Quarterly Performance Report

1 October 2025 to 31 December 2025

Overview

In Quarter 2 of the 2025–26 financial year, the Australian Pesticides and Veterinary Medicines Authority (APVMA) completed 1,358 activities related to regulatory decisions (2863 year-to-date), including 546 product registrations / permits and 20 emergency permits. This represents an increase of almost 100 product registration completions against the first quarter.

We completed 73 compliance investigations and initiated removal of 378 unauthorised products from online marketplaces, remaining on-track to exceed the 2024-25 full-year figure of 1277.

Approvals, Registration and Permits

74.4 per cent of all applications for approval of an active or a label, registration of a product or the issuing of a permit completed in Quarter 2 were completed within statutory timeframes, against a target of 90%, below the previous quarter (80.2 per cent in Quarter 1).

The proportion of permit applications completed within legislative timeframe reduced in Quarter 2 as a strategic temporary redeployment of resources from the agricultural chemicals team concluded, however 132 permits were finalised, which is on par with completions in Quarter 1 (138).

It is important to note that although the timeframe performance percentage has slipped, the APVMA completed 1,358 regulatory decision activities, including 546 product registrations and permits, representing an increase of almost 100 completions compared to Quarter 1. This increase reflects a deliberate focus on finalising overdue applications and priority work, including emergency and minor use permits, which has resulted in a temporary decline in reported timeframe performance.

Table 1: Timeframe Performance Quarter 2 2025-26

Area	Received	In progress	Finalised	Finalised on time
Agricultural Chemicals	187	644	159	60.4%
Veterinary Medicines	230	333	255	84.7%
Permits (excl. emergency permits)	131	273	132	60.6%
Emergency permits [1]	10	8	20	N/A
Active constituents	94	278	80	92.5%
Total	**652**	**1536**	**646**	**74.4%**

While these results are lower than expected, they must be interpreted in context: the completion of large numbers of previously overdue applications negatively impacts headline timeframe measures even as overall throughput increases. Importantly, the APVMA completed the highest number of technical applications in over 18 months and finalised 20 emergency permits during the quarter, demonstrating increased operational activity and responsiveness.

[1] Emergency permits (Item 22) do not have a numerical statutory timeframe. The APVMA must determine the application as soon as is practicable in the circumstances of the case.

During Quarter 2, the APVMA implemented a full pause and redesign of the reference product pathway to strengthen robustness and consistency, while simultaneously increasing rejection rates for poor-quality applications. These improvement activities required additional effort from scientific and regulatory staff and contributed to short-term reductions in timeframe performance. At the same time, the organisation continued to prioritise backlog reduction, supported by redeployment of staff across efficacy and environment areas, the engagement of external scientific reviewers, and the onboarding of additional delegates.

One of our most closely monitored results relates to the processing of technical applications for agricultural chemicals (previously described as pesticides majors). This result, at **36.0 per cent**, displayed the forecast decline anticipated in previous quarters arising from the temporary diversion of resources to the permits team.

During this period, the APVMA finalised 8 products containing new active constituents this quarter, up from 1 last quarter. Evaluation commenced on **12** new applications for a new active constituent with accompanying new product. These were:

- 3 applications for assessment under item 2
- 9 applications for assessment under item 27.

The APVMA is prioritising these assessments, however there is still a backlog that is expected to last until mid-2026, due to the long timeframes and complex nature of these assessments. Many of these products are biological rather than synthetic chemicals. APVMA's foresight activities related to biologicals support our preparedness for these applications.

As at 31 December 2025, the APVMA had a total of 76 item 1, 2 and 27 applications under evaluation. This performance measure and result is likely to fluctuate significantly throughout the year and is heavily impacted by the inherent complexity of these long-timeframe applications.

