



PLANT PROTECTION

Our regulatory expertise at your service



CHEMICAL PRODUCTS - BIO CONTROL PRODUCTS – ADJUVANTS

Staphyt Regulatory team can advise and support you in all regulatory procedures required to obtain approval of active substance and/or product authorisation for Plant Protection Product in Europe, as per Regulation (EC) No. 1007/2009 and as per specific Member State regulations.

Our expertise covers all areas related to substances and products evaluation: physico-chemistry, analysis, toxicology, ecotoxicology, E-fate, biological efficacy, residue, etc.

Staphyt Regulatory offers a complete range of services to successfully register your substances and products, in order to achieve a smooth and rapid entry to the market:

- Regulatory and strategic advice, Data Gap Analysis
- Coordination of conventional, bio-control 🌱 or adjuvants development
- Active substance (CA/MA) and preparation (CP/MP) dossiers
- Draft Registration Report (dRR)
- Biological Assessment Dossier (BAD)
- Risk assessment (Tox, Résidus, E-fate, Ecotox)
- Active substance technical equivalence dossiers
- Contact with Authorities
- Field and laboratory study management (GEP, GLP)
- Trial permit dossier
- CLP classification, SDS and Labelling
- Poison Control Center declaration
- and much more...

+ STAPHYT

A UNIQUE PARTNER FOR A GLOBAL MANAGEMENT OF YOUR DOSSIERS IN EUROPE AND BEYOND.

Staphyt Regulatory teams can rely on Staphyt Agrosociences network for an integrated management of your dossiers including carrying out the trials required for registration (GEP and GLP): efficacy, selectivity, residues, processing, bee safety, operator exposure (OPEX)...



AGROCHEMICALS & BIOCONTROLS



FERTILISERS & BIOSTIMULANTS



BIOCIDES



REACH



HUMAN & VETERINARY MEDECINES



COSMETICS

For further information, do not hesitate to contact us:



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