



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

21 October 2019
EMA/573788/2019
Information Management Division

Monthly statistics report: September 2019

Medicinal products for human use (cumulative figures for the year to date)

This document provides current information related to the volume and evaluation of marketing authorisation and post-authorisation applications for medicinal products for human use received by the European Medicines Agency.

The purpose is only to provide ongoing factual information. Commentaries and analysis are provided in the Agency's annual reports.

Official address Domenico Scarlattilaan 6 • 1083 HS Amsterdam • The Netherlands

Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us

Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000

An agency of the European Union



Table 1. Pre-authorisation: Marketing-authorisation applications*

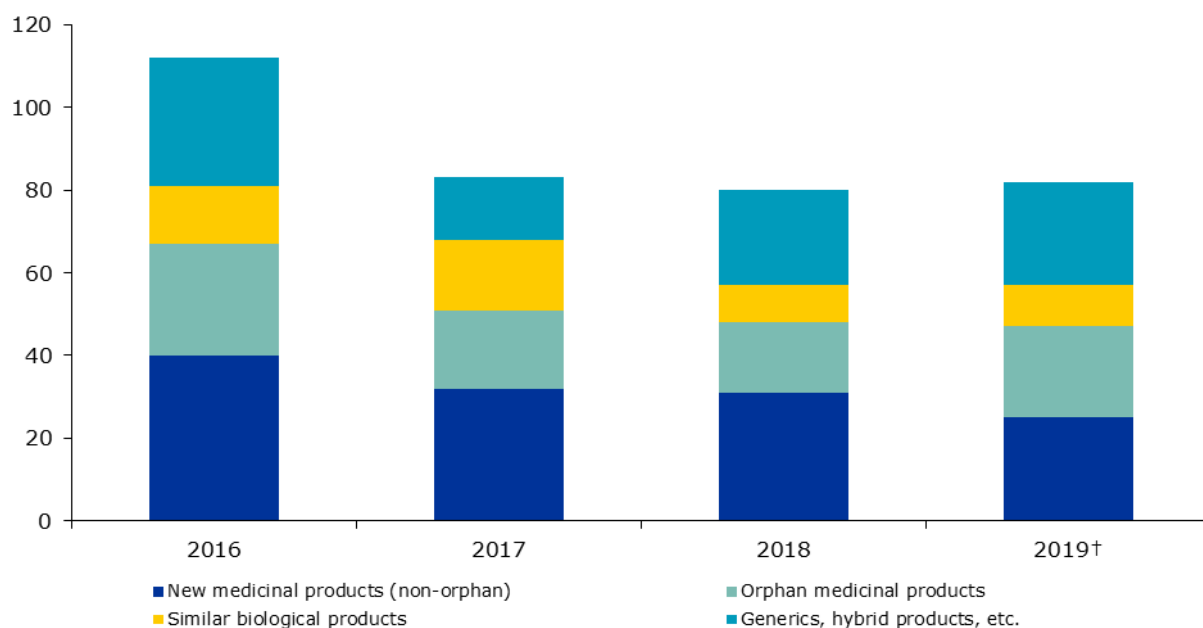
	2016		2017		2018		2019†	
	Started	Finalised	Started	Finalised	Started	Finalised	Started	Finalised
Non-orphan medicinal products								
New products	40	28	32	33	31	34	25	20
Advanced-therapy medicinal products	0	0	0	1	1	0	0	0
Paediatric-use (PUMA) products	1	1	2	1	0	2	0	0
Well-established use, abridged, hybrid and informed consent products	7	5	5	6	5	6	11	8
Generic products	24	22	10	22	18	9	14	11
Similar biological products	14	7	17	14	9	15	10	3
Sub-total product applications	86	63	66	77	64	66	60	42
Orphan medicinal products[◇]								
New products	27	16	19	20	17	20	22	8
Advanced-therapy medicinal products	1	2	4	1	2	3	0	1
Total product applications	114	79	89	101	83	89	82	51

* Finalised applications exclude applications withdrawn prior to opinion.

† Figures for the current year are cumulative, year to date. Figures for preceding years are totals for the year.

◇ These figures reflect the orphan status of the medicinal products at the time of the CHMP opinion. EMA's Committee for Orphan Medicinal Products (COMP) then assesses whether the orphan designation should be maintained.

Marketing authorisation application evaluations started by type of application



† Figures for the current year are cumulative, year to date. Figures for preceding years are totals for the year.