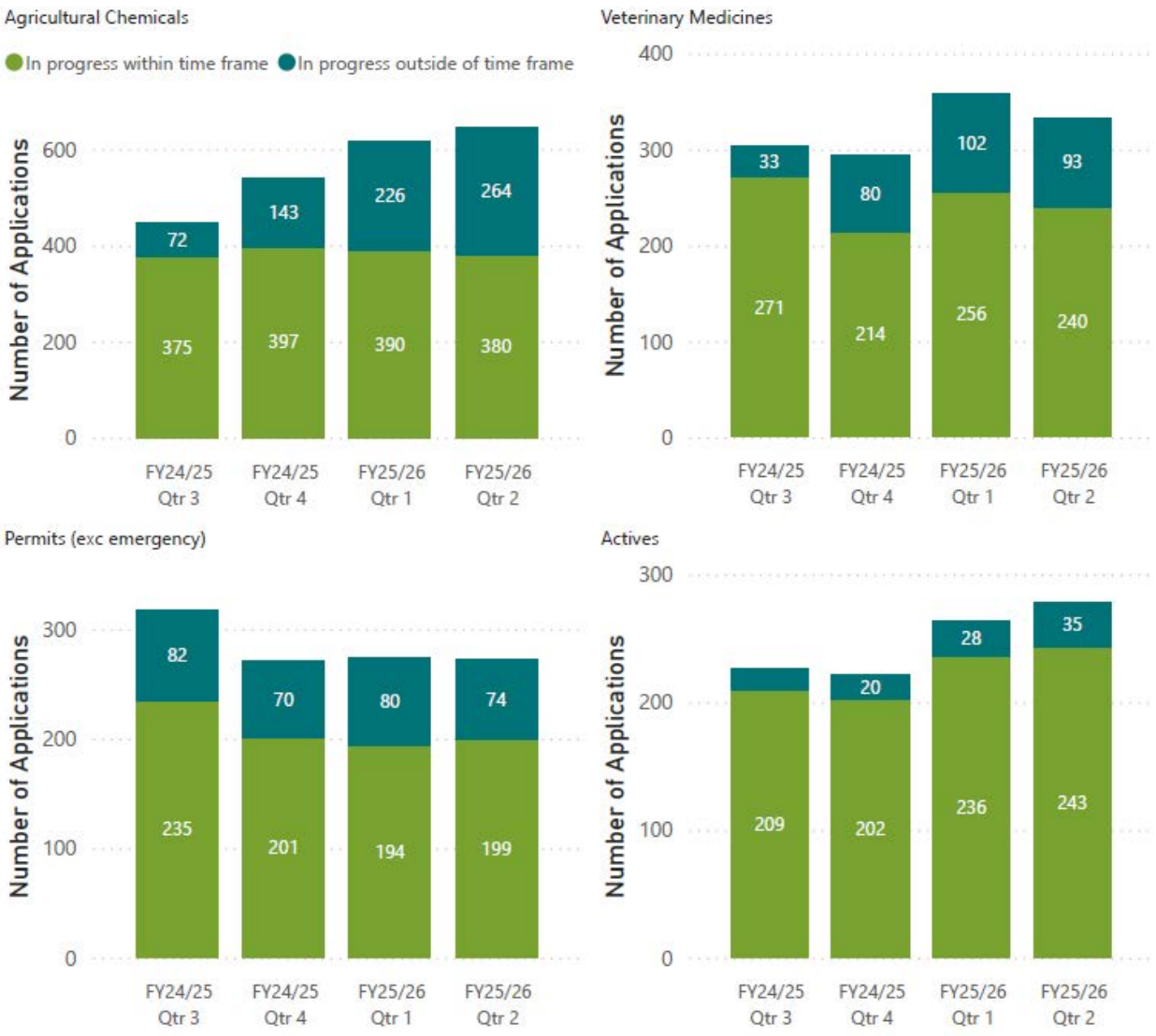


Figure 1: Applications in progress within and outside of timeframe

The number of applications that went overdue during the reporting period decreased in Quarter 2 in both agricultural chemicals and veterinary medicines.

Deficiencies in submitted applications continue to cause delays in timeframe performance. The APVMA is strengthening its requirements for applicants, to make the best use of APVMA resources.

Post market regulatory actions

Chemical reconsideration activity continued apace throughout Quarter 2 with publication of the proposed regulatory decision for anticoagulant rodenticides on 16 December 2025. The APVMA also issued notices proposing to suspend the registration of second-generation anticoagulant rodenticides, due to risks to non-target species identified in the proposed regulatory decision. A final decision on the proposed suspension is expected in Quarter 3, following a 6-week public consultation.

The APVMA made a final decision to suspend certain dimethoate products on 11 November 2025, due to new data showing a potential risk to human health due the combination of significantly

increased Australian consumption of blueberries, blackberries and raspberries and the potential of residues remaining on those crops.

Concurrent resourcing of multiple, extremely complex chemical reviews is a major challenge for the APVMA and impacts capacity to process technical applications for agricultural chemicals. Strategies are being implemented to respond to this pressure point, including expansion of use of external science reviewers and other measures to realise the full value of separating post-market and pre-market activities in the recent restructure.

The proportion of serious **adverse experience reports** received, and assessments completed within 20 business days, is a new reporting metric initiated by the APVMA with performance at **90 per cent** against a 75 per cent target.

Table 2: Adverse Experience Reporting Program

Summary		2024-25^[2]	2024-25 Q2^	2025-26 Q2^
AERP	Performance	43.7%	63.7%	90.0%
	Number of serious individual reports received	2797	1391	1664
	Number of serious individual reports closed	2485	878	1688
	Number of serious adverse incidents received	2423	1253	1368

90.0% of serious adverse experience reports were received and initial assessments completed by Quarter 2 within the 20 working days timeframe. This is significantly up compared to what has been calculated for the same period last financial year (63.7% by Quarter 2 FY2024-25):

By Quarter 2 on a year-on-year basis there has been an approximate 20% increase in serious adverse experience reports (1391 to 1664).

In the second quarter, the APVMA continued its rigorous program of **compliance action**, initiating the removal of **378** unauthorised products from online marketplaces, with **706** removals year-to-date, compliance action remains on track for a full year figure to exceed the 1,277 in 2024–25. Investigations were initiated into **64** compliance matters, including generating 9 forfeiture actions under the *Customs Act 1901.*

Table 3: Summary of investigations

Summary of investigations	2024-25	2025-26 Q1	2025-26 Q2
Commenced	289	96	64
Closed	297	99	73
Remaining open	22	37	18

Table 4: Summary of investigation outcomes

Summary of investigation outcomes	2024-25 Cases	2025-26 Q1 Cases	2025-26 Q2 Cases
Negotiated Compliance / Education	161	15	47

[2] ^This is the first year calculating this metric for serious adverse experience reports. Values for 2024-25 have been calculated for comparison only and were not reported previously.

Formal Warnings	77	36	28
Infringement Notices	10	3	0
Enforceable Undertakings	0	0	0
Civil Proceedings	0	0	0
Prosecutions	0	0	0
Customs Act Forfeiture	56	38	9
No Offence / Insufficient Evidence	32	0	8
Online product removals	1277	328	378

The Manufacturing Quality and Licensing (**MQL**) performance target is 90 per cent of all audit reviews finalised within a 4-month timeframe. 87.8 per cent of all audit reviews completed in Quarter 2 were completed within 4-month timeframe. This is slightly reduced against the previous financial year, with several projects ongoing to ensure consistent timeframe performance and decision making in auditing processes.

