31 January 2023
EMA/2/2023

Key performance indicators (KPIs) to monitor the European clinical trials environment

Metrics on the Clinical Trials Regulation and Clinical Trials Directive

1 – 31 December 2022, edition 9

On the 31 January 2022 the Clinical Trials Regulation (EU) No 536/2014, hereinafter 'CTR', repealing the Clinical Trials Directive 2001/20/EC, hereinafter 'CTD', became applicable and the Clinical Trial Information System (CTIS) was launched. In line with the provisions outlined in Article 97 of the Clinical Trials Regulation, the European Commission shall assess the impact of the Regulation on scientific and technological progress.

This report provides an overview of Key Performance Indicators (KPIs) related to the implementation of the CTR. The Clinical Trials Regulation Metrics report is published on a monthly basis starting in May 2022. The previous report can be found at this link.

This report is published as part of the business change programme Accelerating Clinical Trials EU (ACT EU), involving the European Commission, the Heads of Medicines Agencies (HMA), Clinical Trial Coordination Group (CTCG) and the Agency.

ACT EU seeks to transform how clinical trials are initiated, designed and run. One of the priority actions of ACT EU focusses on monitoring the implementation of the CTR.

The metrics presented in the report reflect the status of applications in CTIS and EudraCT[1] as of 31 December 2022 for Clinical Trial applications (CTA) submitted between 1-31 December 2022[2] as well as cumulative figures.

[1] EudraCT is the (European Union Drug Regulating Authorities Clinical Trials Database) European database for all interventional clinical trials on medicinal products authorised in the European Union (EEA) under the Clinical Trial Directive and outside the EU/EEA if they are part of a Paediatric Investigation Plan (PIP)
[2] The two 'smoke test' trials, submitted to CTIS for testing purposes just before the CTIS launch, are not counted.

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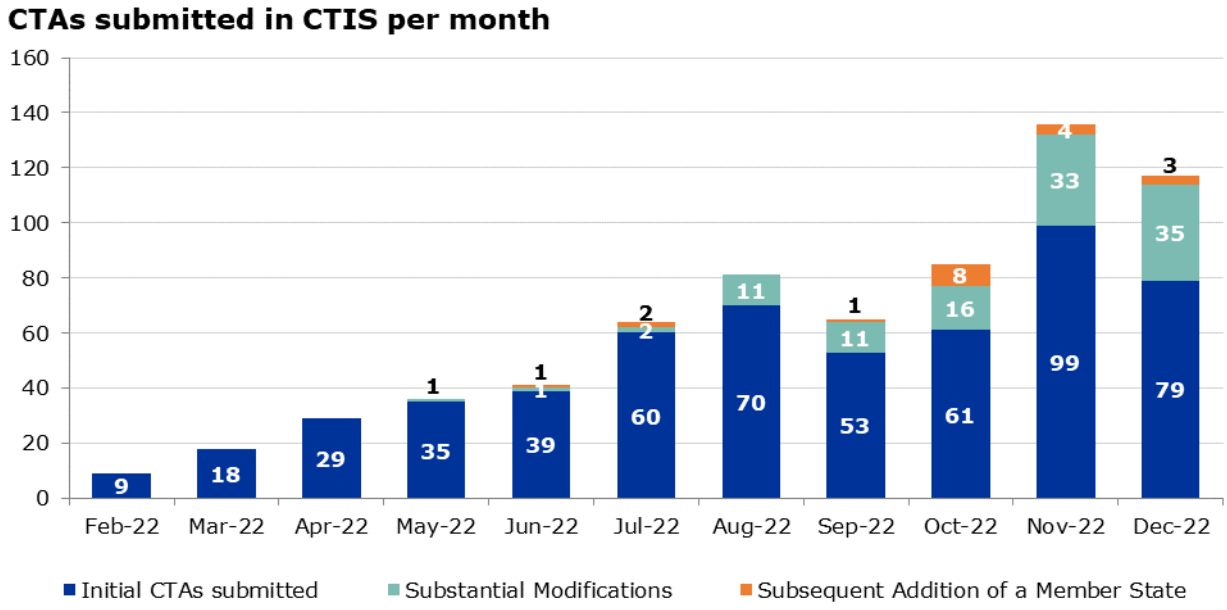
Clinical Trial Information System (CTIS) and EudraCT metrics

This report shows the key performance indicators (KPIs) generated from the two databases containing information on clinical trials in the EU/EEA, namely CTIS and EudraCT.

1.1. Number of clinical trial applications (CTAs) submitted under the Clinical Trials Regulation in CTIS

The graph below shows the cumulative number of clinical trial applications that have been submitted to CTIS since the launch on 31 January 2022.

Specifically, the applications submitted in December 2022 include initial clinical trial applications[3], substantial modifications[4] and addition of a new Member State Concerned[5] applications in the selected period.[6]



Overall, 681 clinical trial applications have been submitted in CTIS since the launch of the system on 31 January 2022, of which 552 are initial clinical trial applications, 110 are substantial modification applications and 19 are applications for the addition of a new Member State Concerned.

Of the submitted applications during December 2022, 5 are re-submissions of previously withdrawn applications.

[3] Initial clinical trials applications are those submitted in accordance with the requirements of Article 5 and Article 11, as applicable, of the Clinical Trials Regulation (EU) No 536/2014

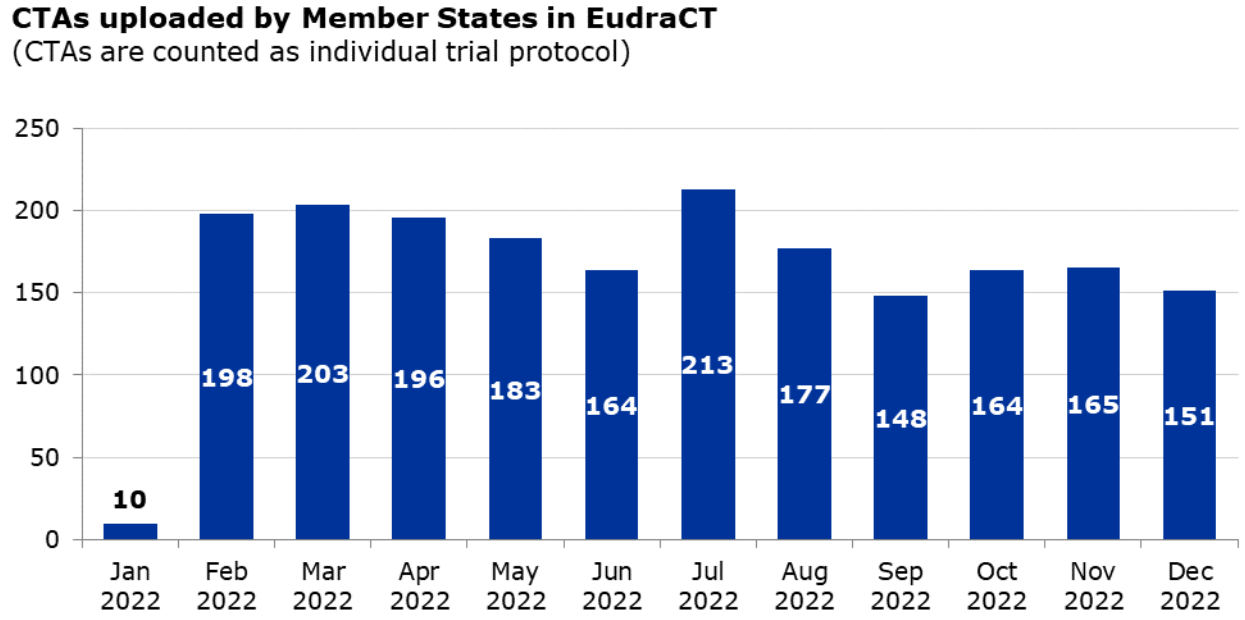
[4] Substantial modifications are those submitted in accordance with the requirements of chapter III of the Clinical Trials Regulation (EU) No 536/2014

[5] Applications to add a new Member States Concerned are submitted in accordance with the requirements of Article 14 of Regulation (EU) No 536/2014

[6] Corrigendum: the graph shows the corrected figures for initial clinical trial applications submitted in July which are 60 and not 58 as displayed in the report edition 4. The incorrect calculation was due to an error in the reporting system.

1.2. CTAs under Clinical Trial Directive (CTD) uploaded by Member States (MSs) in EudraCT, counted as individual clinical trial protocol

The graph below shows the number of CTAs uploaded by the Member States in EudraCT as individual clinical trial protocol, per month during the selected period[7]. Overall 1,972 CTAs have been uploaded in EudraCT.



1.3. Number of ongoing clinical trials (CTs)

CTs under the CTR with at least one positive decision in the EU

The term 'ongoing' refers to clinical trials that have been authorised in at least one Member State Concerned where the recruitment of patients has started at the clinical investigator sites[8].

As of 31 December 2022, 60 clinical trials were reported as ongoing in CTIS.

CTs under the CTD

In EudraCT there are no fields available to capture recruitment status at the site.

