









SMML newsletter

September 2024
Based on data from May-August 2024

Monthly updates around multiple myeloma

















Objectives





The goal of this newsletter is to give an overview of what is happening around multiple myeloma in terms of medical developments. This is achieved by monitoring news articles, PR releases and scientific reports on one hand. Conversations from HCPs, digital opinion leaders and other influencers attending congresses, offering their opinion on treatments and commenting on new developments on the other hand.





Who will benefit from this newsletter?

Any professionals interested in keeping up with developments around multiple myeloma:

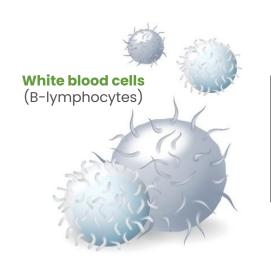
- Marketers
- Business Analysts
- But also HCPs who want to anticipate the future landscape of MM management



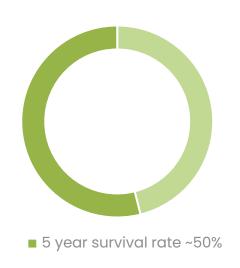
Introduction on Multiple Myeloma

What is Multiple Myeloma?











Read more on Multiple Myeloma:



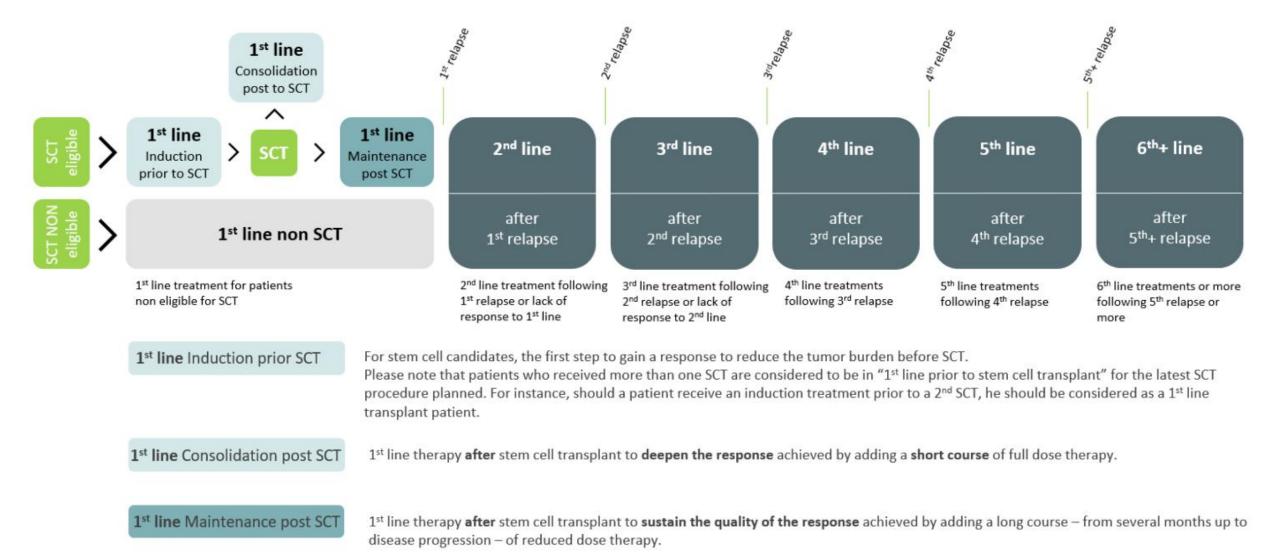
- → Professional version in English
- → Professional version in French



- → Consumer version in English
- → Consumer version in French

MM lines of therapy based on to the following definitions





Drugs approved by European Medicines Agency (EMA)

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+	•	•	•	
	•	•	•	
	•	•	•	

Product Nonograph Click here

Click here

Brand name	Active substance	Manufacturer	Class of therapy	EMA Date of authorisation	Line of treatment	M	
Revlimid [®]	lenalidomide	BMS	Immunomodulating agents (IMiDs)	14/06/2007	All lines		
Thalidomide® / Thalomid® (US)	thalidomide	BMS	Immunomodulating agents (IMiDs)	16/04/2008	All lines		
Imnovid® / Pomalyst® (US)	pomalidomide	BMS	Immunomodulating agents (IMiDs)	05/08/2013	2L+ (after Revlimid and Velcade)		
Pepaxti® / Pepaxto® (US)	melflufen	Oncopeptides AB	Peptide conjugated alkylator	17/08/2022	Triple class exposed / 3L+ / 4L+ / 5L+		

Click here Click here Click here 26/04/2004 1LSCT / 1LNSCT Velcade[®] bortezomib Janssen Proteasome inhibitors (PIs) Click here Kyprolis® carfilzomib 19/11/2015 2L+ / 3L+ **Click here**

21/11/2016

28/08/2015

20/05/2016

11/05/2016

30/05/2020

21/07/2022

09/08/2024 (FDA)

14/08/2024 (FDA)

25/08/2020

26/03/2021

04/12/2016

18/08/2021

25/05/2022

31/10/1991

20/03/2001

26/05/2010 - 13/07/2011

30/07/2009

2L+

All lines

2L+ / 3L+

2L+ / 3L+

Supportive care

Supportive care

2L+ Relapsed / Refractory

Proteasome inhibitors (PIs) Amgen ixazomib Takeda Proteasome inhibitors (PIs) panobinostat

Ninlaro[®]

Farydak[®]

Darzalex®

Empliciti®

Sarclisa[®]

Tecvayli[®]

Talvey[®]

Elrexfio[®]

Abecma®

Carvykti[®]

Aredia[®]

Zometa[®]

Mozobil®

Prolia® + Xgeva®

Blenrep® / Belamaf® (US)

Nexpovio[®] / Xpovio[®] (US)

Venclyxto® / Venclexta® (US)

daratumumab

elotuzumab

isatuximab

selinexor

venetoclax

pamidronate

zoledronate

denosumab

plerixafor

teclistamab-cqyv

talquetamab-tgvs

elranatamab-bcmm

idecabtagene vicleucel

ciltacabtagene autoleucel

belantamab mafodotin-blmf

Pfizer

GSK

AbbVie

Janssen

Novartis

Novartis

Amgen

Genzyme

BMS

Karyopharm Therapeutics

Novartis + (Secura Bio)

Janssen

Histone deacetylase inhibitors (HDACis) Monoclonal antibody against CD38 (Mabs) Monoclonal antibody against SLAMF7 (Mabs)

BMS + (AbbVie) Sanofi Monoclonal antibody against CD38 (Mabs) Bispecific B-cell maturation antigen (BCMA)-directed Janssen CD3 T-cell engager Janssen

Cell-based gene therapy (CAR-T)

Bisphosphonates for bone disease

Bisphosphonates for bone disease

Bisphosphonates for bone disease

Stem cell mobilazor

Bispecific antibody targeting GPRC5D receptor Bispecific B-cell maturation antigen (BCMA)-directed