Table 5: Manufacturing Quality and Licencing (MQL)

Summary		2024-25	2024-25 Q2	2025-26 Q2
MQL	Performance	86.7%	95.7%	87.8%
	Number of audits commenced	86	47	52
	Number of audits closed and reviewed	98	47	49

The **Recalls** performance target is 100 per cent of all voluntary recall notices published to the APVMA website within 3 working days and gazetted within 14 days as per Agvet Code requirements.

Table 6: Recalls

Summary		2024-25	2024-25 Q2	2025-26 Q2
Recalls	Performance	100%	100%	100%
	Number of voluntary recalls	29	13	23

The APVMA is achieving 100 per cent for all Recalls. There has been a significant increase in voluntary recalls compared to the same period last financial year (from 13 to 23). Most products recalled are sterile injectable veterinary chemical products due to sterility or particulate matter concerns from both domestic and overseas manufacturers.

Table 7: Hormonal Growth Promotants

Summary		2024-25	2024-25 Q2	2025-26 Q2
HGP	Performance	99.4%	96.8%	100%
	Number of applications	180	32	36

The Hormonal Growth Promotants (HGPs) performance target is 90 per cent of all notification number applications processed within 5 working days. Most renewal applications (~70-80 per cent) occur in Quarter 4 of each financial year. The APVMA is achieving 100 per cent for HGP applications in Quarter 2.

Concluding comments

Overall, the second quarter report indicates the scale of actions being taken across the agency to fulfil its responsibilities and reveals a range of crucial decisions that have been made on agricultural chemicals and veterinary medicines. The report also builds from the first quarter baseline for measuring improvement in performance as we move into 2026 where we will be progressing the following initiatives:

- Sustainable funding – with a primary focus on cost recovery policy in conjunction with the Department of Agriculture, Fisheries and Forestry
- Implementing critical elements of our ICT Roadmap
- Productivity Initiatives – use of international assessments; and increasing the onus on applicants to submit quality applications
- Structure and resources – maximising the gains inherent in our new organisational structure; and expanded use of external science reviewers.

Looking ahead, timeframe performance is expected to remain under pressure in Quarter 3 and may decline slightly further before improvement is realised. Most assessment areas continue to experience backlog pressures, and it is anticipated that recovery will take more than a year. The APVMA remains focused on increasing capacity, strengthening application quality, prioritising high-risk and high-impact work, and maintaining transparent engagement with stakeholders to provide greater predictability and confidence in regulatory processes.

While achieving reliable timeframes for the work we do is undoubtedly important, the APVMA is also committed to maintaining the trust of the Australian public, as that is critical to maintaining confidence in Australian agricultural chemicals and veterinary medicines.

**In line with other changes to the APVMA quarterly reports, references to 'pesticides' have been updated to refer to 'agricultural chemicals' and references o 'pesticide majors' have been changed to refer to 'technical agricultural chemical applications'.*

Analysis by application type (1 October – 31 December 2025)

Table 8: Summary of activities related to regulatory decisions

Types of regulatory decisions	Received	In progress	Finalised
Pre-application assistance	39	88	50
Product registration—agricultural chemicals	187	644	159
Product registration—veterinary medicines	230	333	255
Permits (exc emergency)	131	273	132
Emergency permits (Item 22)	10	8	20
Actives	94	278	80
Items 8L, 8M, 8P	99	31	89
Technical assessment (Item 25)	5	17	2
Notifiable variations	268	2	266
Import consents	164	24	175
Certificate of export	131	30	128
Interchangeable constituent determination	0	0	0
Total	**1358**	**1728**	**1356**

Applications processed

Table 9: Applications processed

Type	Received	In progress	Finalised	Finalised on time (%)	In progress still on time (%)
Actives	94	278	80	92.5%	87.4%
Agricultural Chemicals	187	644	159	60.4%	59.0%
Emergency permits (Item 22)	10	8	20	N/A	N/A
Permits (exc emergency)	131	273	132	60.6%	72.9%
Veterinary Medicines	230	333	255	84.7%	72.1%
Total	**652**	**1536**	**646**	**74.4%**	**69.5%**

Table 10: Applications processed continued

Type	Finalised on time	Finalised outside timeframe	Went overdue in reporting period
Actives	74	6	14
Agricultural Chemicals	96	63	101
Permits (exc emergency)	80	52	49
Veterinary Medicines	216	39	30
Total	**466**	**160**	**194**

Applications in evaluation

Table 11: Applications in evaluation

Type	Evaluations in progress	Evaluations in progress and overdue
Actives	221	34
Agricultural Chemicals	550	259
Emergency permits (Item 22)	8	2
Permits (exc emergency)	228	74
Veterinary Medicines	257	92
Total	**1264**	**461**

Pre-application assistance

Pre-application assistance (PAA) is an optional, three-tiered, fee-based service for applicants to provide them with advice before they submit an application for registration. PAAs do not have a numerical statutory timeframe; the target timeframe for each tier is based on the anticipated complexity of the proposed application and effort required to provide the assistance.

Table 12: Pre-application assistance

Type	Received	In Progress	Finalised	Processed on time (%)
PAA Tier 1	11	15	7	0.0%
PAA Tier 2	26	65	42	26.2%
PAA Tier 3	2	8	1	0.0%
Total	**39**	**88**	**50**	**22.0%**

Permits

Table 13: Permits

Category Name	Received	In Progress	Finalised	Issued	Refused	Withdrawn or Other
Ag chemical and Vet chemical	0	0	0	0	0	0
Agricultural chemical	103	213	116	95	0	21
Other	0	0	0	0	0	0
Veterinary chemical	28	60	16	14	0	2

Emergency Permits

Emergency Permits (Item 22) do not have a numerical statutory timeframe. The APVMA must determine the application as soon as is practicable in the circumstances of the case. In this reporting period, the APVMA issued 14 emergency permits, taking an average of 85 days, with a range of 3 to 328 days.