1.4. Number of trials for which a decision has been issued under the CTR with/without deferral[9] for the protocol

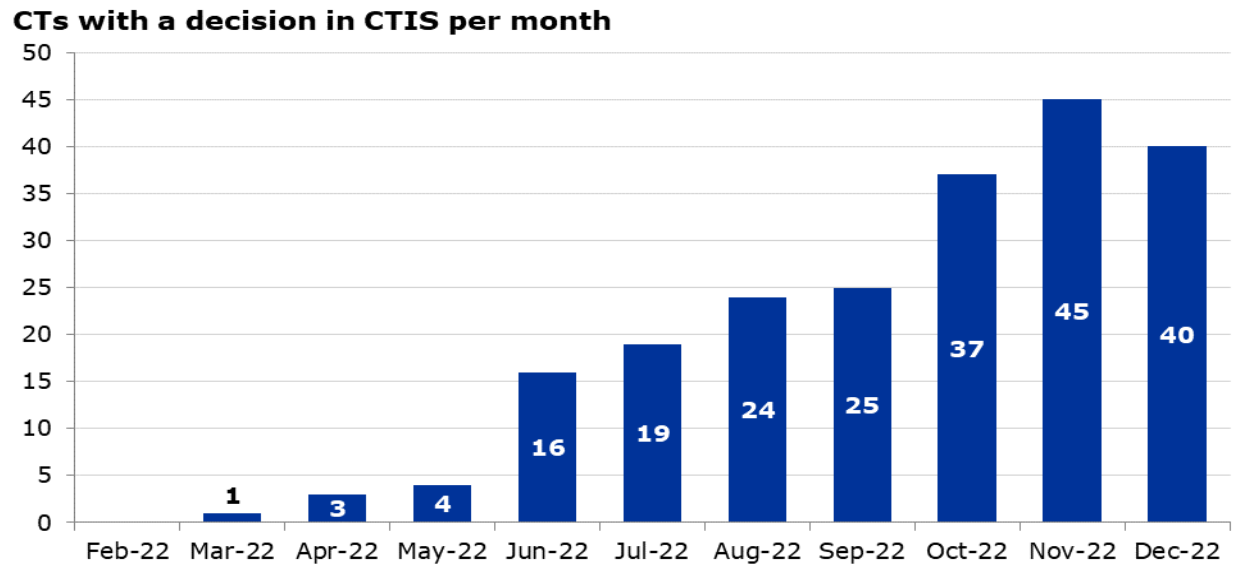
Due to a known issue, mitigation measure has been put in place to prevent publication of clinical trials with deferrals. As a consequence, clinical trials with any type of deferrals with a decision issued mid-August onwards are not available in the public domain. This is a temporary measure until the functionality of the deferral mechanism is restored. Sponsors and EU/EEA Members have the possibility to apply deferrals to clinical trials data, which will be published in due course once the issue is resolved.

Therefore, information on number of deferrals applied for the protocol will be reinstate once the functionality of the deferral mechanism is restored.

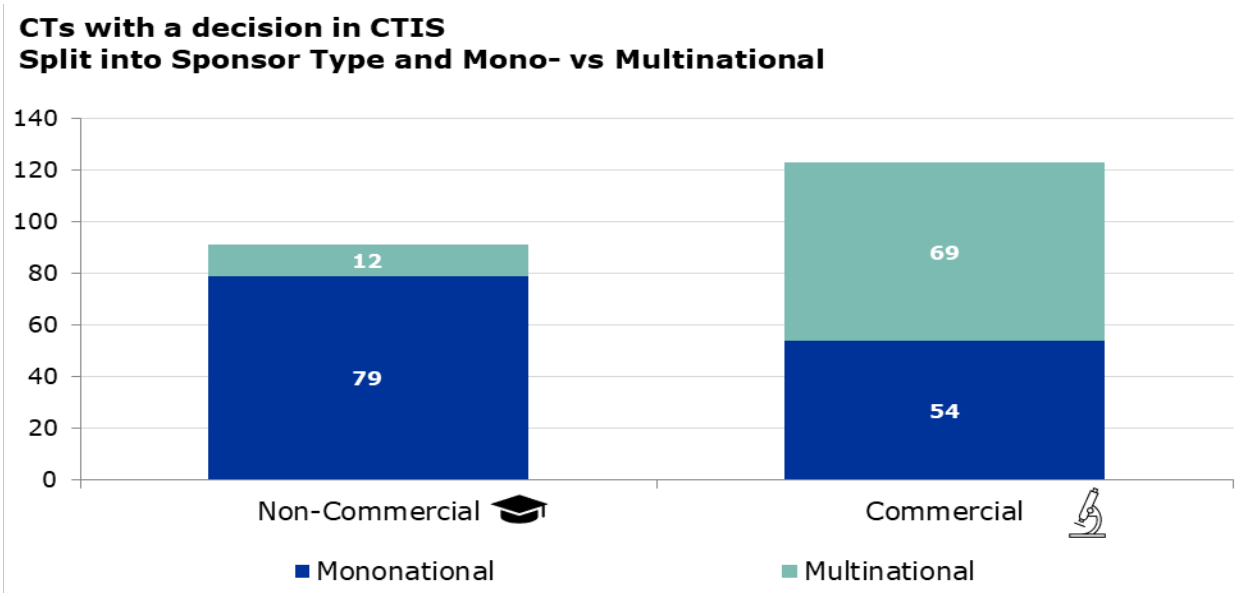
[7] The data for January 2022 in the graph refers to CTA uploaded by the Member State on the 31 January 2022 only.
[8] Details on recruitment status are based on the information reported by the trial sponsor in CTIS
[9] The option to defer the protocol is only available in CTIS.

1.5. Number of mononational-multinational trials for which a decision has been issued by the Member States Concerned (MSC) under the Clinical Trials Regulation, broken down per sponsor type (commercial vs. non-commercial) and average number of MSCs[10]

The graph below shows the number of trials for which a decision has been issued in CTIS by the Member State Concerned, per month, since 31 January 2022. By end December 2022 the cumulative total rose from 174 to a total of 214 clinical trials, which have been authorised, authorised with conditions and not authorised. Out of the total cumulative clinical trials, 5 were reported as ended.



The graph below shows the number of clinical trials for which a decision has been issued by the first MSC, with information whether the trial is a mono- or multinational and in relation to sponsor type.



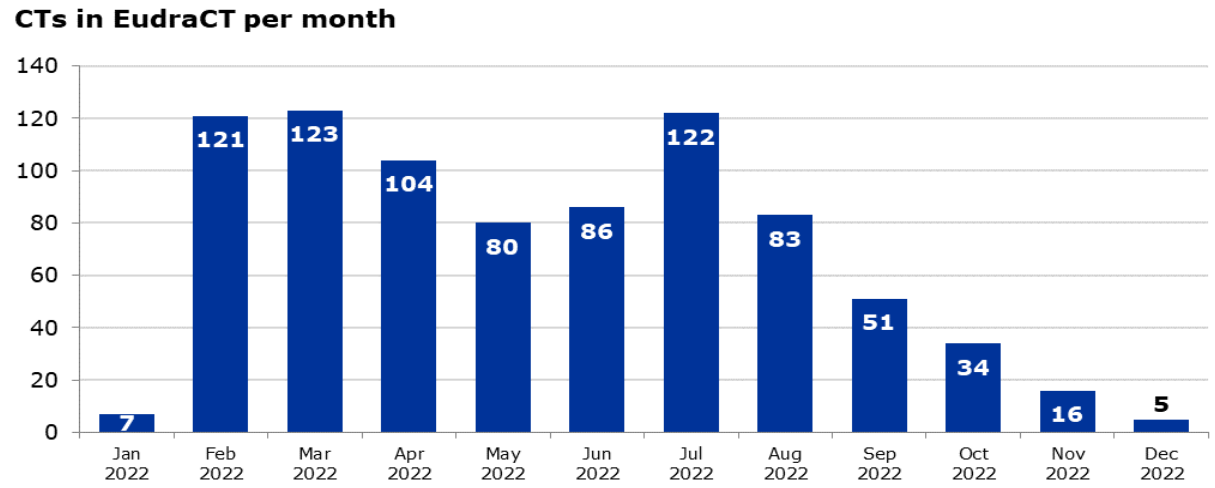
Currently 81 multinational clinical trial have a decision in CTIS with an average of 6 Member States Concerned.

[10] Details on trial sponsor type, commercial *vs* non-commercial are derived based on information reported at the time of registration of an organisation in OMS: Organisation Management Service database, and are not recorded as such in the clinical trial application form. Commercial classification includes for example industry, pharmaceutical company, while non-commercial classification includes values such as academia, health care facility, micro, small and medium enterprises.

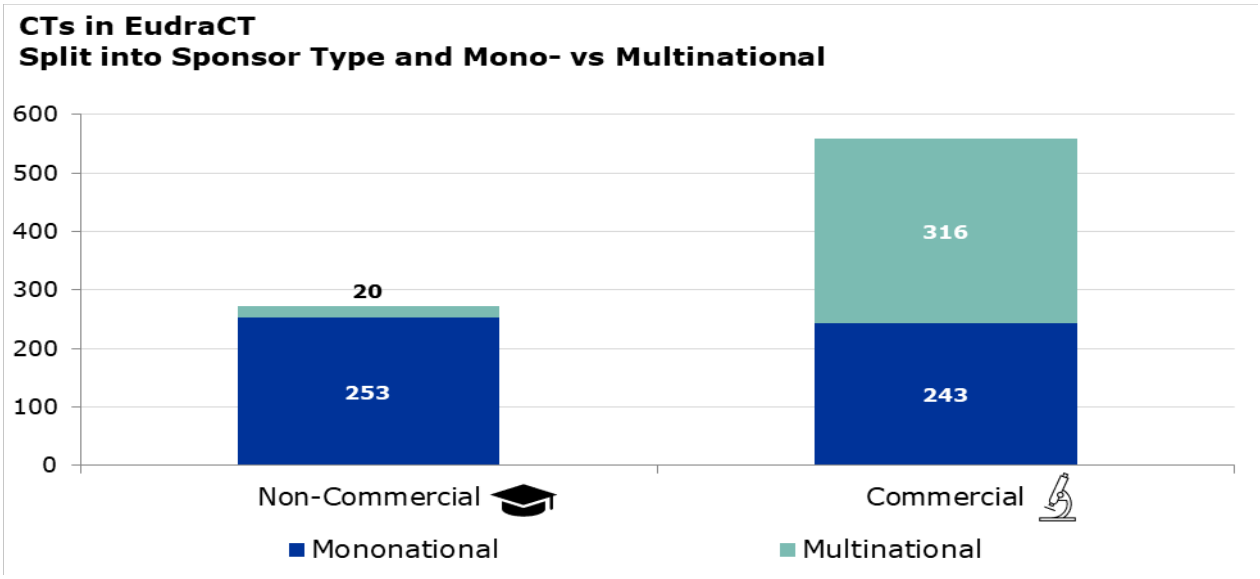
1.6. Number of mononational-multinational trials for which a NCA decision and an Ethics Committee opinion have been issued by the Member States under the Clinical Trials Directive, broken down per sponsor type (commercial vs. non-commercial) and average number of MSs

The graph below shows the number of trials, as individual clinical trials protocols, that received a National Competent Authority decision and an Ethics Committee opinion from the first Member State uploading the CTA in EudraCT, per month, since 31 January 2022[11] displayed by upload date in EudraCT.

