

**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)	12	100%	180.0	9
2	Standard timetable (National Type II variations. New MA - MAPIs and Copycats. New VHRs)	28	100%	120.0	6
3	Shortened timetable (National Renewals (MA and VHR) Type IB variations. New ATC (type B). Out of Scope MRLs)	168	100%	60.0	7
4	Minor timetable (National) Type IA variations. Administrative Type IB variations. New ATC (Type A/S). ATC variations and renewals.	171	100%	30.0	17
5	Batch timetable (National) specific Batch Control.	20	100%	20.0	0
6	Autogenous Vaccines. New & Variations	10	90.0%	45.0	35

**Published Standard Number 1 – Applications (Other)**

Application number	Application Type	Number of applications	Performance
7	Mock ups	403	99.5%
8	Validation	588	100%
9	Issue of authorised documentation	861	100%

**Published Standard Number 1 – Applications (European)**

Application number	Application Type	Number of applications	Performance
10	New Centralised (CAP)	8	100%
11	New Decentralised (DCP)	26	100%
12	New Mutual Recognition (MRP) and New DCP	42	100%
13	MRP Variations (Type IB & II) and Renewals	209	100%
14	Parallel Assessment with EU Procedures	0	n/a
15	Shared Assessment with International Partners	0	n/a
16	Referrals	0	n/a

**Published Standard Number 2 – Public Assessment Reports**

Application number	Application type	Total number	Performance
17	Publishing Summary of Product Characteristics (SPCs)	0	n/a
18	Publishing Public Assessment Reports (PuARs)	0	n/a
19	Updating PuARs	1	100%

**Published Standard Number 3 – Quality of Documentation**

Application number	Application type	Number of applications	Performance
20	Unreturned Documents	1685	97.69%

**Published Standard Number 4 – Product Defects**

Task number	Task	Number of tasks	Performance	Target Days	Average Days
21	Product Defects reports High risk <5 days	0	n/a	n/a	n/a
21	Product Defects reports Low risk <10 days	48	100%	n/a	n/a

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

Application number	Application Type	No of Apps	Performance	Target Days	Average Days
22	Applications for new products	96	99%	15	3
23	All other applications	486	98.7%	-	-
23	All other urgent applications	2	-	2	0
23	All other non-urgent applications	484	-	10	2
24	Instant Certificates (Apr-May)	17,609	-	-	-
25	Export	198	100%	10	5.5
26	Batch Release	1722	99.9%	10	2.7

**Published Standard Number 6 – Pharmacovigilance**

Task number	Task	No.	Performance
27	Human, Animal & Environmental AERs	4955	99.58%
28	PSURs	907	100%
29	Inspections	5	100%

**Published Standard Number 7– Inspections**

<b>Task number</b>	<b>Task</b>	<b>No.</b>	<b>Performance</b>	<b>Target Days</b>	<b>Average Days</b>
30	Inspections within 3 years (GMP)	9	73.9%	n/a	n/a
30	Within 5 years (GDP) of last inspection	14	Joint with above	n/a	n/a
31	Final Inspection Reports	<b>23</b>	100%	90.0	36

**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.