Table 14: Emergency permits

Types of regulatory decisions	Received	In Progress	Finalised	Issued	Refused	Withdrawn or Other
Emergency permits (Item 22)	10	8	20	14	3	3

Technical assessment, notifiable variations, holders and nominated agents

Table 15: Technical assessment, notifiable variations, holders and nominated agents

Types of regulatory decisions	Received	In Progress	Finalised	Finalised on time (%)
Items 8L, 8M, 8P	99	31	89	100.0%
Technical assessment (Item 25)	5	17	2	
Notifiable variations	268	2	266	100.0%

Consents to Import

Consents to import are issued to allow importation – in limited circumstances – of unregistered products or unapproved actives into Australia when a legitimate reason exists for a person or company to have possession of the chemicals in Australia. There is no statutory timeframe for consents to import – the APVMA seeks to process these within 14 days.

Table 16: Consents to import

Types of regulatory decisions	Received	In Progress	Finalised	Finalised on time (%)
Import consents	164	24	175	96.6%

Certificates of Export

Before accepting exports of a chemical product from Australia, many countries require an assurance from the government authority responsible for regulating the product in Australia. This is provided by the APVMA in the form of a Certificate of Export. There is no statutory timeframe for certificates of export – the APVMA seeks to process these within 20 working days. Please note that an application may request multiple certificates.

Table 17: Applications for certificates of export

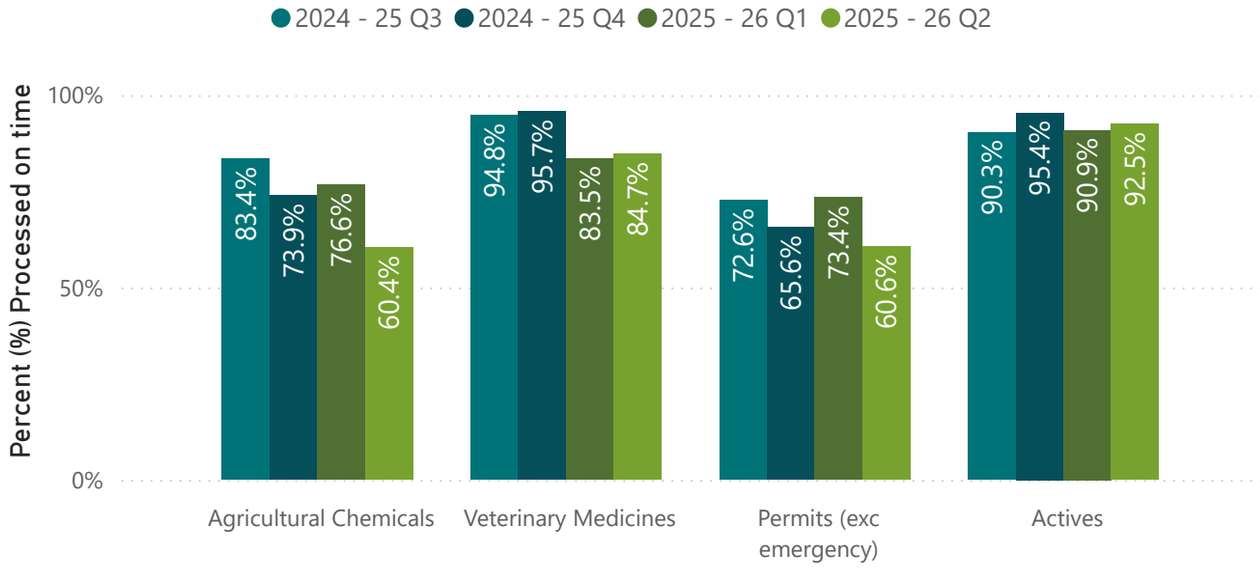
	Received	In Progress	Finalised	Issued	Finalised on time (%)
Applications	131	30	128	128	60.9%

Table 18: Certificates of export

	Received	In Progress	Finalised	Issued	Finalised on time (%)
Certificates	254	48	244	244	66.4%