The numbers of applications with a NCA decision and Ethics Committee opinion may increase overtime, as soon as additional information is provided in EudraCT by the Member States. The cumulative total rose from 768 to a total of 832 clinical trials by end December 2022.



The graph below shows the number of clinical trials for which a decision and an opinion have been issued, with information whether the trial is a mono- or multinational and in relation to sponsor type.

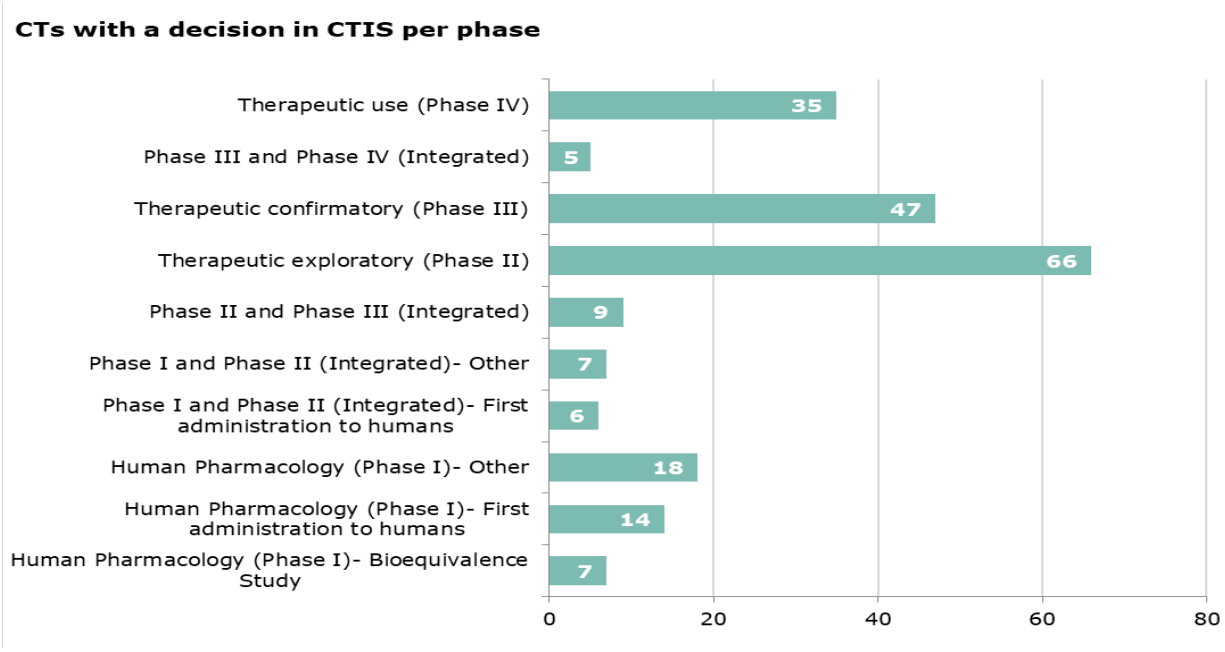


Considering clinical trials for which a decision and an opinion have been issued, on average 3 Member States are involved in multinational trials.

[11] The data for January that appear in the graph below refers to CTA loaded in EudraCT the 31st January having a subsequent decision by the national Competent Authority and Ethic Committee opinion loaded in EudraCT.

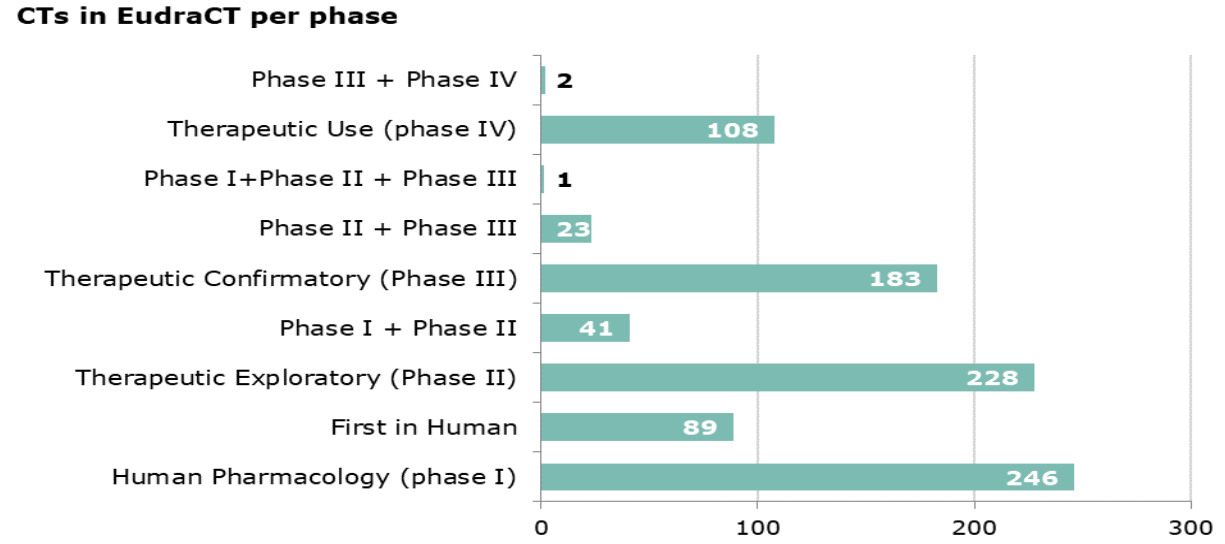
1.7. Number of clinical trials for which a decision has been issued per phase (i.e. I, II, III, IV, as well as CT first in human or combined phases early (I and II)) under CTR[12]

The graph below shows the number of clinical trials for which a decision has been issued, broken down per trial phase.



1.8. Number of clinical trials for which a NCA decision and an Ethics Committee opinion have been issued per phase (i.e. I, II, III, IV, as well as CT first in human or combined phases early (I and II)) under CTD

The graph below shows the number of clinical trials, as individual clinical trial protocol, uploaded in EudraCT as of 31 January 2022, for which a decision by the National Competent Authority and an opinion by the Ethics Committee have been inserted by the first Member States uploading the CTA in EudraCT, broken down per trial phase.