CD3 T-cell engager Antibody-drug conjugates (BCMA)

Nuclear export inhibitor (SINE) B-cell lymphoma 2 (BCL-2)

Cell-based gene therapy (CAR-T)

2L+ (already used in CLL / AML treatments) Triple class exposed / 3L+ / 4L+ / 5L+ L3 / L4+ / L5+

4L+ (1Pi + 1 IMiDs + 1anti-CD38 Mabs)

4L+ (1Pi + 1 IMiDs + 1anti-CD38 Mabs)

4L+ (1Pi + 1 IMiDs + 1anti-CD38 Mabs)

2L+ (4L+ after P, IMiD and MAB (L5+ in Italy)

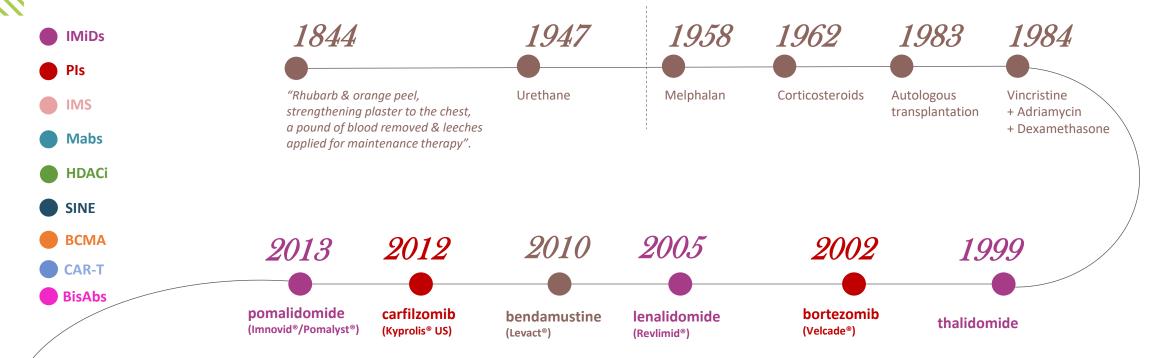
Penta-refractory (2xPI + 2xIMiDs + 1 Mab)

Supportive care / Long-term use (5+ years)

Click here Click here Click here Click here Supportive care / Long-term use (5+ years) **Click here** Prolia® / Xgeva®

MM treatment history timeline







SYNDICATED BY

MM studies at APLUSA







per wave per country per year

4,5K **PATIENTS CHARTS** per wave

18k **PATIENTS CHARTS** per year (in EU5)



Launching of a US pilot wave, Why?

- Needs from Pfizer and GSK for MM US data
- Back data for EU5 for 15 years
- MM US market changing fast
- Different needs in the US market for labs (patients' ethnicity treatments gap as an example)

→ Co-funding with GSK for Q4 2022

APLUSA's added value

- Quarterly + month to month data collection (our competitors → only quarterly data)
- Market adaptability: adjusting our questionnaires and updating the list of treatments by including market trends on a monthly basis → **new CAR-Ts + BisAbs**
- Patient's full treatment history
- Deliverables with Power BI DID



May-August 2024





CONTENTS

01.

News around clinical trials in phase II

- Phase II Relapse/Refractory:
 Bispecific Therapies
- Phase II Relapse/Refractory
 CAR-T

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News around clinical trial s in Phase III

- Phase III Line 1 Triplet Therapy
- Phase III Line 1 Quadruplet
 Therapy
- Phase III Relapse/Refractory
 Doublet Therapy
- Phase III Relapse/Refractory
 Triplet Therapy

Phase III relapse/refractory
 CART

03.

Themes of discussion: HCPs

- SOV Themes of discussion
- Focus on #ASCO2024
- Focus on #EHA2024
- Posts driving most engagement

04.

News articles overtime

- Volume of articles per volume of mentions M ay-June 2024
- Volume of articles per volume of mentions J uly-August 2024
- Articles per themes

05.

Market watch

- Market watch May
- Market watch June
- Market watch July
- Market watch August







SCOPE



The scope of the analysis is focused on mentions coming worldwide.*



There were a total of 120K mentions recorded during the listening period from May 1st, 2024, to June 30th, 2024.



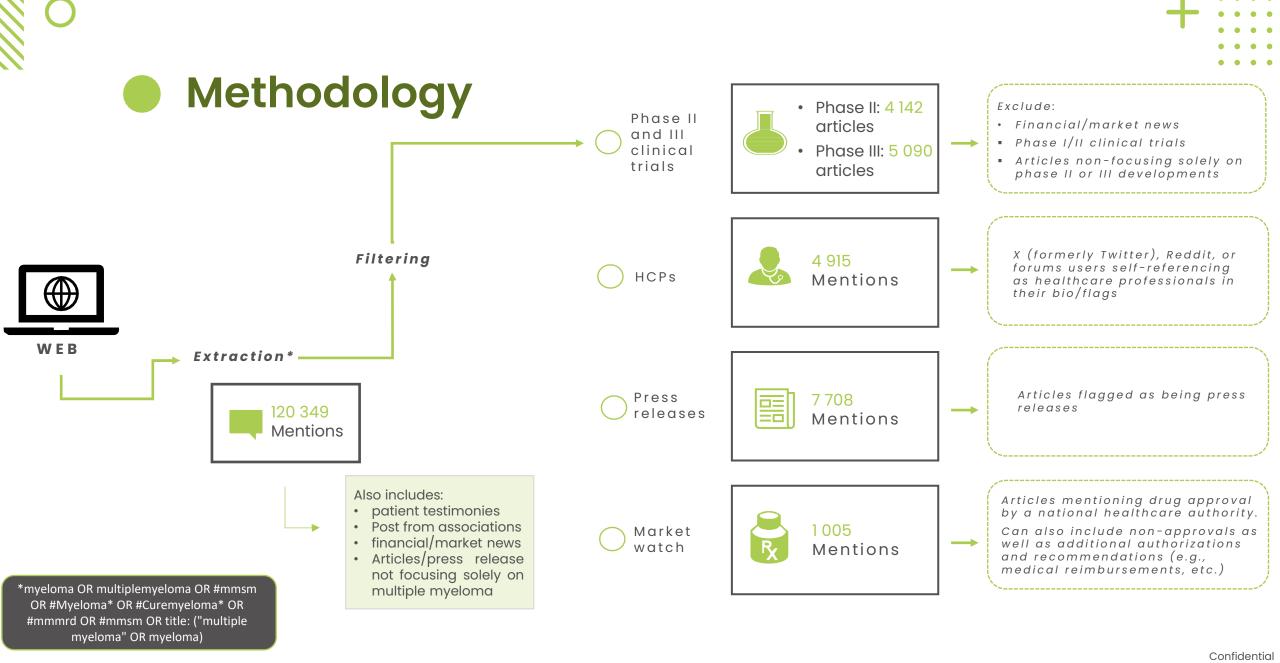
A majority of mentions came from News (71%), X (formerly Twitter) (23%), Instagram (3%), Reddit (2%), forum (1%).



