Published Standard Number 1 – Applications (National)

Standard	Application type	Number of applications	Performance	Target days
1	Complex/Major timetable (National new MA applications) Complex/Major new MA applications, e.g. novel therapies, new actives	35	100%	210/180
2	Standard timetable (National VRA-S variations. New MA Copycats. New VHRs)	315	100%	120
3	Shortened timetable (VRA-R variations. New ATC (type A, B, S), ATC variations.	167	100%	60
4	Minor timetable (National) VNRA variations. Administrative VRA-R variations. ATC renewals.	853	98.9%	30
5	Batch timetable (National) specific Batch Control	186	100%	20
6	Autogenous Vaccines. New & Variations	4	100%	60
7	EU Procedures New & variations	247	98.4%	10/20

Published Standard Number 1 – Applications (Other)

Standard	Application Type	Number of applications	Performance
8	Mock-up period completed within 20 days	102	99%
9	Packaging surveillance	50	100%
10	Validation	1088	99.5%
11	Issue of authorised documentation	2499	99.7%

Published Standard Number 2 – Public Assessment Reports

Standard	Application type	Total number	Performance
12	Publishing Summary of Product Characteristics (SPCs)	10	100%
13	Publishing Public Assessment Reports (PuARs)	38	100%
14	Updating PuARs	12	100%

Published Standard Number 3 – Quality of Documentation

Standard	Application type	Number of applications	Performance
15	Unreturned Documents	4393	96.9%

Published Standard Number 4 – Product Defects

Standard	Task	Number of tasks	Performance
16	Product Defects reports with 10 days	45	100%

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

Standard	Application Type	No of Apps	Performance	Target Days

17	Applications for new products	149	100%	15/25
18	Applications for previously imported products	289	100%	15
19	All other applications	414	99.6%	-
	Urgent	1	-	2
	Non Urgent	413	-	10
20	Instant Import Certificates	28,836	-	-
21	Export	98	99%	10
22	Batch Release Requests (BRRs)	1802	99.9%	10

Published Standard Number 6 – Pharmacovigilance

Standard	Task	No.	Performance
23	Human, Animal & Environmental AERs	1512	99.7%
24	Send final inspection report to MAH	11	100%
25	Number of Benefit-Risk reports validated	1075	95.8%
26	Number of Benefit-Risk reports undergoing full assessment	7	100%
27	Number of standard signal notifications	31	87.1%
28	Number of urgent signal notifications	-	-

Published Standard Number 7– Inspections

Standard	Task	No.	Performance	Target Days
29	Inspections within 3 years (GMP)	13	100%	-
	Within 5 years (GDP) of last inspection	42	Joint with above	-
30	Inspection Deficiency Reports	38	100%	30
31	GMP Certificates to manufacturers	13	100%	90

32	Approval of new Feed business operators and SQP retailer sites	10	100%	45
33	Final inspection report to Veterinary Practice Premises, Feed business operators and SQP retailers	527	100%	30

Published Standard Number 8 – Enforcement

Standard	Task	No.	Performance
34	Publication of GB enforcement notices	21	95.2%
35	Initiate removal of illegal listings	81	98.8%

Published Standard Number 9 – Residues

Standard	Task	No.	Performance
36	Quarterly publication of results	-	100%
37	Sample Testing	19,647	98.8%

Additional information

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