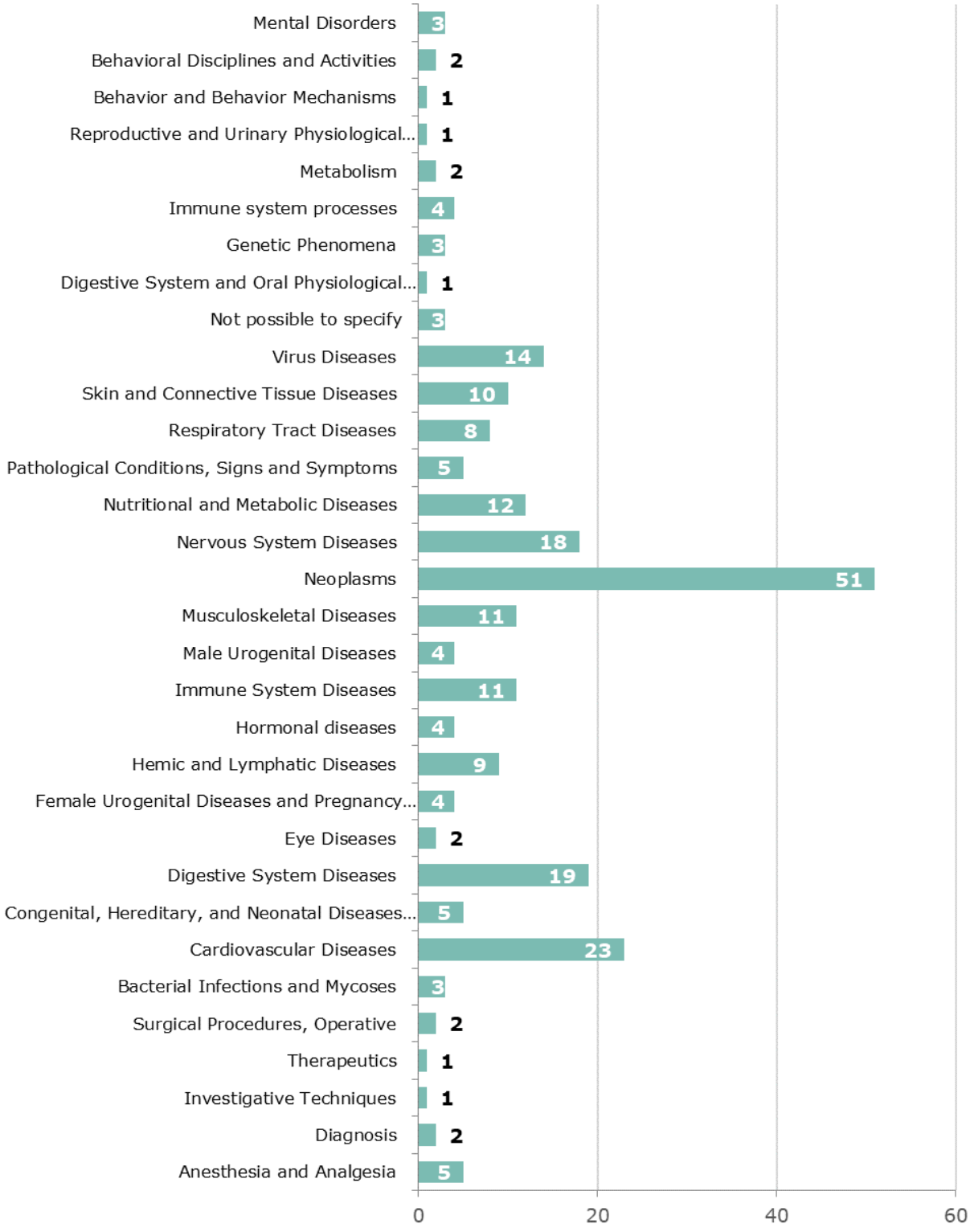


[12] More than one trial phase can be selected for a single trial and it is counted in each trial. The graph shows the applicable trial phases in the selected period.

1.9. Number of trials for which a decision has been issued under CTR, per therapeutic area[13]

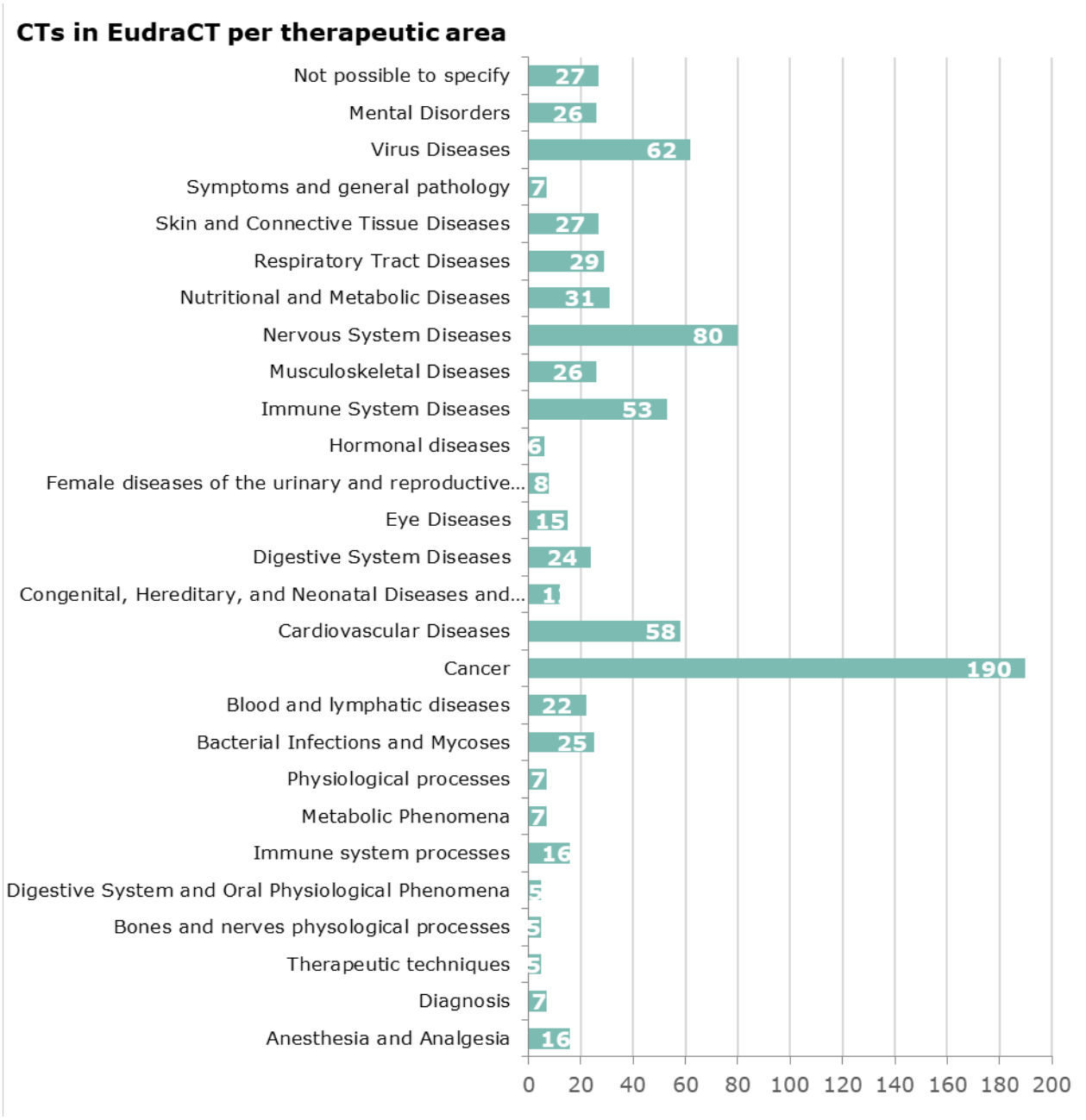
The graph below shows the number of clinical trials for which a decision has been issued in CTIS, broken down per therapeutic area.

CTs with a decision in CTIS per therapeutic area



1.10. Number of trials for which a NCA decision and an Ethics Committee opinion have been issued under CTD, per therapeutic area[14]

The graph below shows the number of clinical trials, as individual clinical trial protocol, for which a decision by the National Competent Authority and an opinion by the Ethics Committee have been inserted by the first Member State uploading the CTA in EudraCT, broken down per therapeutic area.[15]



[13] In case a clinical trial investigates several therapeutic areas, it is counted in each of such identified therapeutic areas.

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[14] In case a clinical trial investigates several therapeutic areas, it is counted in each of such identified therapeutic areas.
. The graph shows the applicable trials therapeutic areas in the selected period displaying only therapeutic areas selected in 5 or more clinical trials, but not less.

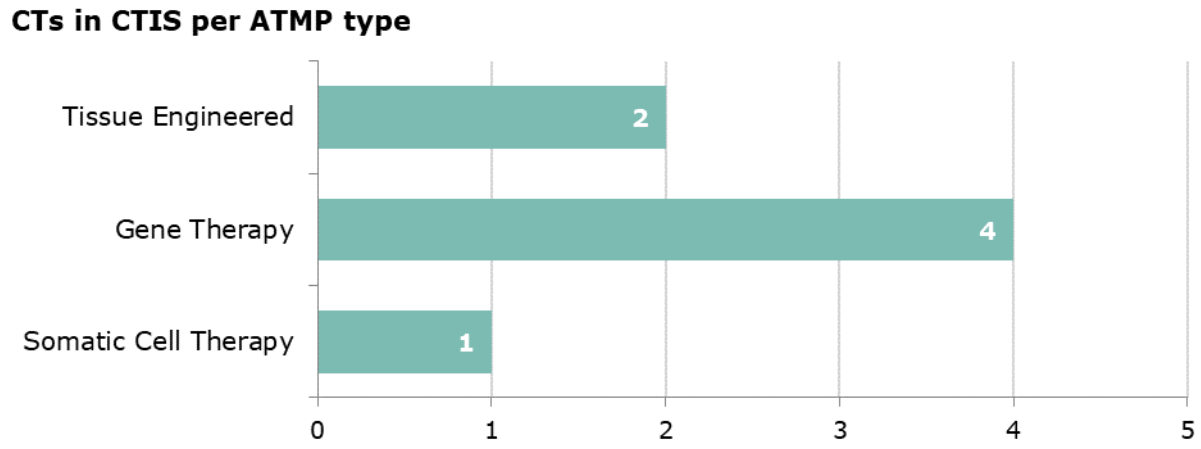
[15] The value 'not possible to specify' in the graph above reflects the fact that section E.1.1.2 of the CTA was not filled in

1.11. Number of trials for which a decision has been issued on Advanced Therapy Medicinal Products (ATMP) under CTR

Seven clinical trials for which a decision has been issued in CTIS include an Advanced Therapy Medicinal Product.