A total of 13K unique authors were identified.

myeloma OR multiplemyeloma OR #mmsm OR #Myeloma OR #Curemyeloma* OR #mmmrd OR #mmsm OR title: ("multiple myeloma" OR myeloma)





SYNDICATED BY

Drugs in clinical trials (Phase II) monitored during the listening period

RR

RR

RR

RR

RR

RR

RR

1L

1L

1L

RR

RR

RR

RR

MajesTEC-1

RedirecTT-1

iMMagine-1

FUMANBA-1

NCT04720313

MagnetisMM-3

MagnetisMM-9

LINKER-MM1

NCT05123131

NCT02969837

NCT03590652

NCT04309981

NCT06105554

Cardamon

MASTER

REBUILD

FORTE

Kite, a Gilead company

Nexcella

Regeneron

Pfizer

Nanjing IASO Biotechnology

MonumenTAL-1



Sponsor: Mario Boccadoro, University of Turin, Italy

Sponsor: University of California, San Diego

Sponsor: Hellenic Society of Hematology

Sponsor: M.D. Anderson Cancer Center

Sponsor: University of Alabama at Birmingham

Sponsor: University College, London

Sponsor: Cancer Trials Ireland

Sponsor: University of Chicago

Sponsor: Sara V. Latorre

Diag	Drugs in clinical trials (Fridse ii) monitored during the listering period			
Sponsor	Trial name + link (ctrl + right click)	Line of treatment	Combination/Monotherapy being studied	Note
AbbVie	NCT03314181	RR	venetoclax, daratumumab and dexamethasone (With and Without Bortezomib)	
Bristol-Myers Squibb	ELOQUENT-3	RR	Elotuzumab, pomalidomide, dexamethasone	
CARsgen Therapeutics	LUMMICAR-2	RR	Monotherapy: zevorcabtagene autoleucel	
Celgene	NCT03989414	1L/RR	bortezomb, dexamethasone, daratumumab, elotuzumab, isatuximab, carfilzomib	
	NCT03989414	IL/RR	mezigdomide, bortezomib, dexamethasone, daratumumab, carfilzomib, elotuzumab, isatuximab	
	KarMMa-2	RR	Monotherapy (one arm with lenalidomide); BMTCTN1902: Monotherapy: Idecabtagene vicleucel	
	NCT03374085	RR	Monotherapy: mezigdomide or in combination with dexamethasone	•
Cellular Biomedicine Group	NCT05521802	RR	Monotherapy: C-CAR088	
GSK	DREAMM-2	RR	Monotherapy: blenrep	
Janssen	GRIFFIN	1L	daratumumab, lenalidomide, bortezomib, and dexamethasone (D-RVd)	
	CARTITUDE-1	RR	Monotherapy: ciltacabtagene autoleucel	
	CARTITUDE-2	RR	Monotherapy: ciltacabtagene autoleucel	

Monotherapy: teclistamab

Monotherapy: talquetamab

Monotherapy: NXC-201

Monotherapy: elranatamab

Monotherapy: daratumumab

teclistamab, talquetamab, daratumumab

Monotherapy: anitocabtagene autoleucel

Monotherapy: equecabtagene autoleucel

Monotherapy: elranatamab or in combination with dexamethasone

Monotherapy: Linvoseltamab (also known as REGN5458)

carfilzomib, lenalidomide, daratumumab, dexamethasone

isatuximab, bortezomib, lenalidomide, dexamethasone

elotuzumab, carfilzomib, lenalidomide, dexamethasone

daratumumab, ixazomib, pomalidomide, dexamethasone

Monotherapy: cesnicabtagene autoleucel (ARI0002h)

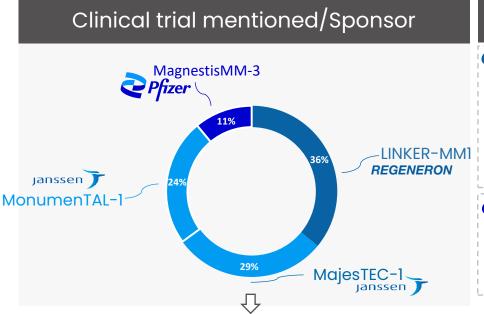
belumosudil mesylate with or without dexamethasone

dexamethasone, lenalidomide, daratumumab, carfilzomib

cyclophosphamide, dexamethasone, carfilzomib

Phase II Relapse/Refractory Bispecific Therapies: Updated Linvoseltamab Data Showcase Continued Deepening of Responses in Patients with Heavily Pre-Treated Multiple Myeloma





Headlines/Hot off the press

Updated Linvoseltamab Data Showcase Continued Deepening of Responses in Patients with Heavily Pre-Treated Multiple Myeloma Click here to read the full article

FDA declines to approve Regeneron's linvoseltamab amid manufacturing issues

Click here to read the full article

ELREXFIO™ Shows Median Overall Survival of More Than Two Years in People with Relapsed or **Refractory Multiple Myeloma**

Click here to read the full article

99166 Johnson & Johnson At ASCO Highlights Long Term Data From Multiple Myeloma Drug, Shows Sustained Deep, Durable Responses In **Pretreated Patients**

Click here to read the full article

TALVEY® (talquetamab-tgvs) demonstrates highly durable, longer-term responses in patients with relapsed or refractory multiple myeloma

Click here to read the full article

Janssen's bispecific antibody drug for multiple myeloma wins nod in Korea

Click here to read the full article

LINKER-MM1

Ctrl + click to access clinical trial:

MajesTEC-1

Ctrl + click to access clinical

MonumenTAL-1

Ctrl + click to access clinical trial:

talquetamab in participants with relapsed or refractory multiple myeloma at the recommended Phase 2 dose(s) (RP2Ds) (Part 3)."

(MajesTEC-1)"

Relapsed or Refractory

MagnetisMM-3

Ctrl + click to access clinical trial: NCT04649359

"Study Of Elranatamab (PF-06863135) Monotherapy in Participants With Multiple Myeloma Who Are Refractory to at Least One PI, One IMiD and One Anti-CD38 mAb"

"Phase 1/2 Study of REGN5458 in Patients With

"A Study of Teclistamab in Participants With

"The purpose of this study is to evaluate the efficacy of

Multiple Myeloma

Relapsed or Refractory Multiple Myeloma"

Combinations

linvoseltamab

teclistamab

talquetamab

elranatamab

including an immunomodulatory agent response rate and overall response rate

Annual Meeting three prior lines of therapy treatment of adult patients with relapsed

B-cell maturation antigen refractory AL

lenalidomide-refractory multiple myeloma who received one statistically significant adults with relapsed complete response

CARTITUDE-2 Cohort D study ciltacabtagene autoleucel study is to characterize the

data from the Phase patients with less than a complete relapsed or refractory multiple myeloma

European Commission CAR-T cell therapy

18 months infusion of CARVYKTI Orphan Drug Designation

efficacy and safety proteasome inhibitor

patients with functional high-risk multiple myeloma multiple myeloma patients

Phase II R/R Bispecifics N= 1 079 mentions Social media listening period: May-August 2024 Scope: worldwide in English



Phase II Relapse/Refractory CAR-T: Legend Biotech Shares Data on the Earliest Use to Date of CARVYKTI® in the Treatment of Multiple Myeloma and Important Subgroup Analyses at ASCO and EHA



Headlines/Hot off the press

Legend Biotech Shares Data on the Earliest Use to Date of CARVYKTI® in the Treatment of Multiple Myeloma and Important Subgroup Analyses at ASCO and EHA

"Legend Biotech Corporation (...), a global leader in cell therapy, announced today results for the first time from the Phase 2 CARTITUDE-2 Cohort D study in multiple myeloma patients. Results showed patients with less than a complete response (CR) after front-line autologous stem cell transplant (ASCT) experienced deep and durable responses following a single infusion of CARYYKTI * with or without lenglidomide maintenance.

