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	49	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)	209	100%	120.0
4	Shortened timetable (Type IB VRA. New ATC (type B). Out of Scope MRLs)	150	99.3%	60.0
5	Minor timetable (National) Type IA VNRA. Administrative Type IB VRA. New ATC (Type A/S). ATC variations.	790	98.9%	30.0
6	Parallel Assessment with EU Procedures	456	99.6%	-
7	Batch timetable (National) specific Batch Control	40	100%	20.0
8	Autogenous Vaccines. New & Variations	2	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)	621	99%
10	Validation	965	94%
11	Issue of authorised documentation	2108	99.7%

Published Standard Number 1 – Applications (European- NI)

Application number	Application Type	Number of applications	Performance
12	New Decentralised (DCP)	18	100%
13	New Mutual Recognition (MRP)	-	-
14	MRP Variations (Type IB & II)	260	100%

Published Standard Number 2 – Public Assessment Reports

Application number	Application type	Total number	Performance
15	Publishing Summary of Product Characteristics (SPCs)	83	100%
16	Publishing Public Assessment Reports (PuARs)	58	100%
17	Updating PuARs	1	100%

Published Standard Number 3 – Quality of Documentation

Application number	Application type	Number of applications	Performance
18	Unreturned Documents	3911	95.1%

Published Standard Number 4 – Product Defects

Task number	Task	Number of tasks	Performance
19	Product Defects reports	33	100%
	High risk <5 days	2	-
	Low risk <10 days	31	-

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	118	98%	15/25
21	Applications for previously imported products	155	100%	15
22	All other urgent applications	144	100%	-
	Urgent	0	-	2
	Non Urgent	144	-	10
23	Instant Import Certificates	22,024	-	-
24	Export	151	100%	10
25	Batch Release	1451	100%	10

Published Standard Number 6 – Pharmacovigilance

Task number	Task	No.	Performance
26	Human, Animal & Environmental AERs	468	97.4%
27	Validate & extract all UK data from PSURs	980	100%
28	PSUR Data fully validated, database closed by 25/12/2025	-	-
29	Send final inspection report to MAH	8	100%
30	Number of Benefit-Risk reports validated	627	100%
31	Number of Benefit-Risk reports undergoing full assessment	2	100%
32	Number of standard signal notifications	51	90.2%
33	Number of urgent signal notifications	-	-

Published Standard Number 7– Inspections

Task number	Task	No.	Performance	Target Days
34	Inspections within 3 years (GMP)	17	100%	-
	Within 5 years (GDP) of last inspection	32	Joint with above	-
35	Inspection Deficiency Reports	44	100%	30.0
36	(GMP) Certificates or (GDP) final reports sent	42	100%	90.0
37	Approval of new Feed business operators and SQP retailer sites	15	100%	45.0
38	Final inspection report to Feed business operators and SQP retailers	146	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	1	100%
41	Sample Testing	15,529	98.1%

Additional information

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