



## International reach through local focus

### APC provides a complete service for product registration in the EU

#### APC can:

- ▶ Prepare a data gap analysis identifying necessary studies
- ▶ Develop a cost effective strategy for re-registration and new registrations
- ▶ Evaluate what uses can successfully be registered according to new EU risk assessments
- ▶ Advise on complex data protection issues
- ▶ Provide expert advice on specific national registration requirements
- ▶ Manage studies, including efficacy and residue trials
- ▶ Prepare, submit and follow-up dossiers
- ▶ Co-ordinate multi-company Task Forces



*For existing national registrations, submission of a complete dossier is usually required within 2 years after inclusion in Annex I. New applications can be submitted at any time after the data have been generated. Active substance data protection expires 5 years after Annex I inclusion for agrochemicals or 10 years after Annex I inclusion for biocides.*

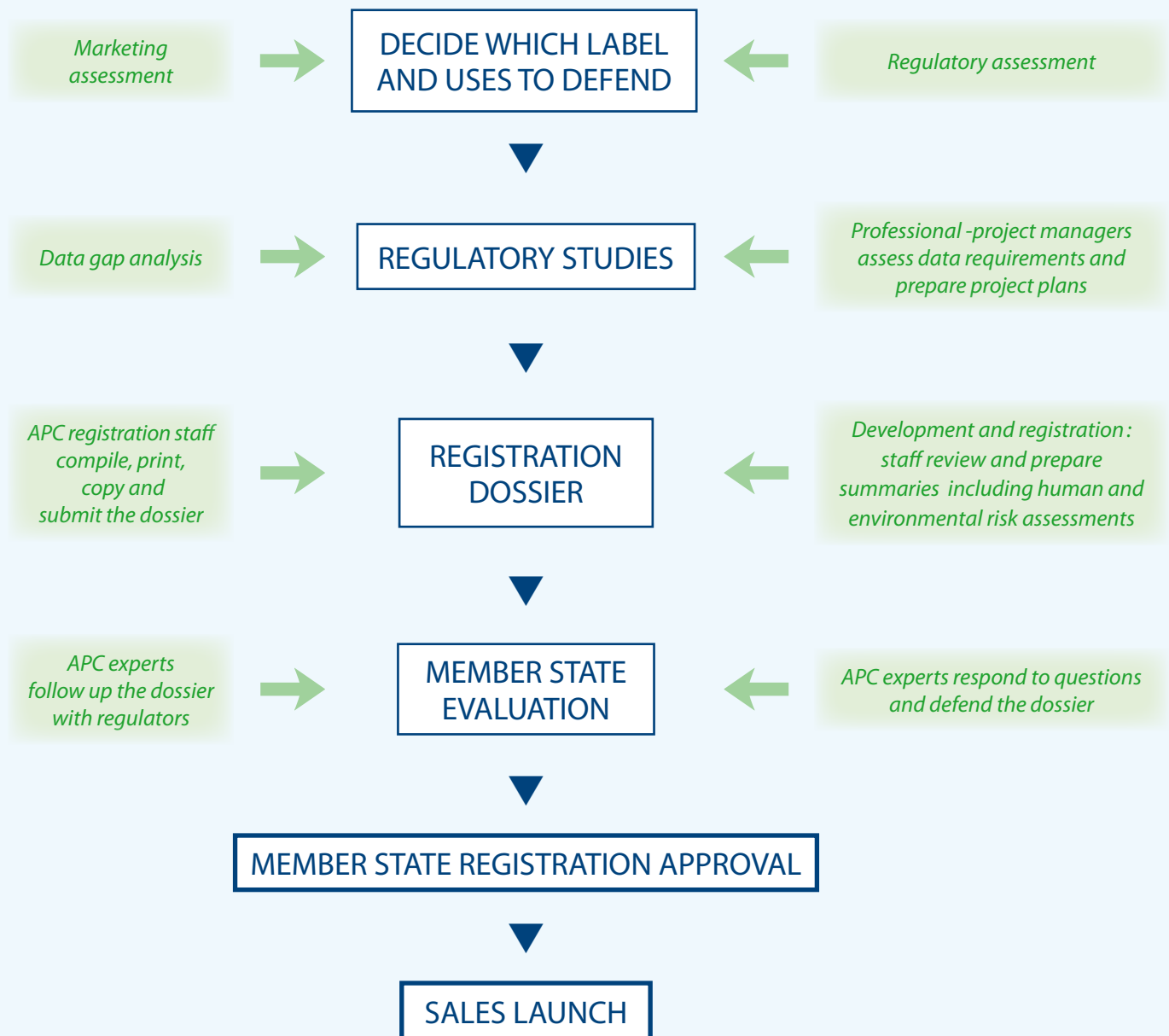
#### A product registration dossier should include:

- ▶ Physical chemical properties
- ▶ Validated methods of analysis
- ▶ Acute toxicology studies and human risk assessment
- ▶ Ecotoxicology studies and environmental risk assessment
- ▶ Efficacy trials
- ▶ Classification and labelling



our team provides a complete service for the plant protection, biocide & chemical industries

APC expertise ensures successful POST ANNEX I REGISTRATION in Member States



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or find our nearest office on [www.apc.eu.com](http://www.apc.eu.com)