Click here to read the full article

Sponsor



CARTITUDE-2

Ctrl + click to access clinical trial: NCT04133636

"A Study of JNJ-68284528, a Chimeric Antigen Receptor T Cell (CAR-T) Therapy Directed Against B-cell Maturation Antigen (BCMA) in Participants With Multiple Myeloma"

Combinations

ciltacabtagene autoleucel



Drugs in clinical trials (Phase III) monitored during the listening period

T	

Note

Sponsor: University of Chicago

Sponsor: Poitiers University Hospital

Sponsor: SWOG Cancer Research Network

Sponsor: European Myeloma Network

Sponsor: European Myeloma Network

Sponsor: University of Leeds

Sponsor: ECOG-ACRIN Cancer Research Group

Sponsor: University of Heidelberg Medical Center

	Diag	5 III CIII II CUI UI UI S	(Filase III) IIIoli	intored during the listering period
Sponsor		Trial name + link (ctrl + right click)	Line of treatment	Combination/Monotherapy being studied
AbbVie	Bellini		RR	venetoclax, bortezomib, dexamethasone

RR

RR

RR

RR

RR

RR

Maintenance

RR

RR

1L

RR

RR

RR

RR

RR

11

RR

RR

RR

1L

Maintenance

1L

1L

1L

1L

1L

1L

CANOVA

KarMMa-3

DREAMM-3

DREAMM 7

DREAMM 8

CARTITUDE-4

AURIGA

CASTOR

MAIA

MajesTEC-3

POLLUX

BOSTON

OCEAN

IMROZ

IKEMA

ATLAS

BENEFIIT

EQUATE

IsKia

DRAMMATIC

GMMG HD7

Myeloma XI **PERSEUS**

ICARIA-MM

TOURMALINE-MM1

TOURMALINE-MM2

MagnetisMM-5

Celgene

GSK

Janssen

Karyopharm

Oncopeptides

Pfizer

Sanofi

Takeda

Other

DETERMINATION

	Sponsor	Trial name + link (ctrl + right click)	Line of treatment	Combination/Monotherapy being studied		
brugs in chinical trials (Friase III) mornitorea daring the listering period						

venetoclax, dexamethasone

daratumumab, lenalidomide

lenalidomide, bortezomib, dexamethasone

belantamab mafodotin, bortezomib and dexamethasone

teclistamab, daratumumab Subcutaneously (SC) (Tec-Dara)

Monotherapy: elranatamab or doublet therapy with daratumumab

belantamab mafodotin Plus pomalidomide and dexamethasone

Monotherapy: Idecabtagene vicleucel

Monotherapy: belantamab mafodotin

Monotherapy: ciltacabtagene autoleucel

Monotherapy: melphalan flufenamide

isatuximab, pomalidomide, dexamethasone

isatuximab, carfilzomib And dexamethasone

ixazomib, lenalidomide, dexamethasone

ixazomib, lenalidomide, dexamethasone

lenalidomide, carfilzomib, dexamethasone

isatuximab, lenalidomide, dexamethasone

daratumumab, bortezomib, lenalidomide, dexamethasone

Isatuximab, lenalidomide, bortezomib, dexamethasone

isatuximab, lenalidomide, carfilzomib, dexamethasone

daratumumab, bortezomib, dexamethasone, lenalidomide

cyclophosphamide, dexamethasone plus lenalidomide with or without carfilzomib

lenalidomide, daratumumab

daratumumab, bortezomib, dexamethasone

daratumumab, lenalidomide, dexamethasone

daratumumab, lenalidomide, and dexamethasone

isatuximab, bortezomib, lenalidomide dexamethasone

Phase III Line 1 Triplet Therapy: DARZALEX® (daratumumab)-based regimens significantly improve clinical outcomes in both transplant-eligible and -ineligible patients who are newly diagnosed with multiple myeloma



Headlines/Hot off the press

DARZALEX® (daratumumab)-based regimens significantly improve clinical outcomes in both transplant-eligible and -ineligible patients who are newly diagnosed with multiple myeloma

"7.5 years median overall survival achieved with DARZALEX®-based regimen in Phase 3 MAIA final analysis is the longest reported in patients ineligible for transplant

Click here to read the full article

11

MAIA

Ctrl + click to access clinical trial:

clinical trial: NCT02252172 "Study Comparing Daratumumab, Lenalidomide, and Dexamethasone With Lenalidomide and Dexamethasone in Participants With Previously Untreated Multiple Myeloma"

Combinations

Sponsor

daratumumab



lenalidomide

dexamethasone

janssen /

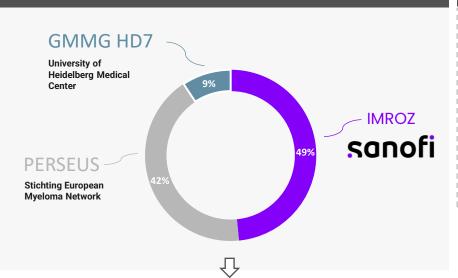




Phase III Line 1 Quadruplet Therapy: ASCO: Sarclisa is first anti-CD38 to significantly improve progression-free survival in combination with VRd for newly diagnosed transplant-ineligible multiple myeloma in phase 3



Clinical trial mentioned/Sponsor



Headlines/Hot off the press

Sarclisa accepted for FDA priority review for the treatment of transplant-ineligible newly diagnosed multiple myeloma

Click here to read the full article

ASCO: Sarclisa is first anti-CD38 to significantly improve progression-free survival in combination with VRd for newly diagnosed transplant-ineligible multiple myeloma in phase 3

Click here to read the full article

DARZALEX® (daratumumab)-based regimens significantly improved clinical outcomes in both transplant-eligible and -ineligible patients who are newly diagnosed with multiple myeloma

Click here to read the full article

FDA approves J&J's multiple myeloma therapy
Click here to read the full article

66 Isatuximab/VRd Improves PFS vs VRd Alone in Transplant-Eligible NDMM

Click here to read the full article

IMROZ

Ctrl + click to access clinical trial:

