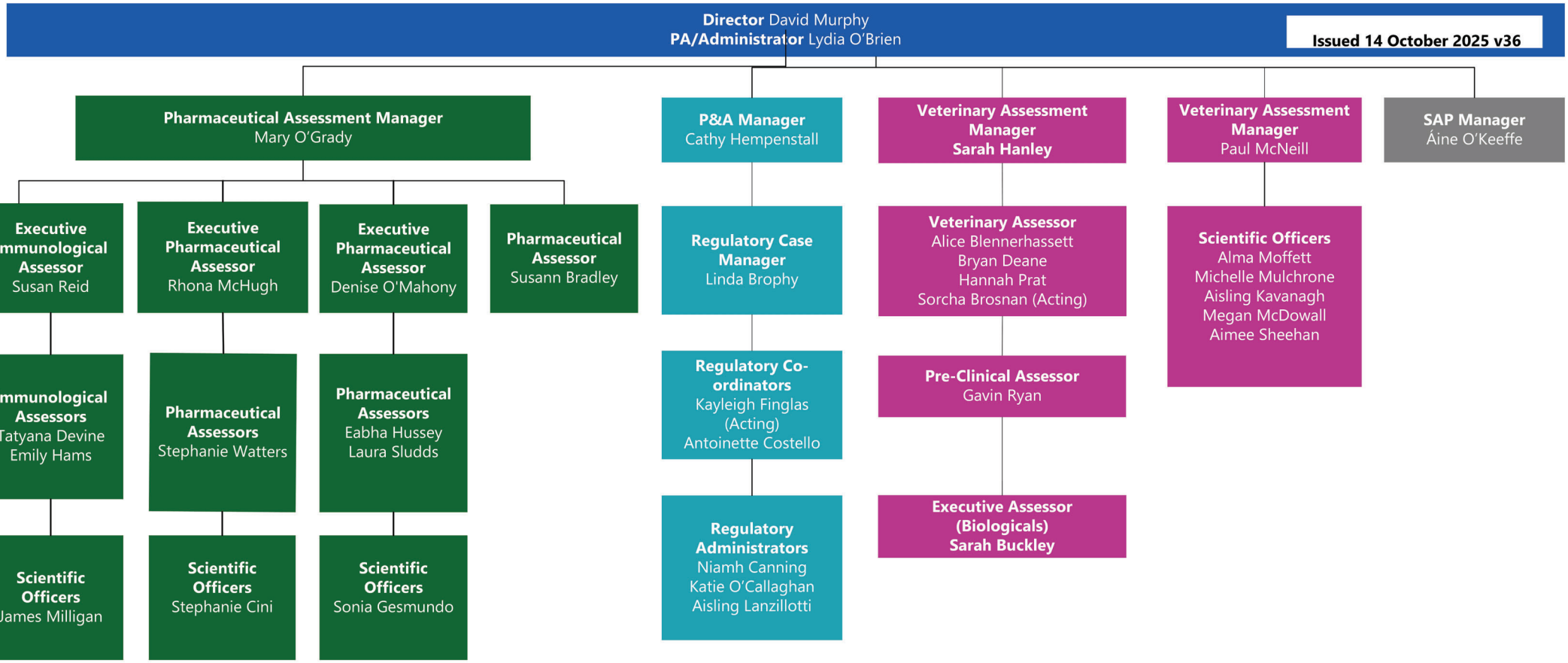
VS department structure



Summary of activities:

- Assessment of the quality data in marketing authorisation applications for veterinary medicinal products
- Assessment of the quality data in applications to vary veterinary medicinal products authorisations
- Assessment of applications for registration of homeopathic veterinary products
- Assessment of product labelling
- Preparing publications on quality issues
- Assessment of data submitted in EU applications for scientific advice

Summary of activities:

- All national and EU marketing authorisation applications for medicinal products for veterinary use
- EU applications for scientific advice on medicinal products
- Applications for clinical field trials using medicinal products
- Classification enquiries
- Scheduling and capacity planning

Summary of activities:

- Assessment of the safety and efficacy data in applications for veterinary medicinal products
- Assessment of classification enquiries
- Assessment of data submitted in EU applications for scientific advice
- Assessment of post-marketing pharmacovigilance data
- Evaluation and follow-up of adverse reaction reports
- Preparing publications on safety/pharmacovigilance issues

Summary of activities:

- All applications submitted under Directive 2010/63/EU relating to the conduct of research and regulatory studies in animals
- Clinical field trial applications