Published Standard Number 1 – Applications (National)

Application number	Application type	Number of applications	Performance	Target days
1	Complex timetable (National new MA applications) Complex new MA applications, e.g. novel therapies, new actives	59	100%	210.0
2	Major timetable (National) New MA applications (excl. MAPI and Copycats)	2	100%	180.0
3	Standard timetable (National Type II VRA. New MA - Copycats. New VHRs)	293	100%	120.0
4	Shortened timetable (Type IB VRA. New ATC (type B). Out of Scope MRLs)	213	99.5%	60.0
5	Minor timetable (National) Type IA VNRA. Administrative Type IB VRA. New ATC (Type A/S). ATC variations.	1426	99%	30.0
6	Parallel Assessment with EU Procedures	629	99.7%	-
7	Batch timetable (National) specific Batch Control	48	100%	20.0
8	Autogenous Vaccines. New & Variations	4	100%	45.0

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)	888	99%
10	Validation	1494	96.1%
11	Issue of authorised documentation	3113	99.8%

Published Standard Number 1 – Applications (European- NI)

Application number	Application Type	Number of applications	Performance
12	New Decentralised (DCP)	21	100%
13	New Mutual Recognition (MRP)	1	100%
14	MRP Variations (Type IB & II)	349	99.7%

Published Standard Number 2 – Public Assessment Reports

Application number	Application type	Total number	Performance
15	Publishing Summary of Product Characteristics (SPCs)	112	100%
16	Publishing Public Assessment Reports (PuARs)	89	100%
17	Updating PuARs	3	100%

Published Standard Number 3 – Quality of Documentation

Application number	Application type	Number of applications	Performance
18	Unreturned Documents	6285	95.5%

Published Standard Number 4 – Product Defects

Task number	Task	Number of tasks	Performance
19	Product Defects reports	58	100%
	High risk <5 days	13	-
	Low risk <10 days	45	-

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

Application number	Application Type	No of Apps	Performance	Target Days
20	Applications for new products	186	99%	15/25
21	Applications for previously imported products	238	99.5%	15
22	All other urgent applications	215	100%	-
	Urgent	0	-	2
	Non Urgent	215	-	10
23	Instant Import Certificates	32,369	-	-
24	Export	225	100%	10
25	Batch Release	2164	99.9%	10

Published Standard Number 6 – Pharmacovigilance

Task number	Task	No.	Performance
26	Human, Animal & Environmental AERs	650	96.2%
27	Validate & extract all UK data from PSURs	1339	100%
28	PSUR Data fully validated, database closed by 25/12/2025	-	-
29	Send final inspection report to MAH	13	100%
30	Number of Benefit-Risk reports validated	1082	100%
31	Number of Benefit-Risk reports undergoing full assessment	6	100%
32	Number of standard signal notifications	76	90.8%
33	Number of urgent signal notifications	-	-

Published Standard Number 7– Inspections

Task number	Task	No.	Performance	Target Days
34	Inspections within 3 years (GMP)	26	100%	-
	Within 5 years (GDP) of last inspection	51	Joint with above	-
35	Inspection Deficiency Reports	76	100%	30.0
36	(GMP) Certificates or (GDP) final reports sent	71	100%	90.0
37	Approval of new Feed business operators and SQP retailer sites	27	100%	45.0
38	Final inspection report to Feed business operators and SQP retailers	225	100%	30.0

Published Standard Number 8 – Enforcement

Task number	Task	No.	Performance
39	Quarterly VMR Breaches	2	100%

Published Standard Number 9 – Residues

Task number	Task	No.	Performance
40	Quarterly Non-Compliance Data	2	100%
41	Sample Testing	21,719	98.5%

Additional information

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