NCT03319667

PERSEUS

Ctrl + click to access clinical trial:

NCT03710603

"A Phase 3 Study Comparing Daratumumab, VELCADE (Bortezomib), Lenalidomide, and Dexamethasone (D-VRd) vs VELCADE, Lenalidomide, and Dexamethasone (VRd) in Subjects With Previously Untreated Multiple Myeloma Who Are Eligible for High-dose Therapy"

A Phase 3 Randomized, Open-label, Multicenter Study

Lenalidomide and Dexamethasone Versus Bortezomib

Diagnosed Multiple Myeloma Not Eligible for Transplant

in Combination With Bortezomib (Velcade®

Assessing the Clinical Benefit of Isatuximab (SAR650984)

Lenalidomide and Dexamethasone in Patients With Newly

GMMG HD7

Ctrl + click to access clinical trial:

NCT03617731

"Trial on the Effect of Isatuximab to Lenalidomide/Bortezomib/Dexamethasone (RVd) Induction and Lenalidomide Maintenance in Patients With Newly Diagnosed Myeloma"

Combinations

isatuximab + dexamethasone
- lenalidomide bortezomib

daratumumab + dexamethasone lenglidomide + bortezomib

isatuximab + lenalidomide + bortezomib + dexamethasone combination with bortezomib risk of disease progression or death

patients with multiple myeloma eligible for an autologous stem transplant-ineligible newly diagnosed multiple myeloma

sustained MRD negativity induction and consolidation in patients newly diagnosed transplant-ineligible multiple myeloma

Food and Drug Administration approved DARZALEX FASPRO FDA approved DARZALEX FASPRO

newly diagnosed patients

treatment of patients providing them with a new treatment induction therapy

complete response

patients with transplant-ineligible



SYNDICATED

Phase III Relapse/Refractory Doublet Therapy: ELREXFIO™ Shows Median Overall Survival of More Than Two Years in People with Relapsed or Refractory Multiple Myeloma



Headlines/Hot off the press



ELREXFIO™ Shows Median Overall Survival of More Than Two Years in People with Relapsed or Refractory Multiple Myeloma

"Pfizer Inc. announced detailed overall survival results from the Phase 2 MagnetisMM-3 study of ELREXFIO™ in patients with heavily pretreated relapsed or refractory multiple myeloma."

Click here to read the full article



Ctrl + click to access clinical trial:

NCT05020236

"AN OPEN-LABEL, 3-ARM, MULTICENTER, RANDOMIZED PHASE 3 STUDY TO EVALUATE THE EFFICACY AND SAFETY OF ELRANATAMAB (PF-06863135) MONOTHERAPY AND ELRANATAMAB + DARATUMUMAB VERSUS DARATUMUMAB + POMALIDOMIDE + DEXAMETHASONE IN PARTICIPANTS WITH RELAPSED/REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 1 PRIOR LINE OF THERAPY INCLUDING LENALIDOMIDE AND A PROTEASOME INHIBITOR"

Sponsor



Combinations

elranatamab

daratumumab





Phase III Relapse/Refractory Triplet Therapy: GSK Reignites Blenrep Blockbuster Aspirations With Data from **Phase III Studies**



Headlines/Hot off the press

🍊 GSK's Blenrep Combo Therapy Cuts Risk Of Disease Progression Or Death 🦡 By Around 50% In Pretreated Multiple Myeloma

Click here to read the full article

GSK Reignites Blenrep Blockbuster Aspirations With Data from Phase III Studies

Click here to read the full article

GSK Blenrep Combos for Multiple Myeloma Accepted for EU Review

Click here to read the full article

DREAMM-8

Ctrl + click to access clinical trial: NCT04484623

"A Phase III, Multicenter, Open-Label, Randomized Study to Evaluate the Efficacy and Safety of Belantamab Mafodotin in Combination With Pomalidomide and Dexamethasone (B-Pd) Versus Pomalidomide Plus Bortezomib and Dexamethasone (PVd) in Participants With Relapsed/Refractory Multiple Myeloma"

Sponsor



Combinations

belantamab mafodotin

dexamethasone

pomalidomide

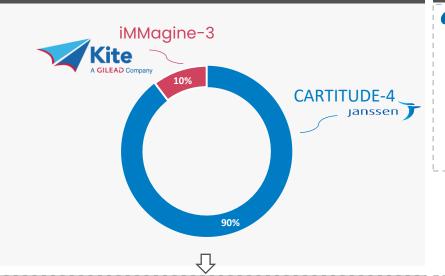




Phase III relapse/refractory CART: Legend's Phase III Win Cements Carvykti as Earlier-Line Multiple Myeloma Treatment



Clinical trial mentioned/Sponsor



Headlines/Hot off the press

66 CARVYKTI® significantly improved progressionfree survival and deepened responses versus
two standard therapies for patients with
functional high-risk multiple myeloma
Click here to read the full article

Legend's Phase III Win Cements Carvykti as Earlier-Line Multiple Myeloma Treatment

Click here to read the full article

Arcellx and Kite Continue Momentum with

Advances in Anito-Cel Multiple Myeloma Program

Click here to read the full article

CARTITUDE-4
Ctrl + click to access clinical

trial: NCT04181827

"A Study Comparing JNJ-68284528, a CAR-T Therapy Directed Against B-cell Maturation Antigen (BCMA), Versus Pomalidomide, Bortezomib and Dexamethasone (PVd) or Daratumumab, Pomalidomide and Dexamethasone (DPd) in Participants With Relapsed and Lenalidomide-Refractory Multiple Myeloma"

Combinations

ciltacabtagene autoleucel

iMMagine-3 Ctrl + click to access clinical trial:

"The goal of this study (iMMagine-3) is to compare the study drug, anitocabtagene autoleucel to standard of care therapy (SOCT) in participants with relapsed/refractory multiple myeloma who have received 1 to 3 prior lines of therapy, including an anti-CD38 monoclonal antibody and an immunomodulatory drug."

anitocabtagene autoleucel

autologous stem cell transplant including an immunomodulatory agent chimeric antigen receptor

three prior lines of therapy
received one to three prior improvement in overall survival new freatments recently announced Food and Drug Administration ciltacabtagene autoleucel CAR-T cell therapy
received at least one prior multiple myeloma patients second interim analysis B-cell maturation antigen
including a proteasome inhibitor
single infusion one prior line of therapy

relapsed and refractory multiple myeloma

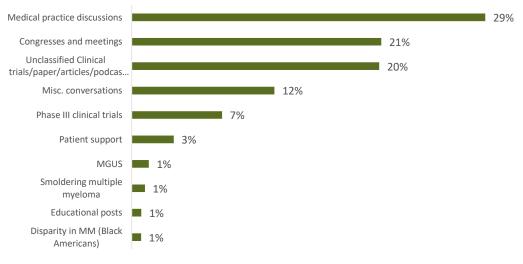
NC106413498

Phase III N= 2 210 mentions Social media listening period: May-August 2024 Scope: worldwide in English

