



4 November 2024
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Human Medicines Division

CAT quarterly highlights and approved ATMPs

November 2024

This report provides information on Advanced Therapy Medicinal Products (ATMPs) approvals and extension of indications of authorised ATMPs, as well as statistical data on product-related activities.

The period covered by this report is: August - October 2024.

Advanced therapy medicinal products approvals

No new approvals during the reporting period

Extension of indication of authorised ATMPs

No procedures during the reporting period

Overview of product-related activities

The Committee discussed ongoing evaluation procedures for ATMPs and other related procedures as summarised in the following tables:

Initial Evaluation of Marketing Authorisation Applications (MAA) for ATMP						
	2009-2020	2021	2022	2023	2024*	Total
Submitted MAAs	32	3	1	3	6	45
Positive draft Opinion	18 ⁱ	2	6	1	1	28 [#]
Negative draft opinions	4 ^{i,ii,iii}	0	0	0	0	4
Withdrawals	8 ^{ii, iv}	0	1 ^v	1 ^{vi}	0	10
Ongoing MAAs						7

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Corresponding to 27 ATMPs (see List of authorised ATMPs)

ⁱ One negative draft opinion and two positive draft opinions for the Glybera

ⁱⁱ Negative draft opinion and withdrawal for the Cerepro

ⁱⁱⁱ Two negative draft opinions for Heparesc

^{iv} Luxceptar, Roctavian, Artobend

^v Sitoiganap

^{vi} Lumevoq

Variations (Type II) for authorised ATMP						
	2009-2020	2021	2022	2023	2024*	Total
Positive opinion	78	32	47	49	27	233

Scientific recommendation on advanced therapy classification ¹						
	2009-2020	2021	2022	2023	2024*	Total
Submitted	489	66	51	43	33	682
Adopted	483	61	46	52	33	675

Scientific advice procedure for ATMPs						
	2009-2020	2021	2022	2023	2024*	Total
Number of procedures	442	64	53	57	46	682

PRIME ² Eligibility for ATMPs						
	2016-2020	2021	2022	2023	2024*	Total
Discussed	91	14	10	13	12	140
Granted	39	7	4	9	6	65

* Period: January – October 2024

¹ More information on the scientific recommendation on advanced therapy classification and the summaries of ATMP classification can be found on the [ATMP classification webpage](#).

It is stressed that the scientific recommendation on advanced therapy classification does not amount to any endorsement of the plausibility of the product, including the mode of action or therapeutic indication(s) claimed by the applicant.

² PRiority MEdicines (PRIME) scheme. PRIME was set up in March 2016 to provide early and enhanced scientific and regulatory support to medicines that have the potential to significantly address patients' unmet medical needs. More information can be found at the [PRIME webpage](#).

List of authorised ATMPs

NAME	Type of ATMP	Authorisation Date	Orphan	PRIME	Comment
Chondroelect	TEP	5/10/2009	No	No	MA withdrawn July 2016
Glybera	GTMP	25/10/2012	Yes	No	MA not renewed (MA ended Oct. 2017)
MACI	TEP, combined ATMP	27/06/2013	No	No	MA not renewed (MA ended June 2018)
Provenge	CTMP	6/09/2013	No	No	MA withdrawn May 2015
Holoclax	TEP	17/02/2015	Yes	No	
Imlygic	GTMP	16/12/2015	No	No	
Strimvelis	GTMP	26/05/2016	Yes	No	
Zalmoxis	CTMP	18/08/2016	Yes	No	MA withdrawn Oct. 2019
Spherox	TEP	10/07/2017	No	No	
Alofisel	CTMP	23/03/2018	Yes	No	
Yescarta	GTMP	23/08/2018	Yes	Yes	
Kymriah	GTMP	23/08/2018	Yes	Yes	
Luxturna	GTMP	22/11/2018	Yes	No	
Zynteglo	GTMP	29/05/2019	Yes	Yes	MA withdrawn March 2022
Zolgensma	GTMP	18/05/2020	Yes	Yes	
Libmeldy	GTMP	17/12/2020	Yes	No	
Tecartus	GTMP	14/12/2020	Yes	Yes	
Skysona	GTMP	16/07/2021	Yes	Yes	MA withdrawn Nov. 2021
Abecma	GTMP	18/08/2021	Yes	Yes	
Breyanzi	GTMP	4/04/2022	No	Yes	
Carvykti	GTMP	25/05/2022	Yes	Yes	
Upstaza	GTMP	18/07/2022	Yes	No	

NAME	Type of ATMP	Authorisation Date	Orphan	PRIME	Comment
Roctavian	GTMP	24/08/2022	Yes	Yes	
Ebvallo	CTMP	16/12/2022	Yes	Yes	
Hemgenix	GTMP	20/02/2023	Yes	Yes	
Casgevy	GTMP	9/02/2024	Yes	Yes	
Durveqtix	GTMP	24/07/2024	No	Yes	

More information on authorised products can be found on: www.ema.europa.eu (type in the product name in the search box)

Abbreviations: ATMP: advanced therapy medicinal product; GTMP: gene therapy medicinal product; CTMP: cell therapy medicinal product; TEP: tissue engineered product; MA: Marketing authorisation

