Published Standard Number 1 – Applications (National)

Application number	Application type	Number of applications	Performance	Target days	Average days
1	Major timetable (National) New MRLs. All other MA applications (excl. MAPI and Copycats)	5	100%	180.0	19
2	Standard timetable (National Type II variations. New MA - MAPIs and Copycats. New VHRs)	20	100%	120.0	12
3	Shortened timetable (National Renewals (MA and VHR) Type IB variations. New ATC (type B). Out of Scope MRLs)	129	99.2%	60.0	6
4	Minor timetable (National) Type IA variations. Administrative Type IB variations. New ATC (Type A/S). ATC variations and renewals.	245	97.6%	30.0	20
5	Parallel Assessment with EU Procedures	295	100%	-	12
6	Shared Assessment with International Partners	0	-	-	0
7	Batch timetable (National) specific Batch Control	16	100%	20.0	1
8	Autogenous Vaccines. New & Variations	2	100%	45.0	44

Published Standard Number 1 – Applications (Other)

Application number	Application Type	Number of applications	Performance
9	Mock-up period completed within 20 days (or up to 40 days for parallel applications involving different QRD sources)	289	96.5%
10	Validation	712	100%
11	Issue of authorised documentation	923	100%

Published Standard Number 1 – Applications (European - NI)

Application number	Application Type	Number of applications	Performance
12	New Decentralised (DCP)	16	100%
13	New Mutual Recognition (MRP)	2	100%
14	MRP Variations (Type IB & II) and Renewals	162	100%

Published Standard Number 2 – Public Assessment Reports

Application number	Application type	Total number	Performance
15	Publishing Summary of Product Characteristics (SPCs)	30	96.7%
16	Publishing Public Assessment Reports (PuARs)	0	-
17	Updating PuARs	3	100%

Published Standard Number 3 – Quality of Documentation

Application number	Application type	Number of applications	Performance
18	Unreturned Documents	1541	97.8%

Published Standard Number 4 – Product Defects

Task number	Task	Number of tasks	Performance	Target Days	Average Days
19	Product Defects reports	22	100%		
	High risk <5 days	1	-		
	Low risk <10 days	21	-		

Published Standard Number 5 – Import, Export and Batch Release Schemes

Application number	Application Type	No of Apps	Performance	Target Days	Average Days
20	Applications for new pharmaceutical products	37	100%	15	-
21	Applications for new Immunological products	5	100%	25	-
22	Applications for previously imported products	119	100%	15	-
23	All other urgent applications	264	99.6%	2	-
	Urgent	0			
	Non Urgent	264			
24	Instant Import Certificates	14,274	-	-	-
25	Export	137	100%	10	6.5
26	Batch Release	1188	100%	10	3.7

Published Standard Number 6 – Pharmacovigilance

Task number	Task	No.	Performance
27	Human, Animal & Environmental AERs	3740	99.8%
28	PSURs	634	100%
29	Inspections	9	100%

Published Standard Number 7– Inspections

Task number	Task	No.	Performance	Target Days
30	Inspections within 3 years (GMP)	14	100%	-
	Within 5 years (GDP) of last inspection	11	Joint with above	-
31	Inspection Deficiency Reports	26	100%	30.0
32	(GMP) Certificates or (GDP) final reports sent	29	100%	90.0
33	Approval of new Feed business operators and SQP retailer sites	25	100%	45.0
34	Final inspection report to Feed business operators and SQP retailers	175	99.2%	30.0

Our inspection procedures enable us to extend our GMP inspections beyond 3 years and our GDP inspections beyond 5 years where there are exceptional circumstances, provided a documented risk-assessment is carried out. Risk-assessments have been conducted for all sites where it has not been possible for us to inspect them within 3 years due to covid-19 related restrictions.

Key:

100% Excellent

>97% - 100% Excellent, but some targets missed

92% - 97% Effective

< 91% Ineffective

Additional information

The VMD continuously monitors all targets and puts in place countermeasures, where possible, to ensure targets are met.

However, sometimes a performance standard may fall into the effective or ineffective category and there are a number of reasons why this may happen, e.g. high volume of applications, staff resource, complexity of applications requiring additional input, etc.