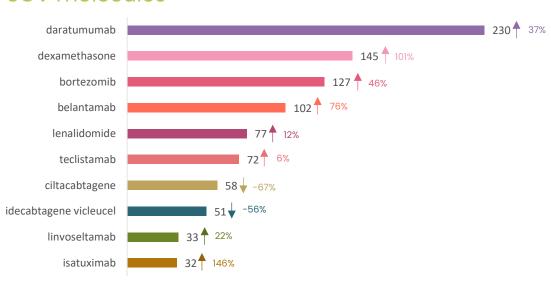


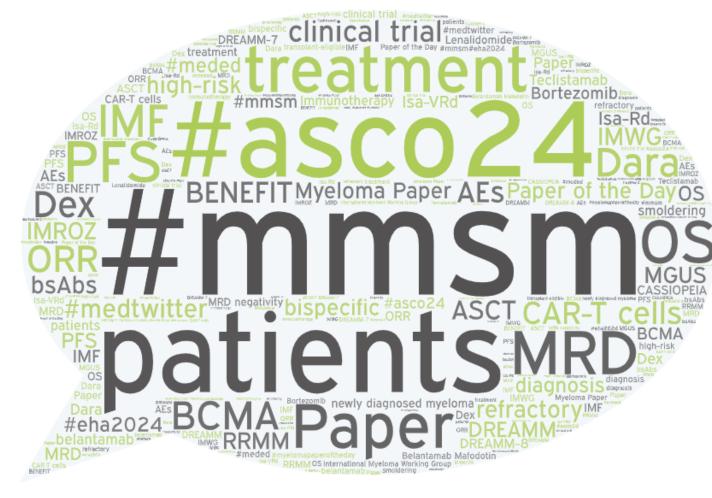
Congresses, most notably ASCO 2024 dominated the listening period. daratumumab was the most talked about molecule.

SOV topics of conversations



SOV molecules









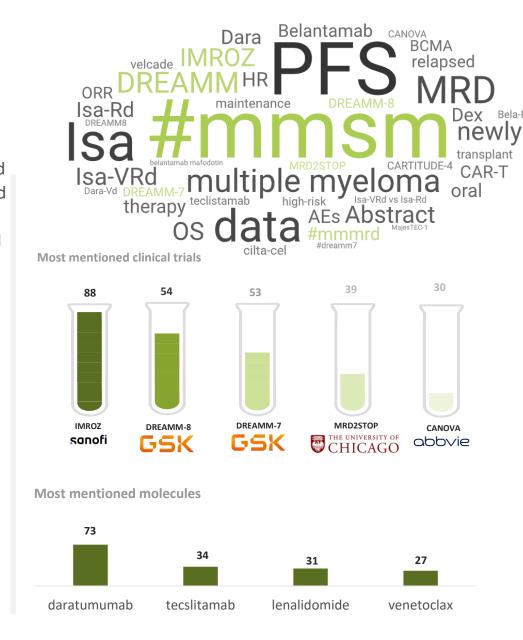
Focus on #ASCO2024: well received results for IMROZ

This year's ASCO conference was heavily focused on monoclonal antibody and more particularly daratumumab and antibody drug conjugate studied in the DREAMM-7 and 8 trials.

The **IMROZ** clinical trial, the most mentioned showing **improved PFS and OS** with the quad therapy **Isa-Vrd** vs Vrd alone in transplant ineligible NDMM patients was well received by HCPs.

ASCO 2023
1,119 mentions

ASCO 2024
981 mentions



First of the @NEJM #ASCO24 myeloma papers that I will be highlighting.

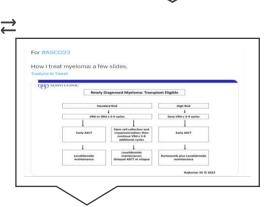
Congrats @dra_y_hungria @mvmateos @thanosdimop et al.
nejm.org/doi/full/10.10...

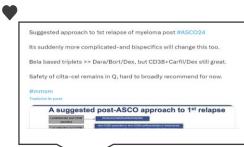
Traduire le post

I of 15

Belantamb Mafodotin, Bortezomib, and
Dexamethasone for Multiple Myeloma

Top posts from HCPs per engagement





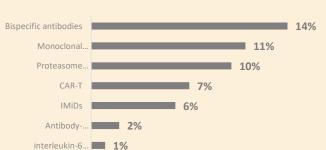
HCPs multiple myeloma ASCO 2024 universe N= 981 mentions Social media listening period: May 24th, 2024, to June 11th, 2024 Scope: worldwide in English

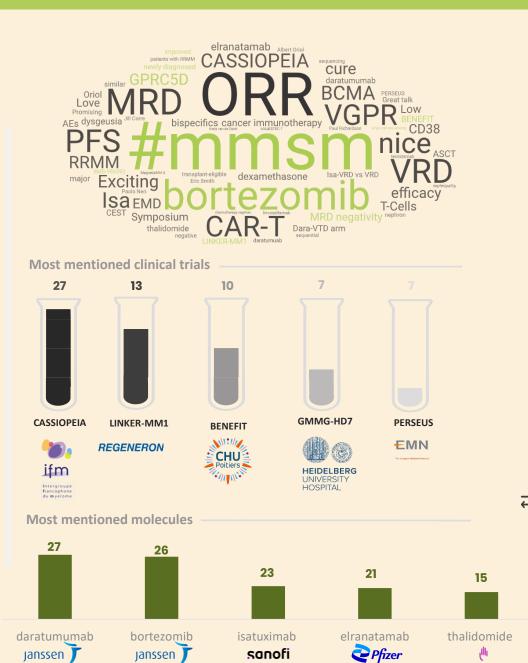
Focus on #EHA2024: CASSIOPEIA trial continues to impress

Long-term follow-up of the **CASSIOPEIA** clinical trial was presented during EHA 2024 and was the most mentioned clinical trial when it comes to multiple myeloma.

Results showed that more than **80%** of the patients were still alive after **8 years** in the daratumumab-VTD arm of the trial.