1.12. Number of trials for which a decision has been issued, with ATMP of type "gene therapy", "somatic cell therapy" and "tissue engineered therapy" under CTR

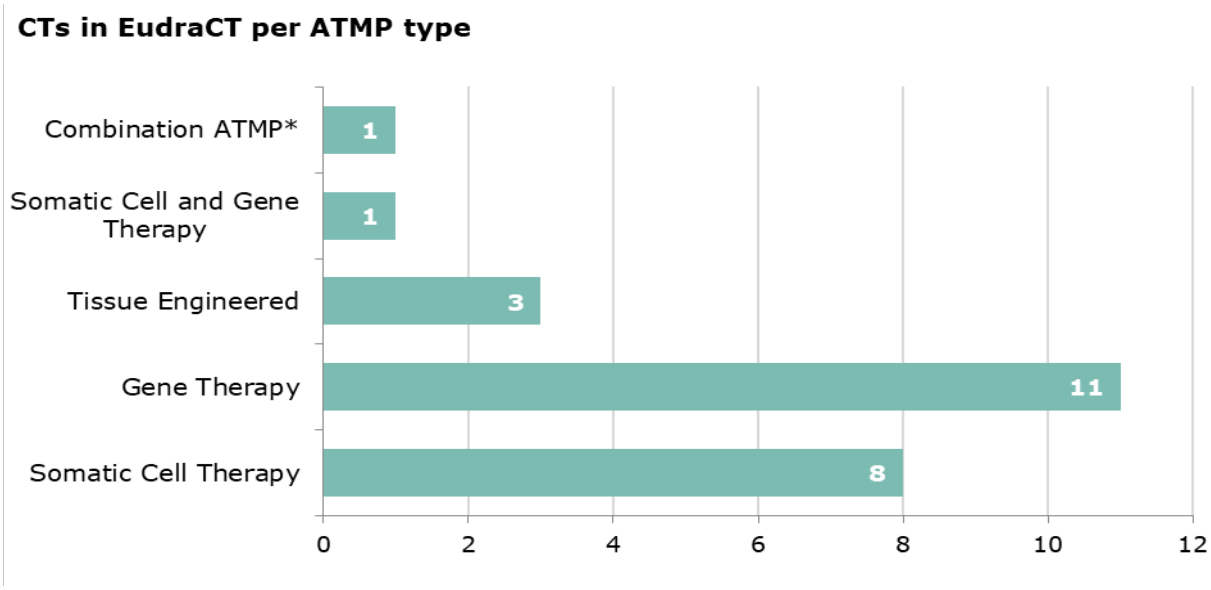
The graph below shows the ATMP type for the 7 clinical trials for which a decision has been issued in CTIS, which include an Advanced Therapy Medicinal Product.



1.13. Number of trials for which a NCA decision and an Ethics Committee opinion have been issued, with ATMP of type "gene therapy", "somatic cell therapy" and "tissue engineered therapy" under CTD

There were 24 clinical trials in EudraCT since 31 January 2022, with a decision and an opinion issued by 31 December 2022 by the first Member State uploading the CTA in EudraCT, including an advanced therapy medicinal products.

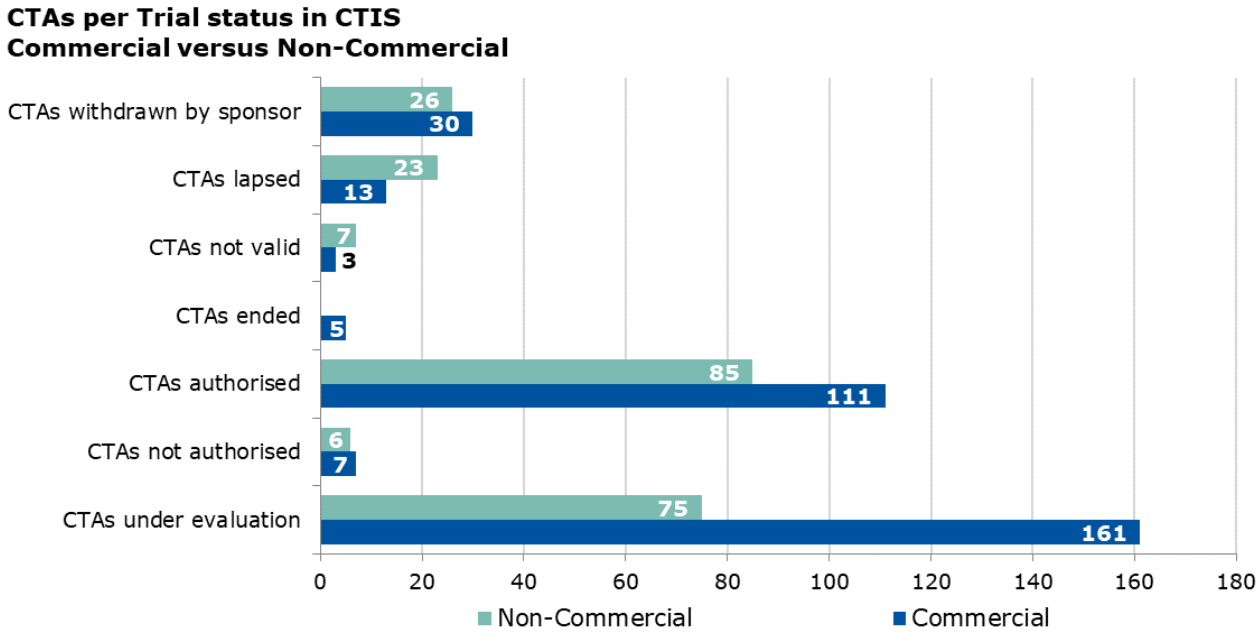
The graph below shows the number of clinical trials per ATMP type as reported in EudraCT.



* Combination ATMP with a medical device.

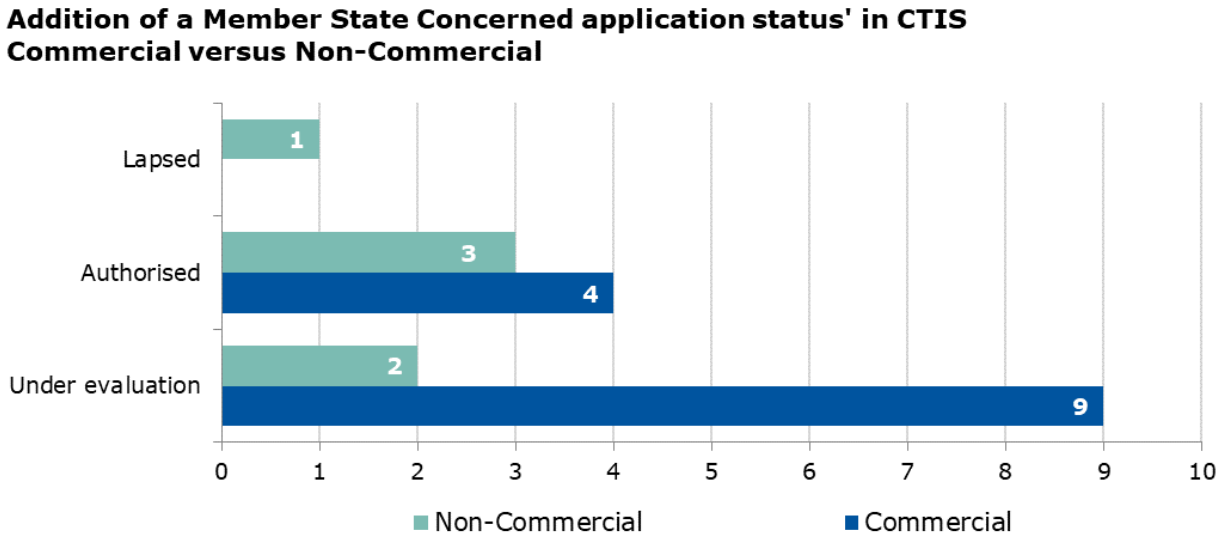
1.14. Number of clinical trial applications under the CTR per applicable trial status during the selected period, broken down per sponsor type: non- commercial/commercial

The graph below shows the number of initial clinical trial applications, per applicable overall trial status[16] and information of sponsor type submitted in CTIS since 31 January 2022.



1.15. Art 14 applications to add a new Member State Concerned: (re-)submission, authorisation, rejection, lapsed and withdrawn dossiers

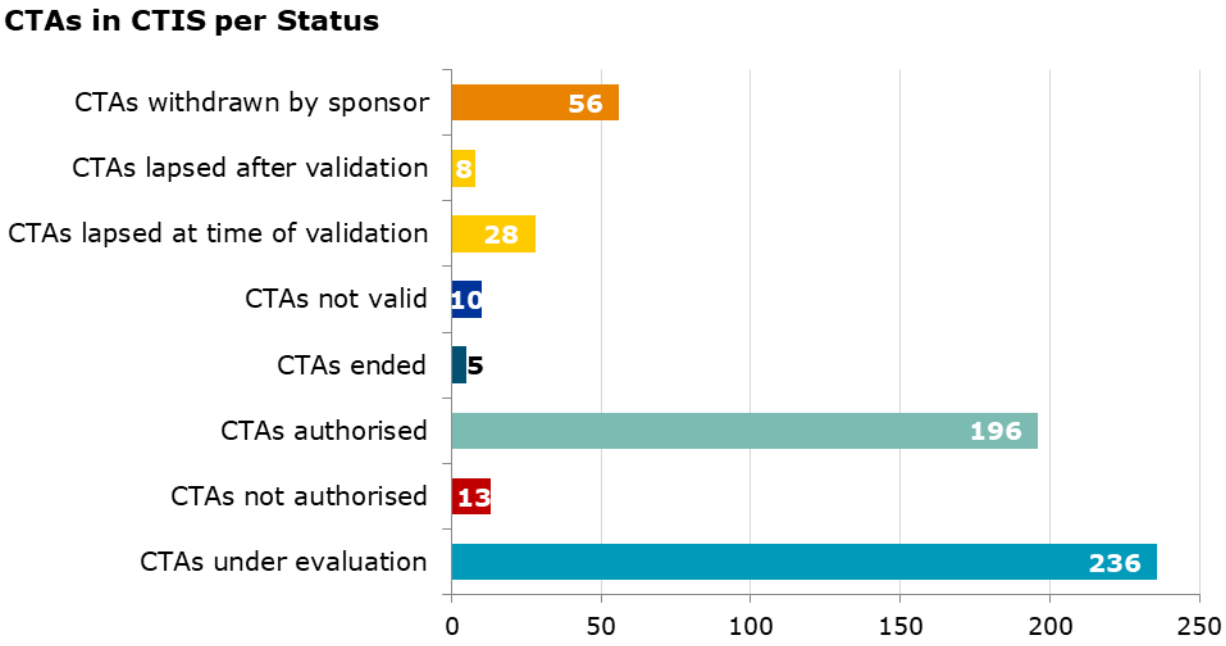
As of 31 December 2022 19 clinical trial applications for the addition of a new MSC, foreseen under Article 14 of Regulation (EU) No 536/2014, have been submitted in CTIS for 12 clinical trials. The below graph provides overview status per application submitted until 31 December 2022.