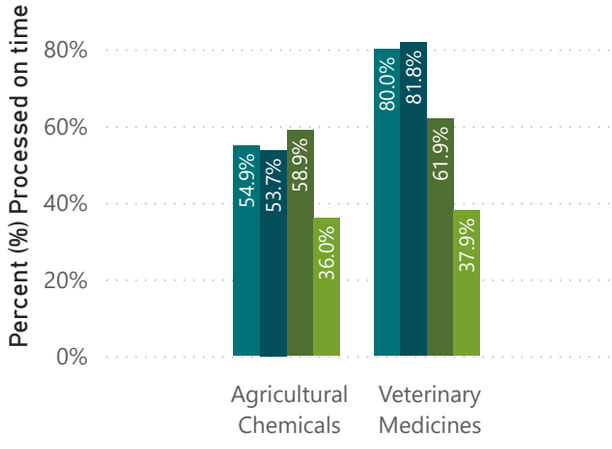
Applications in detail

Timeframe Performance

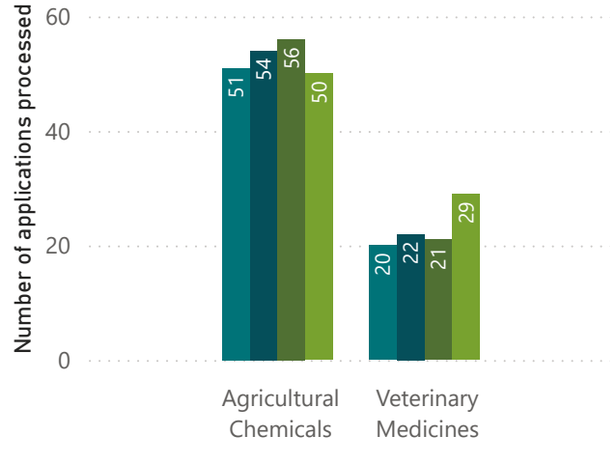


Major and Non-Technical Applications

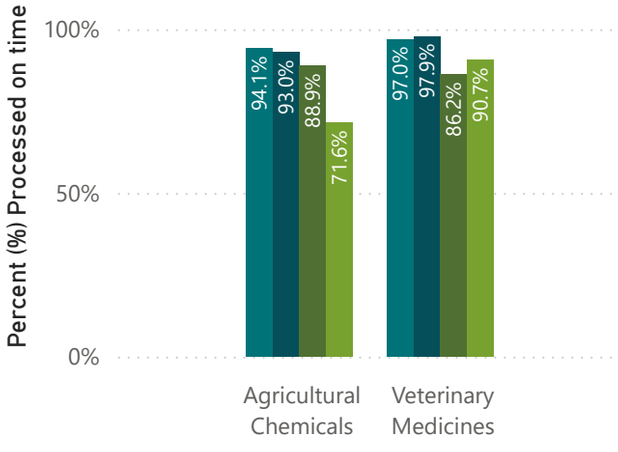
Major - timeframe performance



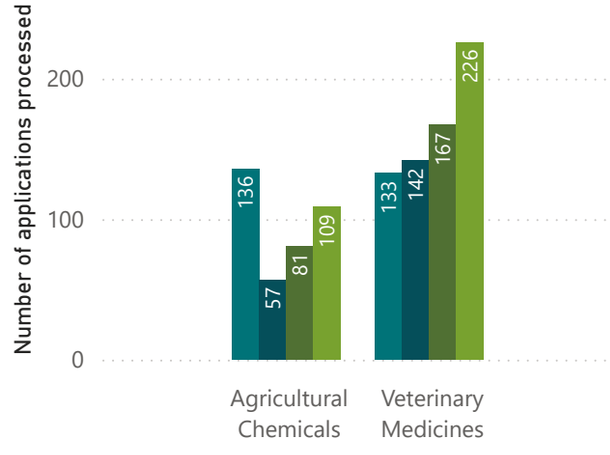
Major - number processed



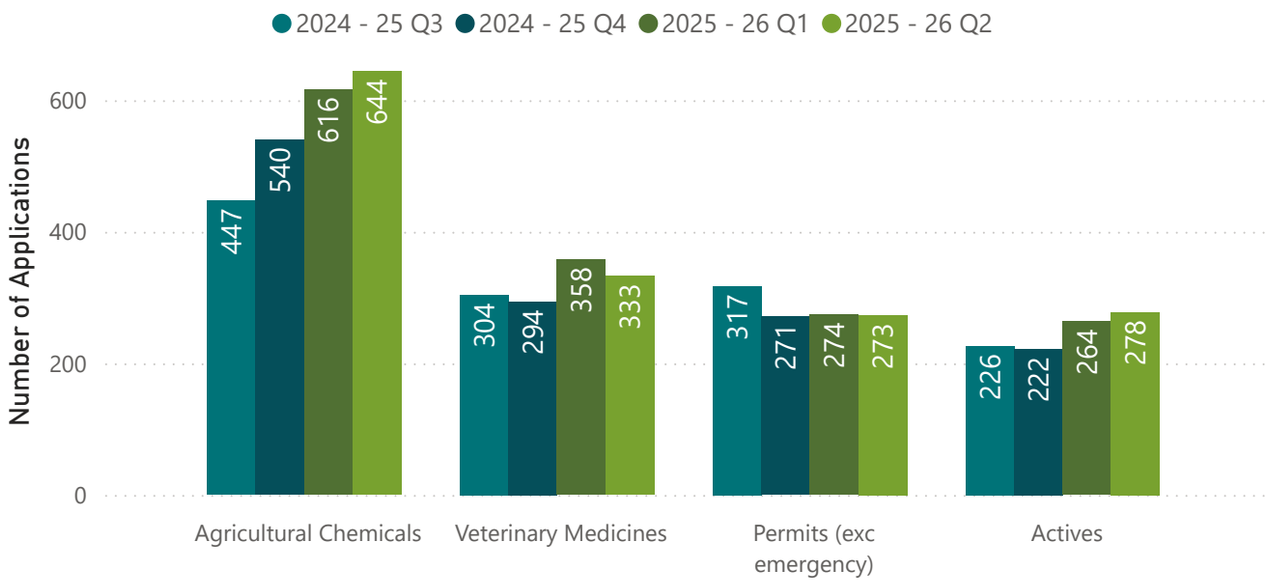
Non-technical - timeframe performance



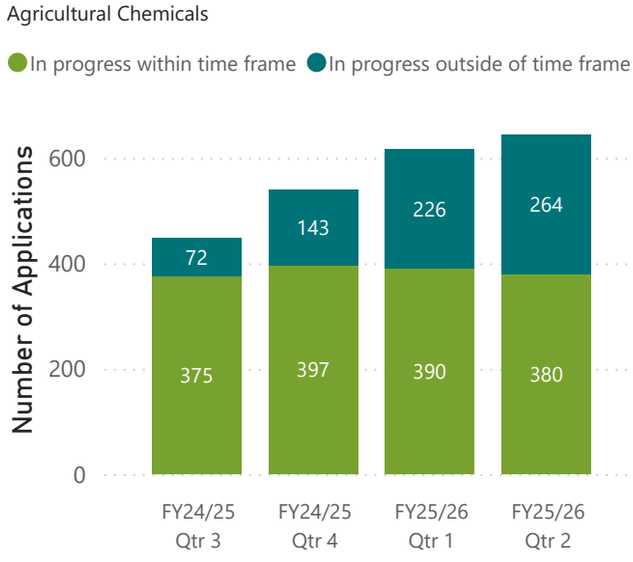
Non-technical - number processed

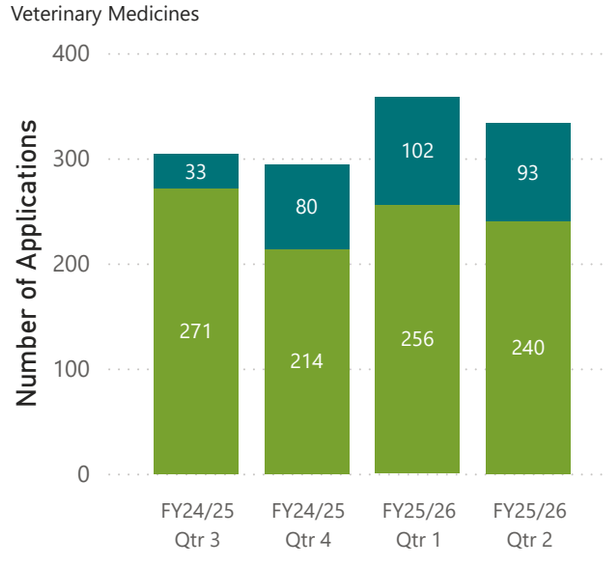


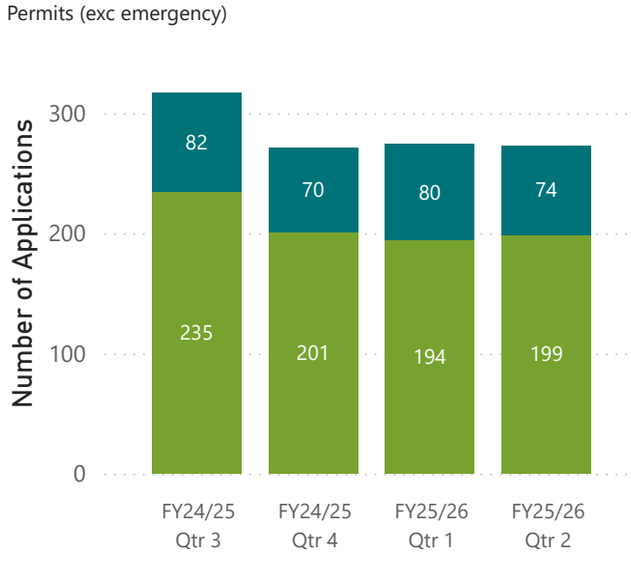
Work in progress

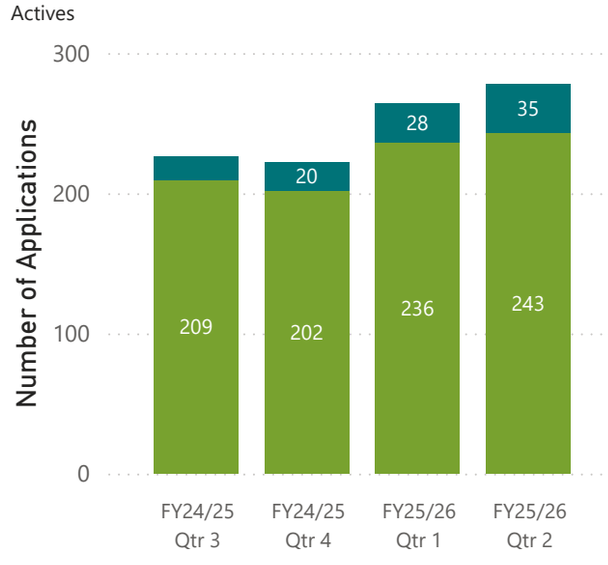


Work in progress still within time frame









Application by item number

Table 19: All applications by Item Number

Type of application	Received	In Progress	Finalised	Processed on time (%)
Products	**417**	**977**	**414**	**75.4%**