Table 2. Pre-authorisation: Outcome of the evaluation of marketing authorisation applications*

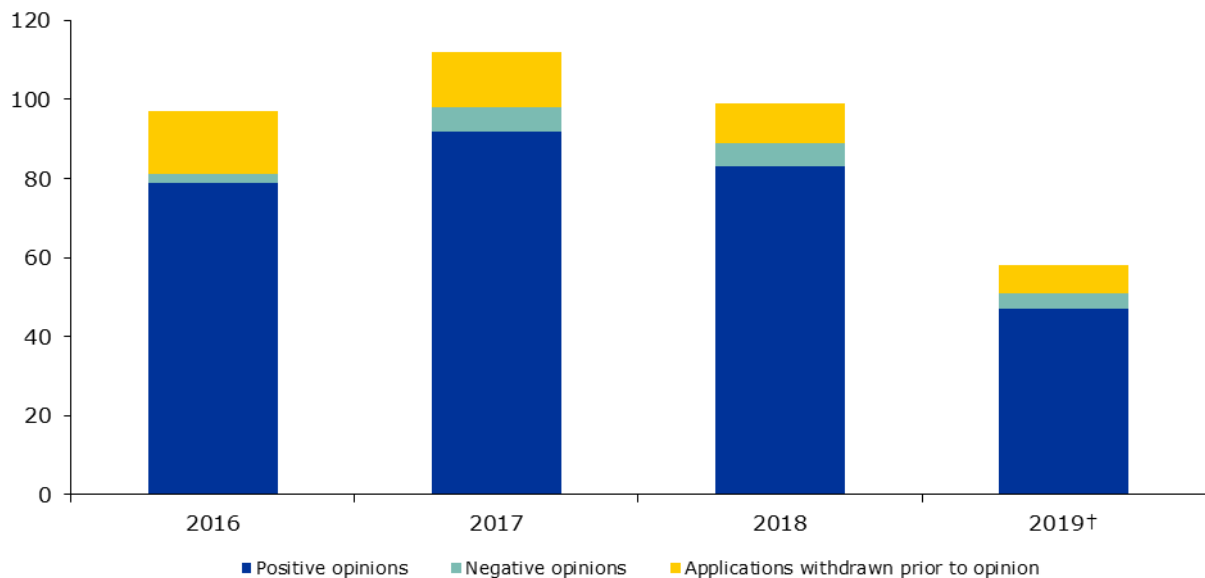
	2016	2017	2018	2019 [†]
Positive opinions	79	92	83	47
Opinions recommending conditional marketing authorisation**	7	3	1	6
Opinions under exceptional circumstances**	1	2	3	1
Negative opinions	2	6	6	4
Opinions after accelerated assessment**	7	7	4	2
Applications withdrawn prior to opinion	16	14	10	7
Re-examinations requested	2	5	5	3
Re-examination - Positive opinions	2	0	1	0

* Applicants can request a re-examination. The first four rows present the outcome of the evaluation before a re-examination (or a re-consideration). The final row shows the number of changes from a negative to a positive opinion following a re-examination or a re-consideration.

** Included in the figures for positive opinions.

[†] Figures for the current year are cumulative, year to date. Figures for preceding years are totals for the year.

Pre-authorisation: Outcome of the evaluation of marketing authorisation applications



[†] Figures for the current year are cumulative, year to date. Figures for preceding years are totals for the year.

Table 3. Scientific services

	2016		2017		2018		2019 [†]	
	Started	Finalised	Started	Finalised	Started	Finalised	Started	Finalised
Compassionate-use opinions	0	0	0	0	0	0	0	0
Art. 58 (WHO) scientific opinions	0	1	1	0	1	1	0	0
Opinions on ancillary medicinal substances in medical devices*	0	0	2	1	0	1	0	0
Plasma master file (includes initial certification, variations and annual re-certification)	19	22	22	24	19	18	15	18

* Consultation in accordance with Council Directive 93/42/EEC concerning medical devices as amended by Directive 2000/70/EC as regards medical devices incorporating stable derivatives of human blood or plasma and Directive 2001/14/EC.

[†] Figures for the current year are cumulative, year to date. Figures for preceding years are totals for the year.

Table 4. Post-authorisation: Variations, renewals and annual reassessments

	2016		2017		2018		2019 [†]	
	Started	Finalised	Started	Finalised	Started	Finalised	Started	Finalised
Type IA variations	3,019	2,934	3,080	3,069	3,433	3,347	2,853	2,759
Type IB variations	2,000	1,988	2,054	1,975	2,164	2,063	1,698	1,731
Type II variations	1,185	1,131	1,133	1,116	1,119	1,041	824	873
Extensions of marketing authorisation	25	16	21	25	20	19	14	9
Annual reassessments	25	19	19	22	22	22	15	15
Renewals*	107	89	94	90	90	101	42	65

* Includes renewals of conditional marketing authorisations.

[†] Figures for the current year are cumulative, year to date. Figures for preceding years are totals for the year.

Post-authorisation: Variations, renewals and annual reassessments