[16] Overall trial status is the status per application and not per individual Member State Concerned.

1.16. Number of CTAs Article 5 of CTR [full dossier initial applications] per applicable trial status during the reporting period, at EU, at MS level and with Reporting Member State (RMS) details

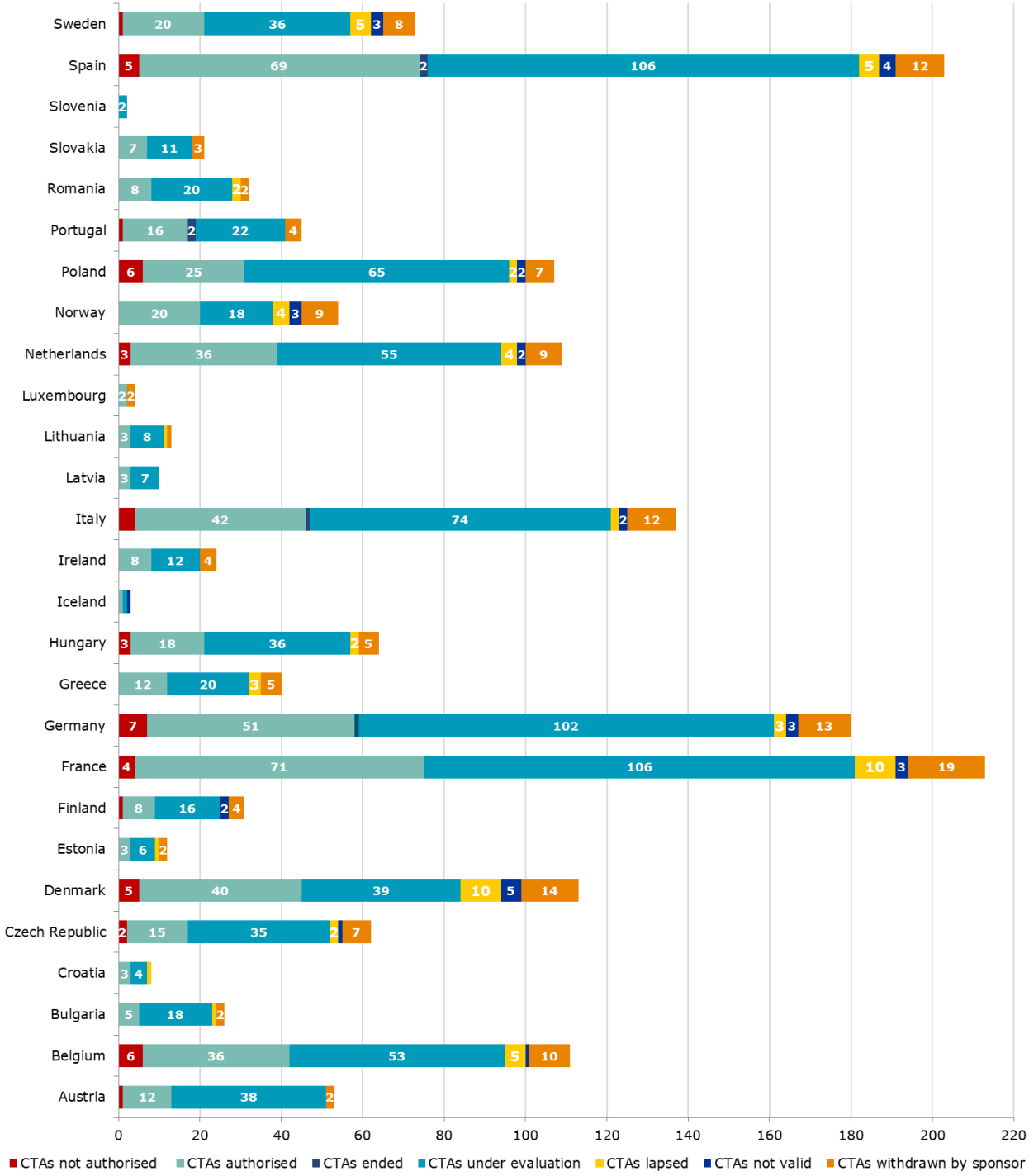
The graph below shows the number of initial clinical trial applications with full dossier, submitted in accordance with Article 5 of CTR since 31 January 2022, per applicable overall status at EU level.



It can be noted an increase of 16% of clinical trial applications under evaluation, and authorisation of clinical trial applications have increased by 22% compared to the previous reporting period.

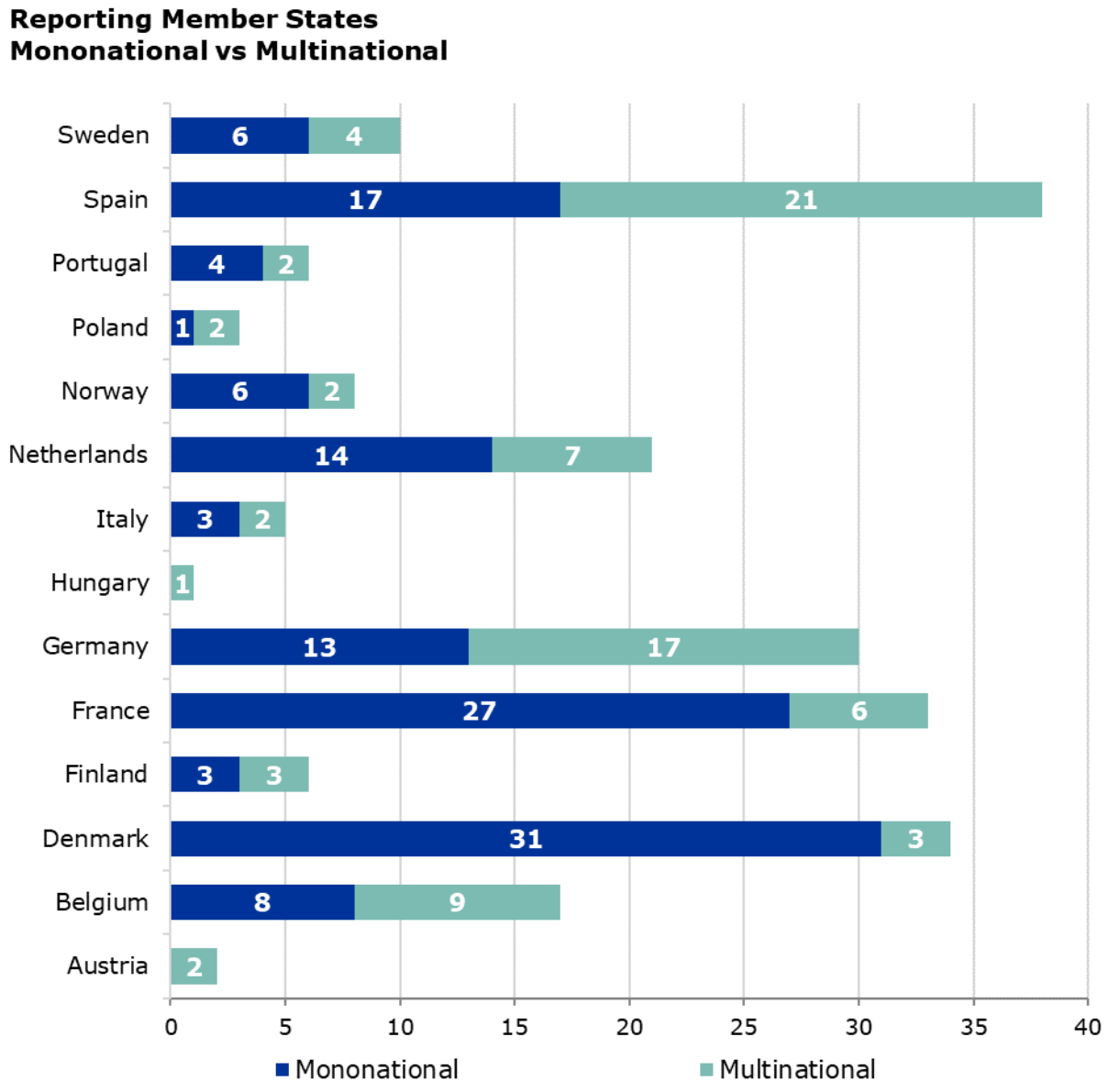
The graph below shows the number of initial clinical trial applications with full dossier, submitted in accordance with Article 5 of CTR since 31 January 2022, per applicable status at the level of the Member States Concerned[17].

Member States Concerned



[17] In multinational clinical trials the same application has been submitted to multiple Member State Concerned, and it is counted in the graph in each applicable MSC.

The graph below shows the distribution of appointment of Reporting Member State (RMS)[18], amongst the applicable Member States Concerned, for clinical trial applications on which a decision has been issued for mono- and multinational trials.



1.17. Number of CTA Article 11 of CTR [partial dossier initial applications with later Part II submission] per applicable trial status during the reporting period, at EU and at MS level

Partial initial applications submitted to CTIS in line with the requirements of Article 11 of the Regulation (EU) No 536/2014 will be considered for future reporting.

