

VS department structure

Director David Murphy
PA/Administrator Lydia O'Brien

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Pharmaceutical Assessment Manager
Mary O'Grady

P&A Manager
Cathy Hempenstall

Veterinary Assessment Manager
Sarah Hanley

Veterinary Assessment Manager
Paul McNeill

SAP Manager
Áine O'Keeffe

Executive Immunological Assessor
Susan Reid

Executive Pharmaceutical Assessor
Rhona McHugh

Executive Pharmaceutical Assessor
Denise O'Mahony

Pharmaceutical Assessor
Susann Bradley

Regulatory Case Manager
Linda Brophy

Veterinary Assessor
Alice Blennerhassett
Bryan Deane
Hannah Prat
Sorcha Brosnan (Acting)

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

Immunological Assessors
Tatyana Devine
Emily Hams

Pharmaceutical Assessors
Stephanie Watters

Pharmaceutical Assessors
Eabha Hussey
Laura Sluuds

Regulatory Co-ordinators
Kayleigh Finglas (Acting)
Antoinette Costello

Pre-Clinical Assessor
Gavin Ryan

Scientific Officers
James Milligan

Scientific Officers
Stephanie Cini

Scientific Officers
Sonia Gesmundo

Regulatory Administrators
Niamh Canning
Katie O'Callaghan
Aisling Lanzillotti

Executive Assessor (Biologicals)
Sarah Buckley

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