VS department structure

Director David Murphy
PA/Administrator Lydia O'Brien

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Pharmaceutical Assessment Manager
Mary O'Grady

- **Executive Immunological Assessor** Susan Reid
  - **Immunological Assessors** Tatyana Devine, Emily Hams
    - **Scientific Officers** James Milligan
- **Executive Pharmaceutical Assessor** Rhona McHugh
  - **Pharmaceutical Assessors** Stephanie Watters, Eilish Bonner
    - **Scientific Officers** Stephanie Cini
- **Executive Pharmaceutical Assessor** Denise O'Mahony
  - **Pharmaceutical Assessors** Eabha Hussey, Laura Sludds, Luke Bowden
    - **Scientific Officers** Sonia Gesmundo
- **Pharmaceutical Assessor** Susann Bradley

P&A Manager
Cathy Hempenstall

- **Regulatory Case Manager** Linda Brophy
- **Regulatory Co-ordinators** Sinead Agar, Antoinette Costello
- **Regulatory Administrators** Kayleigh Finglas, Chloe Boyd (Acting), Aisling Lanzillotti

Veterinary Assessment Manager
Paul McNeill

- **Veterinary Assessor** Alice Blennerhassett, Bryan Deane, Hannah Pratt, Sorcha Brosnan
- **Pre-Clinical Assessor** Gavin Ryan
- **Executive Assessor (Biologicals)** Sarah Buckley

Veterinary Assessment Manager
Sarah Hanley

- **Scientific Officers** Alma Moffett, Michelle Mulchrone, Aisling Kavanagh, Megan McDowall, Aimee Sheehan

SAP Manager
Áine O'Keeffe

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