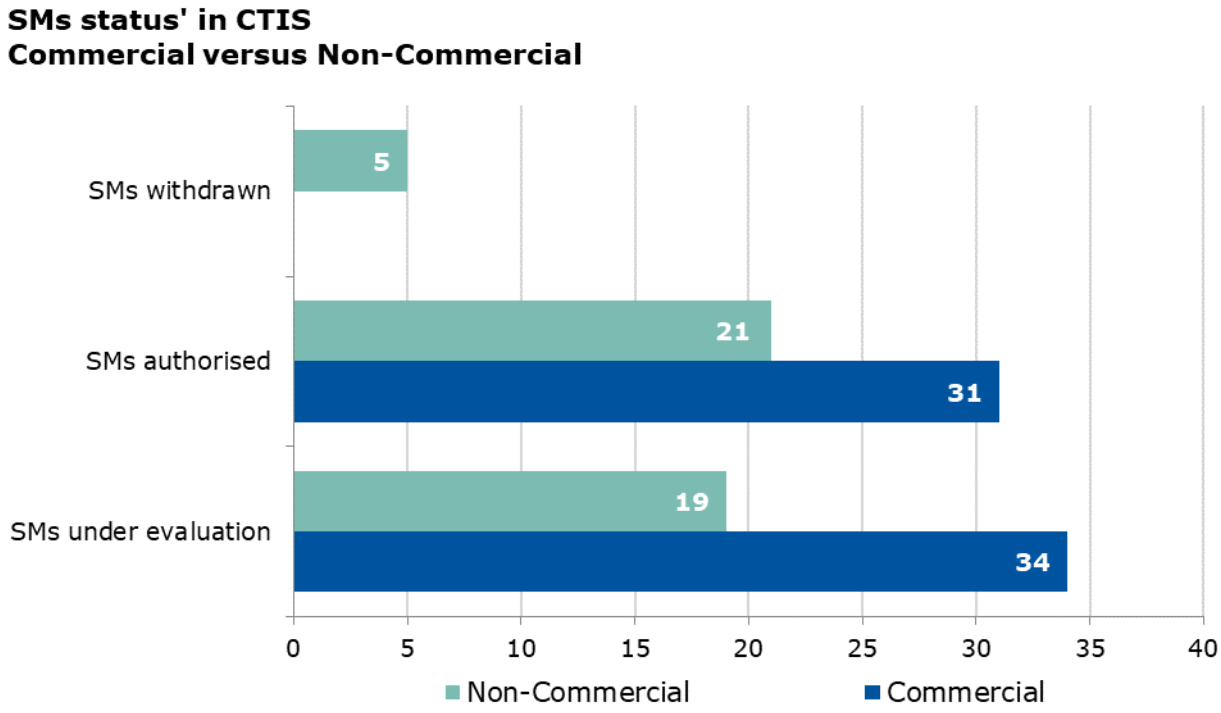
[18] RMS is the Reporting Member State appointed in line with the requirements of Article 5 of the Clinical Trials Regulation (EU) No 536/2014

1.18. Average time from submission to reporting date[19] (Article 11 and Article 5 of CTR), and to first decision (Article 5 of CTR) for initial applications and Substantial Modifications part I or part I and II

On average it took 84 calendar days to issue a decision, during the selected period, for the 214 initial clinical trial applications. More details can be found in Annex I.

1.19. Number of submitted, validated, authorised, rejected, lapsed and withdrawn Substantial Modification (SM) applications, related to part I / II / I and II, by sponsor type

As of 31 December 2022 110 distinct applications for substantial modifications, foreseen in chapter II of Regulation (EU) No 536/2014, were submitted in CTIS for 72 clinical trials.



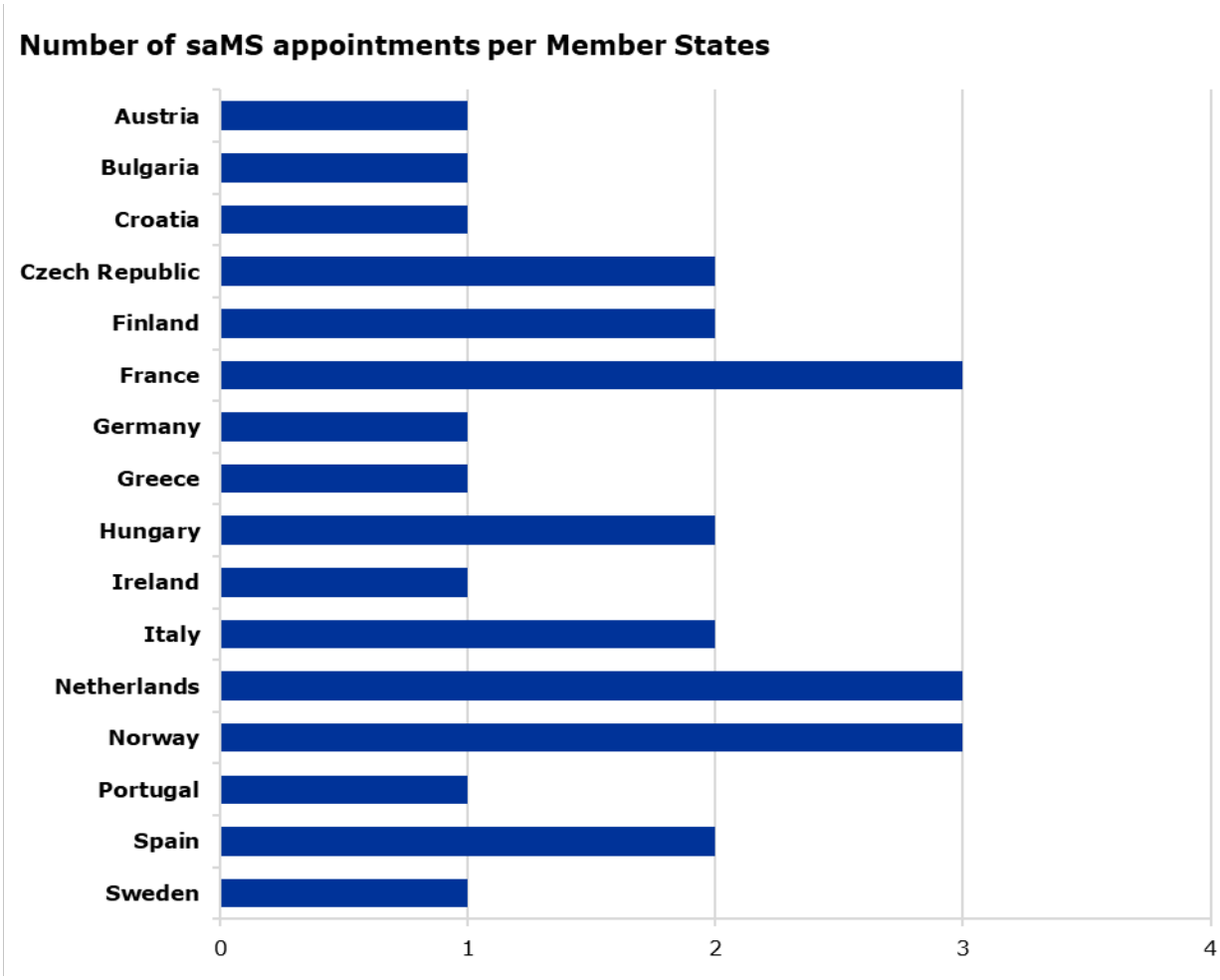
1.20. Number of active substances (ASs) in CTR EU trials (mononational and multinational AS)

During the reporting period saMS[20] were appointed for 27 active substances.

1.21. Number of safety assessing Member State (saMS)-ships per MS

The role of safety assessing Member State Concerned (saMS) will be applicable only for active substances investigated in clinical trials in two or more MSC. During the selected reporting period, 16 saMSs have been appointed for the active substances listed in point 1.20 above.

[19] The reporting date is equal to the date of the RMS conclusion on part I assessment
[20] Safety Assessing Member State


Number of saMS appointments per Member States
Austria
Bulgaria
Croatia
Czech Republic
Finland
France
Germany
Greece
Hungary
Ireland
Italy
Netherlands
Norway
Portugal
Spain
Sweden
0
1
2
3
4

Annex I
Average time from submission to decision for initial CTAs

The table below shows the number of calendar days since the submission of the initial clinical trial application to CTIS up to the time of the first decision of the Member States Concerned.

Please consider that due dates for tasks completion, including decision, in CTIS takes into account rules such as: allowing 2 consecutive working days, the due date cannot fall on weekend nor on a bank holiday in addition to counting the calendar days.

The list of clinical trials below includes resubmitted applications.

