



EUROPEAN MEDICINES AGENCY  
SCIENCE MEDICINES HEALTH

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Information Management Division

## Monthly statistics report: April 2020

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.

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

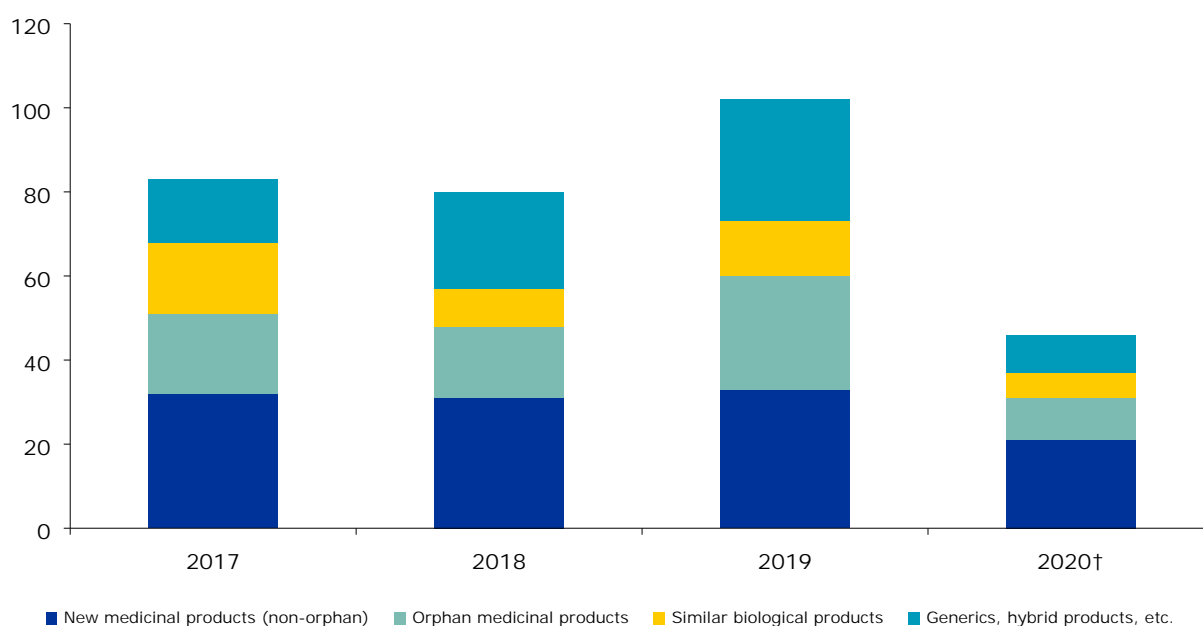
	2017		2018		2019		2020†	
	Started	Finalised	Started	Finalised	Started	Finalised	Started	Finalised
<b>Non-orphan medicinal products</b>								
New products	32	33	31	34	33	31	21	14
Advanced-therapy medicinal products	0	1	1	0	0	0	0	0
Paediatric-use (PUMA) products	2	1	0	2	0	0	1	0
Well-established use, abridged, hybrid and informed consent products	5	6	5	6	12	8	7	3
Generic products	10	22	18	9	17	15	2	6
Similar biological products	17	14	9	15	13	5	6	3
Sub-total product applications	66	77	64	66	75	59	37	26
<b>Orphan medicinal products<sup>◇</sup></b>								
New products	19	20	17	20	27	11	10	6
Advanced-therapy medicinal products	4	1	2	3	1	1	3	1
<b>Total product applications</b>	<b>89</b>	<b>98</b>	<b>83</b>	<b>89</b>	<b>103</b>	<b>71</b>	<b>50</b>	<b>33</b>

\* Finalised applications exclude applications withdrawn prior to opinion.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\*

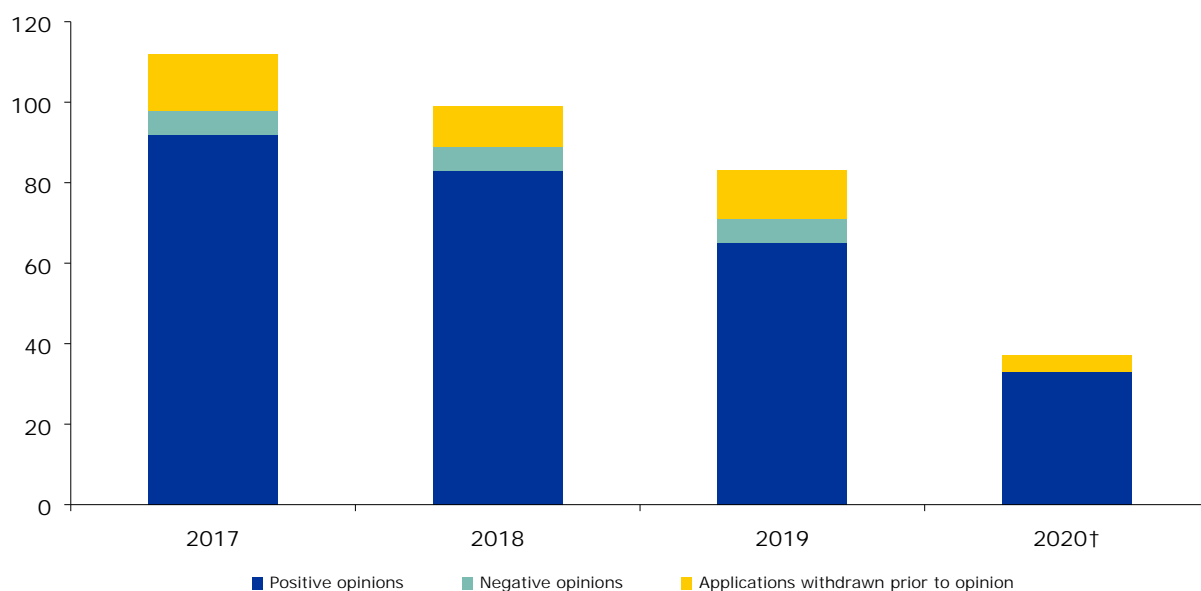
	2017	2018	2017	2020†
Positive opinions	92	83	65	33
Opinions recommending conditional marketing authorisation**	3	1	8	2
Opinions under exceptional circumstances**	2	3	1	0
Negative opinions	6	6	6	0
Opinions after accelerated assessment**	7	4	3	1
Applications withdrawn prior to opinion	14	10	12	4
Re-examinations requested	5	5	4	0
Re-examination - Positive opinions	0	1	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

	2017		2018		2019		2020 <sup>†</sup>	
	Started	Finalised	Started	Finalised	Started	Finalised	Started	Finalised
Compassionate-use opinions	0	0	0	0	0	0	1	1
Art. 58 (WHO) scientific opinions	1	0	1	1	0	0	0	0
Opinions on ancillary medicinal substances in medical devices*	2	1	0	1	0	0	0	0
Plasma master file (includes initial certification, variations and annual re-certification)	22	24	19	18	17	21	6	8

\* 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.

<sup>†</sup> 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

	2017		2018		2019		2020†	
	Started	Finalised	Started	Finalised	Started	Finalised	Started	Finalised
Type IA variations	3,080	3,069	3,433	3,347	3,886	3,849	1,276	1,162
Type IB variations	2,054	1,975	2,164	2,063	2,425	2,279	769	866
Type II variations	1,133	1,116	1,119	1,041	1,123	1,108	383	417
Extensions of marketing authorisation	21	25	20	19	27	19	15	10
Annual reassessments	19	22	22	22	25	23	6	9
Renewals*	94	90	90	101	107	85	39	44

\* 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