1	0	0	0	N/A
2	3	31	1	0.0%
3	0	0	0	N/A
4	0	0	0	N/A
5	6	25	2	50.0%
6	1	15	1	0.0%
7	17	214	47	21.3%
8	46	74	40	90.0%
9	0	0	4	75.0%
10	47	223	36	30.6%
10A	6	6	7	85.7%
11	0	1	0	N/A
12	112	157	75	88.0%
13	1	0	2	100.0%
13A	102	0	160	100.0%
14	67	184	32	50.0%
27	9	45	7	14.3%
27V	0	2	0	N/A
Actives	**94**	**278**	**80**	**92.5%**
1	0	1	0	N/A
2	4	35	1	0.0%
5	0	0	0	N/A
6	0	0	0	N/A
10	0	0	0	N/A
14	0	0	0	N/A
15	0	3	0	N/A
16	0	0	0	N/A
17	63	149	41	97.6%
18	4	20	6	83.3%
24	15	37	14	92.9%
24V	2	10	15	100.0%
27	6	23	3	33.3%
27V	0	0	0	N/A
Permits	**141**	**281**	**152**	**60.6%**
19	8	10	11	90.9%
20	61	77	47	72.3%
21	50	153	48	45.8%
22	10	8	20	N/A
23	9	30	19	47.4%
112A	3	3	7	71.4%
Other	**0**	**0**	**0**	**N/A**
24	0	0	0	N/A
Ingredient determination	**0**	**0**	**0**	**N/A**
28	0	0	0	N/A
29	0	0	0	N/A
Total	**652**	**1536**	**646**	**74.4%**