#	Submission date	Decision date	Days to Decision
1	09/02/2022	07/04/2022	57.00
2	15/02/2022	02/06/2022	107.00
3	15/02/2022	27/04/2022	71.00
4	20/02/2022	30/05/2022	99.00
5	24/02/2022	16/06/2022	112.00
6	28/02/2022	28/03/2022	28.00
7	03/03/2022	27/06/2022	116.00
8	04/03/2022	31/05/2022	88.00
9	04/03/2022	21/06/2022	109.00
10	07/03/2022	28/06/2022	113.00
11	10/03/2022	19/05/2022	70.00
12	15/03/2022	27/04/2022	43.00
13	16/03/2022	27/06/2022	103.00
14	24/03/2022	30/06/2022	98.00
15	24/03/2022	14/06/2022	82.00
16	28/03/2022	19/07/2022	113.00
17	28/03/2022	02/11/2022	219.00
18	29/03/2022	22/06/2022	85.00
19	31/03/2022	26/07/2022	117.00
20	01/04/2022	01/06/2022	61.00
21	05/04/2022	18/07/2022	104.00
22	06/04/2022	12/07/2022	97.00
23	08/04/2022	28/06/2022	81.00
24	14/04/2022	28/07/2022	105.00
25	14/04/2022	28/07/2022	105.00
26	18/04/2022	27/05/2022	39.00
27	19/04/2022	03/08/2022	106.00
28	19/04/2022	05/07/2022	77.00
29	21/04/2022	28/07/2022	98.00
30	22/04/2022	28/07/2022	97.00
31	22/04/2022	05/08/2022	105.00
32	22/04/2022	08/07/2022	77.00
33	22/04/2022	01/08/2022	101.00

#	Submission date	Decision date	Days to Decision
34	26/04/2022	28/06/2022	63.00
35	28/04/2022	24/08/2022	118.00
36	28/04/2022	16/08/2022	110.00
37	29/04/2022	05/08/2022	98.00
38	29/04/2022	08/08/2022	101.00
39	02/05/2022	07/06/2022	36.00
40	03/05/2022	24/08/2022	113.00
41	04/05/2022	04/07/2022	61.00
42	05/05/2022	01/08/2022	88.00
43	05/05/2022	08/07/2022	64.00
44	06/05/2022	23/06/2022	48.00
45	06/05/2022	28/10/2022	175.00
46	08/05/2022	04/07/2022	57.00
47	09/05/2022	18/07/2022	70.00
48	10/05/2022	04/08/2022	86.00
49	10/05/2022	26/07/2022	77.00
50	12/05/2022	31/08/2022	111.00
51	12/05/2022	01/09/2022	112.00
52	13/05/2022	26/08/2022	105.00
53	17/05/2022	09/09/2022	115.00
54	18/05/2022	01/09/2022	106.00
55	18/05/2022	28/06/2022	41.00
56	20/05/2022	08/07/2022	49.00
57	20/05/2022	25/08/2022	97.00
58	23/05/2022	28/10/2022	158.00
59	25/05/2022	11/08/2022	78.00
60	25/05/2022	16/09/2022	114.00
61	30/05/2022	06/09/2022	99.00
62	30/05/2022	22/06/2022	23.00
63	31/05/2022	23/09/2022	115.00
64	02/06/2022	10/08/2022	69.00
65	03/06/2022	21/09/2022	110.00
66	03/06/2022	26/09/2022	115.00
67	03/06/2022	01/08/2022	59.00
68	03/06/2022	01/08/2022	59.00
69	03/06/2022	27/09/2022	116.00
70	07/06/2022	27/09/2022	112.00
71	08/06/2022	27/09/2022	111.00
72	09/06/2022	23/08/2022	75.00
73	09/06/2022	11/08/2022	63.00
74	13/06/2022	09/08/2022	57.00
75	14/06/2022	15/09/2022	93.00

#	Submission date	Decision date	Days to Decision
76	15/06/2022	12/08/2022	58.00
77	16/06/2022	09/09/2022	85.00
78	19/06/2022	03/08/2022	45.00
79	22/06/2022	12/08/2022	51.00
80	22/06/2022	08/07/2022	16.00
81	23/06/2022	26/09/2022	95.00
82	23/06/2022	15/09/2022	84.00
83	24/06/2022	12/09/2022	80.00
84	24/06/2022	12/09/2022	80.00
85	24/06/2022	03/10/2022	101.00
86	24/06/2022	20/10/2022	118.00
87	27/06/2022	12/10/2022	107.00
88	28/06/2022	10/10/2022	104.00
89	28/06/2022	14/10/2022	108.00
90	01/07/2022	13/10/2022	104.00
91	01/07/2022	07/10/2022	98.00
92	04/07/2022	13/10/2022	101.00
93	04/07/2022	12/09/2022	70.00
94	05/07/2022	26/09/2022	83.00
95	08/07/2022	26/10/2022	110.00
96	09/07/2022	26/09/2022	79.00
97	11/07/2022	23/08/2022	43.00
98	12/07/2022	28/07/2022	16.00
99	12/07/2022	24/10/2022	104.00
100	13/07/2022	10/11/2022	120.00
101	13/07/2022	18/10/2022	97.00
102	14/07/2022	20/10/2022	98.00
103	15/07/2022	08/11/2022	116.00
104	15/07/2022	24/10/2022	101.00
105	15/07/2022	19/10/2022	96.00
106	15/07/2022	14/10/2022	91.00
107	15/07/2022	12/09/2022	59.00
108	17/07/2022	08/11/2022	114.00
109	18/07/2022	11/10/2022	85.00
110	18/07/2022	24/10/2022	98.00
111	18/07/2022	10/10/2022	84.00
112	18/07/2022	29/09/2022	73.00
113	19/07/2022	03/10/2022	76.00
114	20/07/2022	22/09/2022	64.00
115	20/07/2022	27/10/2022	99.00
116	20/07/2022	03/10/2022	75.00
117	21/07/2022	03/11/2022	105.00

#	Submission date	Decision date	Days to Decision
118	21/07/2022	09/11/2022	111.00
119	21/07/2022	07/10/2022	78.00
120	21/07/2022	26/07/2022	5.00
121	25/07/2022	14/11/2022	112.00
122	25/07/2022	12/10/2022	79.00
123	26/07/2022	24/10/2022	90.00
124	27/07/2022	09/11/2022	105.00
125	28/07/2022	03/10/2022	67.00
126	28/07/2022	19/09/2022	53.00
127	29/07/2022	10/11/2022	104.00
128	01/08/2022	21/11/2022	112.00
129	02/08/2022	23/11/2022	113.00
130	02/08/2022	12/10/2022	71.00
131	03/08/2022	31/10/2022	89.00
132	03/08/2022	14/10/2022	72.00
133	03/08/2022	14/11/2022	103.00
134	04/08/2022	12/10/2022	69.00
135	04/08/2022	02/11/2022	90.00
136	05/08/2022	16/09/2022	42.00
137	05/08/2022	25/10/2022	81.00
138	09/08/2022	29/11/2022	112.00
139	10/08/2022	09/11/2022	91.00
140	10/08/2022	27/10/2022	78.00
141	11/08/2022	06/12/2022	117.00
142	11/08/2022	06/12/2022	117.00
143	11/08/2022	28/11/2022	109.00
144	12/08/2022	06/12/2022	116.00
145	12/08/2022	29/11/2022	109.00
146	12/08/2022	21/11/2022	101.00
147	12/08/2022	06/12/2022	116.00
148	12/08/2022	05/12/2022	115.00
149	12/08/2022	25/11/2022	105.00
150	15/08/2022	20/10/2022	66.00
151	16/08/2022	29/11/2022	105.00
152	16/08/2022	06/12/2022	112.00
153	17/08/2022	06/12/2022	111.00
154	18/08/2022	15/11/2022	89.00
155	19/08/2022	08/11/2022	81.00
156	19/08/2022	13/12/2022	116.00
157	19/08/2022	06/12/2022	109.00
158	19/08/2022	17/11/2022	90.00
159	22/08/2022	27/10/2022	66.00

#	Submission date	Decision date	Days to Decision
160	22/08/2022	24/11/2022	94.00
161	22/08/2022	22/11/2022	92.00
162	23/08/2022	06/12/2022	105.00
163	25/08/2022	20/12/2022	117.00
164	25/08/2022	15/12/2022	112.00
165	25/08/2022	15/11/2022	82.00
166	25/08/2022	09/12/2022	106.00
167	26/08/2022	11/11/2022	77.00
168	26/08/2022	25/10/2022	60.00
169	26/08/2022	18/11/2022	84.00
170	26/08/2022	10/11/2022	76.00
171	29/08/2022	07/11/2022	70.00
172	29/08/2022	01/12/2022	94.00
173	31/08/2022	19/12/2022	110.00
174	31/08/2022	19/12/2022	110.00
175	31/08/2022	28/11/2022	89.00
176	31/08/2022	20/12/2022	111.00
177	01/09/2022	11/11/2022	71.00
178	09/09/2022	22/12/2022	104.00
179	10/09/2022	28/11/2022	79.00
180	12/09/2022	02/12/2022	81.00
181	13/09/2022	19/12/2022	97.00
182	14/09/2022	05/12/2022	82.00
183	15/09/2022	14/11/2022	60.00
184	16/09/2022	14/12/2022	89.00
185	17/09/2022	06/12/2022	80.00
186	19/09/2022	24/11/2022	66.00
187	19/09/2022	29/11/2022	71.00
188	21/09/2022	16/11/2022	56.00
189	21/09/2022	14/11/2022	54.00
190	21/09/2022	04/11/2022	44.00
191	23/09/2022	06/10/2022	13.00
192	26/09/2022	06/12/2022	71.00
193	27/09/2022	22/12/2022	86.00
194	29/09/2022	22/11/2022	54.00
195	30/09/2022	14/12/2022	75.00
196	03/10/2022	03/11/2022	31.00
197	03/10/2022	14/11/2022	42.00
198	05/10/2022	22/12/2022	78.00
199	10/10/2022	15/11/2022	36.00
200	10/10/2022	30/11/2022	51.00
201	12/10/2022	19/12/2022	68.00

#	Submission date	Decision date	Days to Decision
202	17/10/2022	02/12/2022	46.00
203	20/10/2022	21/12/2022	62.00
204	26/10/2022	05/12/2022	40.00
205	28/10/2022	14/12/2022	47.00
206	31/10/2022	23/12/2022	53.00
207	31/10/2022	15/12/2022	45.00
208	01/11/2022	18/11/2022	17.00
209	03/11/2022	20/12/2022	47.00
210	04/11/2022	20/12/2022	46.00
211	09/11/2022	18/12/2022	39.00
212	15/11/2022	15/12/2022	30.00
213	23/11/2022	16/12/2022	23.00
214	08/12/2022	20/12/2022	12.00