EHA 2023 260 mentions EHA 2024 363 mentions Topics of discussions (type of treatment)





First results of BMS-986393 GPRC5D CAR T-Cells in 1-3 prior lines of therapy presented at #EHA2024

ORR 96%, CR 42%
Low rates of dysgeusia/nail/skin changes
55% triple class refractory pts, 29% EMD
Median 2 prior lines of therapy

Top posts from HCPs per engagement

87% of responses ongoing



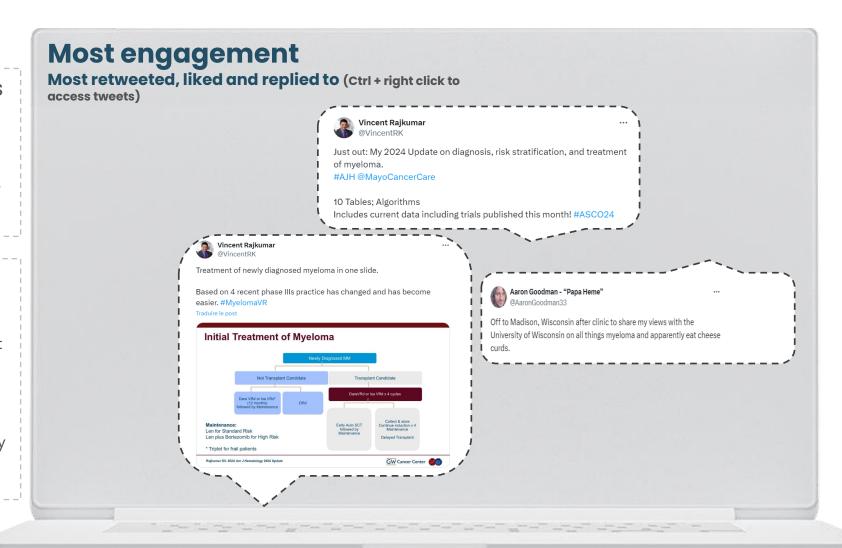
Always nice to see long-term follow up results! Dr. Moreau pointed out that >80% of patients in Dara-VTD arm of CASSIOPEIA are alive at 8 years despite half of them getting no maintenace! Exciting prospects indeed for limited duration therapy in #MultipleMyeloma! #EHA24

Engagement continues to be dominated by Vincent Rajkumar with educational posts.

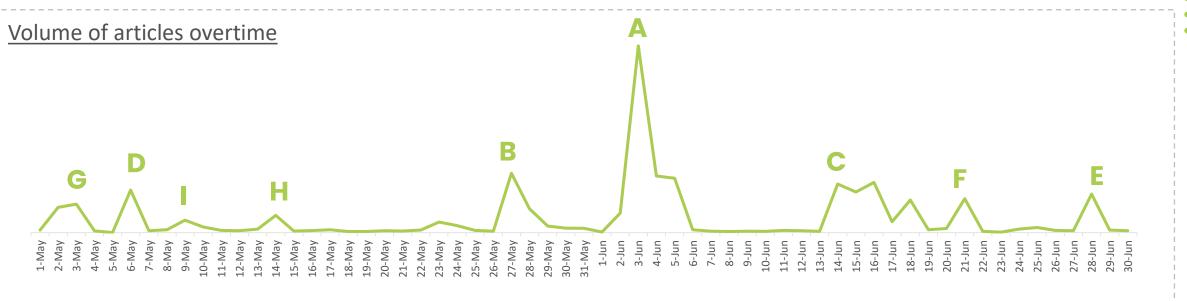
As he often does around the times of congresses, the influential Dr.
Rajkumar shared his recommendations on medical practices and treatment of newly diagnosed patients.

The first post redirects to an article titled "Multiple myeloma: 2024 update on diagnosis, risk-stratification, and management" available here. It was published in the American journal of hematology on June 28th, 2024.

The second post is related to the first and give an overview of the updated recommendation for newly diagnosed patients.







"CARVYKTI®▼ (ciltacabtagene autoleucel; cilta-cel) significantly improved progression-free survival and deepened responses versus two standard therapies for patients with functional high-risk multiple myeloma" Link

"Legend Biotech Shares Data on the Earliest Use to Date of CARVYKTI® in the Treatment of Multiple Myeloma and Important Subgroup Analyses at ASCO and EHA" Link

"GSK's Blenrep Combo Therapy Cuts Risk Of Disease Progression Or Death By Around 50% In Pretreated Multiple Myeloma" Link

"ASCO: Sarclisa is first anti-CD38 to significantly improve progression-free survival in combination with VRd for newly diagnosed transplant-ineligible multiple myeloma in phase 3" Link

"DARZALEX® (daratumumab)-based regimens significantly improved clinical outcomes in both transplant-eligible and -ineligible patients who are newly diagnosed with multiple myeloma" Link

"Johnson & Johnson At ASCO Highlights Long Term Data From Multiple Myeloma Drug, Shows Sustained Deep, Durable Responses In Pretreated Patients" Link

"TECVAYLI® ▼ (teclistamab) shows sustained deep and durable responses in patients with relapsed or refractory multiple myeloma" Link

- "Sarclisa accepted for FDA priority review for the treatment of transplant-ineligible newly diagnosed multiple myeloma" Link
- "Updated Linvoseltamab Data Showcase Continued **Deepening of Responses** in Patients with Heavily Pre-Treated Multiple Myeloma" Link

"BioLineRx Announces Poster Presentation on Economic Model Data for APHEXDA® (motixafortide) as part of CD34+ Hematopoietic Stem Cell Mobilization in Patients with Multiple Myeloma at ISPOR 2024" Link

"Comprehensively Covering Multiple Myeloma: IASO Bio's GPRC5D CAR-T Product RD118 Receives IND **Approval from NMPA**" Link

- "Starton Therapeutics Announces 100% Partial Response or Better in Multiple Myeloma Patients in Interim, Top-line Data from STAR-LLD Phase 1b Clinical" Link
- "Cleveland Clinic's Efficient, Lifesaving Treatment for Multiple Myeloma Giving Hope to Lives "Link





Media (press releases): volume and articles per peak July-August 2024



"DARZALEX FASPRO® (daratumumab and hyaluronidasefihj)-based quadruplet regimen approved in the U.S. for patients with newly diagnosed multiple myeloma who are transplant-eligible" Link

"New research shows how machine learning could revolutionize diagnosis and treatment of multiple myeloma and sepsis" Link

"FDA approves J&J's multiple myeloma therapy" Link

"Legend Biotech Announces Positive Overall Survival Results of Landmark Phase 3 CARTITUDE-4 Trial in Multiple Myeloma" Link

Media (press release, excluding financial news) N= 2411 mentions Social media listening period: July- August 2024 Scope: worldwide in English

- "GenScript Subsidiary Legend Biotech Achieves Breakthrough with Cilta-cel Approval in China, Offering New Hope for Multiple Myeloma Patients" Link
- "Sarclisa induction treatment demonstrated significantly improved progression-free survival in patients with newly diagnosed multiple myeloma eligible for transplant" Link
 - "Sanofi's **Sarclisa wins** in transplant-eligible multiple myeloma trial" Link
- "The Multiple Myeloma Research Foundation, Road to Victories, begins this August through the California Wine Country" Link

- "Multiple Myeloma Research Foundation (MMRF) Announces New Publication on **CoMMpass**SM Study in Nature Genetics Defining Distinct Subtypes of Multiple Myeloma and High-Risk Genetic Markers" Link
- "Join a movement to accelerate a cure for multiple myeloma! The Multiple Myeloma Research Foundation® Walk/Run" Link
- "Bionano Announces Publication of First Multi-Site Study to Analyze the **Utility of OGM in Multiple** Myeloma" Link

SYNDICATED

Articles per theme

Antibody-drug conjugates

"GSK's Blenrep Combo Therapy Cuts Risk Of Disease **Progression Or Death** By Around 50% In Pretreated Multiple Myeloma" Link

Bispecific antibody

"Johnson & Johnson At ASCO Highlights Long Term Data From Multiple Myeloma Drug, Shows Sustained Deep, Durable Responses In Pretreated Patients" Link

"TECVAYLI® ▼ (teclistamab) shows sustained deep and durable responses in patients with relapsed or refractory multiple myeloma" Link

"Updated Linvoseltamab Data Showcase Continued Deepening of Responses in Patients with Heavily Pre-Treated Multiple Myeloma" Link

CAR T-cell therapy

"Comprehensively Covering Multiple Myeloma: IASO Bio's GPRC5D CAR-T Product RD118 Receives IND Approval from NMPA" Link

"GenScript Subsidiary Legend Biotech Achieves Breakthrough with Cilta-cel Approval in China, Offering New Hope for Multiple Myeloma Patients" Link

CAR T-cell therapy (cont.)