Table 20: Agricultural chemical applications by item number

Type of application	Commenced	In Progress	Finalised	Processed on time (%)
Products	**187**	**644**	**159**	**60.4%**
1	0	0	0	N/A
2	1	12	0	N/A
3	0	0	0	N/A
4	0	0	0	N/A
5	6	21	2	50.0%
6	1	13	1	0.0%
7	7	164	31	12.9%
8	42	62	30	96.7%
9	0	0	4	75.0%
10	38	166	23	30.4%
10A	4	4	7	85.7%
11	0	1	0	N/A
12	56	81	35	97.1%
13	0	0	1	100.0%
13A	1	0	1	100.0%
14	30	109	21	42.9%
27	1	9	3	33.3%
27V	0	2	0	N/A
Permits	**110**	**219**	**130**	**59.5%**
19	2	4	10	90.0%
20	52	64	43	74.4%
21	42	122	43	41.9%
22	7	6	14	N/A
23	4	20	13	38.5%
112A	3	3	7	71.4%
Other	**0**	**0**	**0**	**N/A**
24	0	0	0	N/A
Total	**297**	**863**	**289**	**60.0%**

Table 21: Veterinary medicines applications by item number

Type of application	Commenced	In Progress	Finalised	Processed on time (%)
Products	**230**	**333**	**255**	**84.7%**
1	0	0	0	N/A
2	2	19	1	0.0%
3	0	0	0	N/A
4	0	0	0	N/A
5	0	4	0	N/A
6	0	2	0	N/A
7	10	50	16	37.5%
8	4	12	10	70.0%
9	0	0	0	N/A
10	9	57	13	30.8%
10A	2	2	0	N/A
11	0	0	0	N/A
12	56	76	40	80.0%
13	1	0	1	100.0%
13A	101	0	159	100.0%
14	37	75	11	63.6%
27	8	36	4	0.0%
27V	0	0	0	N/A
Permits	**31**	**62**	**22**	**68.8%**
19	6	6	1	100.0%
20	9	13	4	50.0%
21	8	31	5	80.0%
22	3	2	6	N/A
23	5	10	6	66.7%
112A	0	0	0	N/A
Other	**0**	**0**	**0**	**N/A**
24	0	0	0	N/A
Total	**261**	**395**	**277**	**83.8%**

Average decision time

Table 22: All applications by Item Number

	Standard			Extended		
Type of Application	Finalised	Average Duration	Assessment Period	Finalised	Average Duration	Assessment Period
Products	**354**	**3.3**		**60**	**15.2**	
1	0	N/A	18	0	N/A	25
2	0	N/A	~	1	33.4	23.1
3	0	N/A	18	0	N/A	25
4	0	N/A	18	0	N/A	25
5	0	N/A	8	2	12.7	12
6	1	9.9	8	0	N/A	12
7	41	6.4	3	6	9.6	5
8	37	3.1	3	3	4.7	5
9	4	2.2	2	0	N/A	4
10	13	10.7	8.4	23	19.0	10.7
10A	7	2.0	2.0	0	N/A	~
11	0	N/A	10	0	N/A	14
12	67	2.7	3	8	5.6	5
13	2	2.9	3	0	N/A	5
13A	160	0.8	1	0	N/A	2
14	20	10.4	6.8	12	12.7	9.8
27	2	36.6	26.8	5	28.0	24.1
27V	0	N/A	~	0	N/A	~
Actives	**59**	**5.9**		**21**	**9.6**	
1	0	N/A	18	0	N/A	25
2	0	N/A	~	1	33.4	23.1
5	0	N/A	8	0	N/A	12
6	0	N/A	8	0	N/A	12
10	0	N/A	~	0	N/A	~
14	0	N/A	~	0	N/A	~
15	0	N/A	14	0	N/A	20
16	0	N/A	9	0	N/A	13
17	32	6.7	7	9	9.5	10
18	5	7.0	7	1	N/A	10
24	8	2.9	4.1	6	4.6	8.6
24V	13	3.3	0.0	2	4.7	0.0
27	1	36.6	26.8	2	22.6	20.2
27V	0	N/A	~	0	N/A	~
Permits	**126**	**6.0**		**26**	**10.6**	
19	11	2.6	3	0	N/A	5
20	46	3.9	3	1	N/A	5
21	37	7.8	5.3	11	14.0	11.2
22	11	1.6	N/A	9	5.3	N/A
23	14	7.1	4.6	5	13.6	7.6
112A	7	19.0	0.0	0	N/A	N/A
Other	**0**	**N/A**		**0**	**N/A**	
24	0	N/A	~	0	N/A	~
Ingredient determination	**0**	**N/A**		**0**	**N/A**	
28	0	N/A	~	0	N/A	~
29	0	N/A	~	0	N/A	~
Total	**539**	**4.2**		**107**	**13.1**	

Notes for average decision times

Average decision time is the average time taken to complete processing from commencement to when a regulatory decision is made or the average time is calculated by adding the decision times and dividing these by the number of applications finalised in that period in that category. Decision time is calculated by subtracting the lodgement date from the processed date and dividing by 30.4375 to get the decision time in months (accounting for leap years). The assessment period for each application type and item number is set in the legislation - this is the standard decision time. However, there are 2 conditions which allow the APVMA to vary or extend the assessment period:

- A change in the category of an application may be made once in the life of the application under regulation 70B of the Agvet Code.
- One extension of 1.33 times the original assessment period (rounded up) plus one month may be applied under section 159 of the Agvet Code.

