Principles: N/A

Experimental

No results