"CARVYKTI®▼ (ciltacabtagene autoleucel; cilta-cel) significantly improved progression-free survival and deepened responses versus two standard therapies for patients with functional high-risk multiple myeloma" Link

"Legend Biotech Shares Data on the Earliest Use to Date of CARVYKTI® in the Treatment of Multiple Myeloma and Important Subgroup Analyses at ASCO and EHA" Link

Clinical trials and other studies

"New research shows how machine learning could revolutionize diagnosis and treatment of multiple myeloma and sepsis" Link

"Bionano Announces Publication of First Multi-Site Study to Analyze the Utility of OGM in Multiple Myeloma" Link

"Multiple Myeloma Research Foundation (MMRF) Announces New Publication on **CoMMpass**SM Study in Nature Genetics Defining Distinct Subtypes of Multiple Myeloma and High-Risk Genetic Markers" Link

"Starton Therapeutics Announces 100% Partial Response or Better in Multiple Myeloma Patients in Interim, Top-line Data from STAR-LLD Phase 1b Clinical" Link

Hematopoietic Stem Cell Mobilizer

"BioLineRx Announces Poster Presentation on Economic Model Data for APHEXDA® (motixafortide) as part of CD34+ Hematopoietic Stem Cell Mobilization in Patients with Multiple Myeloma at ISPOR 2024" Link

Misc.

"Cleveland Clinic's Efficient, Lifesaving Treatment for Multiple Myeloma Giving Hope to Lives "Link



"DARZALEX® (daratumumab)-based regimens significantly improved clinical outcomes in both transplant-eligible and -ineligible patients who are newly diagnosed with multiple myeloma" Link

"ASCO: Sarclisa is first anti-CD38 to significantly improve progression-free survival in combination with VRd for newly diagnosed transplant-ineligible multiple myeloma in phase 3" Link

"Sarclisa accepted for FDA priority review for the treatment of transplant-ineligible newly diagnosed multiple myeloma" Link

"Legend Biotech Announces Positive Overall Survival Results of Landmark Phase 3 CARTITUDE-4 Trial in Multiple Myeloma" <u>Link</u>

"FDA approves J&J's multiple myeloma therapy" Link

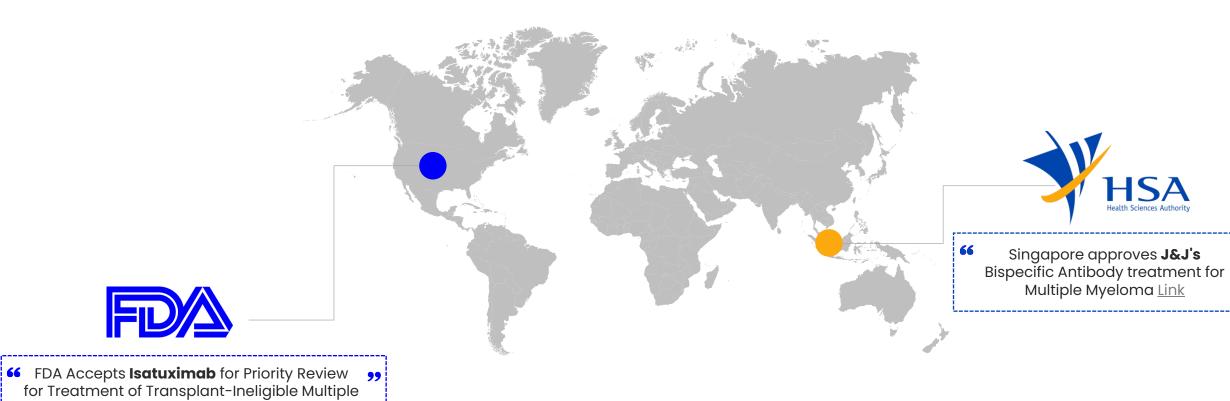
Multiple myeloma awareness

"The Multiple Myeloma Research Foundation, Road to Victories, begins this August through the California Wine Country" Link

"Join a movement to accelerate a cure for multiple myeloma! The Multiple Myeloma Research Foundation® Walk/Run" Link

May 2024





Myeloma <u>Link</u>

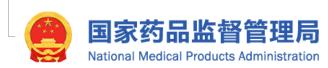
June 2024







Janssen's bispecific antibody drug for multiple myeloma wins nod in Korea Link



IASO Biotechnology Secures NMPA Approval for Physical Secures NMPA Approval for IND Application of RD118 CAR-T Therapy Targeting GPRC5D for Relapsed/Refractory Multiple Myeloma Link

July 2024



National Institute for Health and Care Excellence

J&J's Tecvayli recommended by 🤧 NICE to treat relapsed and refractory multiple myeloma



FDA **approves J&J's** multiple myeloma therapy Link

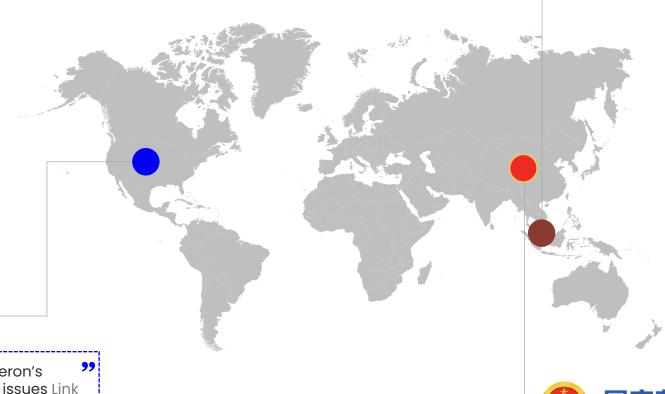
EUROPEAN MEDICINES AGENCY SCIENCE MEDICINES HEALTH

GSK **Blenrep** Combos for Multiple Myeloma Accepted for EU Review



August 2024





Malaysia **approves** Antengene's **XPOVIO NDA** for multiple myeloma Link

FDA declines to approve Regeneron's linvoseltamab amid manufacturing issues Link



GenScript Subsidiary Legend Biotech Achieves Breakthrough with Cilta-cel Approval in China, Offering New Hope for Multiple Myeloma Patients