N/A means no applications were finalised for that item in the reporting period, the application was withdrawn or voided, or there is no statutory assessment period.

~ applies to items that have a variable legislated assessment period because they are modular in nature. Each application will have a different expected duration, as determined by the mix of modules required for the application.

Table 23: Agricultural chemical product applications by Item Number

	Standard			Extended		
Type of Application	Finalised	Average Duration	Assessment Period	Finalised	Average Duration	Assessment Period
Products	**135**	**5.3**		**24**	**19.1**	
1	0	N/A	18	0	N/A	25
2	0	N/A	~	0	N/A	~
3	0	N/A	18	0	N/A	25
4	0	N/A	18	0	N/A	25
5	0	N/A	8	2	12.7	12
6	1	9.9	8	0	N/A	12
7	31	6.4	3	0	N/A	5
8	29	2.6	3	1	4.0	5
9	4	2.2	2	0	N/A	4
10	8	13.1	8.2	15	20.5	10.8
10A	7	2.0	2.0	0	N/A	~
11	0	N/A	10	0	N/A	14
12	34	2.5	3	1	4.6	5
13	1	2.9	3	0	N/A	5
13A	1	N/A	1	0	N/A	2
14	17	10.6	6.7	4	17.5	11.0
27	2	36.6	26.8	1	49.6	39.5
27V	0	N/A	~	0	N/A	~
Permits	**108**	**6.4**		**22**	**11.7**	
19	10	2.6	3	0	N/A	5
20	42	3.9	3	1	N/A	5
21	32	8.2	5.3	11	14.0	11.2
22	8	1.8	N/A	6	5.4	N/A
23	9	8.4	4.7	4	17.3	7.0
112A	7	19.0	0.0	0	N/A	N/A
Other	**0**	**N/A**		**0**	**N/A**	
24	0	N/A	~	0	N/A	~
Total	**243**	**5.7**		**46**	**15.6**	

Table 24: Veterinary medicine product applications by Item Number

	Standard			Extended		
Type of Application	Finalised	Average Duration	Assessment Period	Finalised	Average Duration	Assessment Period
Products	**219**	**1.9**		**36**	**12.7**	
1	0	N/A	18	0	N/A	25
2	0	N/A	~	1	33.4	23.1
3	0	N/A	18	0	N/A	25
4	0	N/A	18	0	N/A	25
5	0	N/A	8	0	N/A	12
6	0	N/A	8	0	N/A	12
7	10	6.3	3	6	9.6	5
8	8	4.7	3	2	5.0	5
9	0	N/A	2	0	N/A	4
10	5	7.9	8.6	8	16.6	10.5
10A	0	N/A	~	0	N/A	~
11	0	N/A	10	0	N/A	14
12	33	3.0	3	7	5.8	5
13	1	N/A	3	0	N/A	5
13A	159	0.8	1	0	N/A	2
14	3	7.9	8.0	8	10.3	9.2
27	0	N/A	N/A	4	22.6	20.2
27V	0	N/A	~	0	N/A	~
Permits	**18**	**3.5**		**4**	**5.4**	
19	1	2.6	3	0	N/A	5
20	4	3.9	3	0	N/A	5
21	5	4.3	5.0	0	N/A	~
22	3	0.8	N/A	3	5.1	N/A
23	5	4.0	4.4	1	6.3	8.7
112A	0	N/A	N/A	0	N/A	N/A
Other	**0**	**N/A**		**0**	**N/A**	
24	0	N/A	~	0	N/A	~
Total	**237**	**2.0**		**40**	**11.9**	

Average decision time – Cumulative quarters to date

Table 25: Average decision time, all applications for completed quarters this financial year

Type of application	Number finalised	Average duration	Assessment period
Products	**739**	**4.9**	**18**
1	0	N/A	18
2	1	33.4	23.1
3	0	N/A	18
4	0	N/A	18
5	3	15.7	8
6	2	9.9	8
7	60	6.8	3
8	62	3.2	3
9	4	2.2	2
10	64	15.5	9.9
10A	22	2.0	2.0
11	0	N/A	10
12	146	3.3	3
13	16	2.8	3
13A	273	0.8	1
14	78	10.0	7.7
27	8	26.8	22.6
27V	0	N/A	~
Actives	**146**	**7.1**	**18**
1	0	N/A	18
2	1	33.4	23.1
5	0	N/A	8
6	0	N/A	8
10	0	N/A	~
14	0	N/A	~
15	0	N/A	14
16	0	N/A	9
17	81	8.2	7
18	10	7.0	7
24	30	4.2	5.8
24V	20	3.1	0.0
27	4	29.6	23.5
27V	0	N/A	~
Permits	**308**	**5.9**	**3**
19	18	2.6	3
20	125	3.3	3
21	86	10.3	6.8
22	37	3.8	N/A
23	34	7.1	4.8
112A	8	15.8	0.0
Other	**0**	**N/A**	**~**
24	0	N/A	~
Ingredient determination	**0**	**N/A**	**~**
28	0	N/A	~
29	0	N/A	~
Total	**1193**	**5.5**	