

Title	European regulatory system for veterinary medicinal products
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## Project description:

The doctoral program will be implemented in a multidisciplinary perspective, geared toward applied research activity at the national regulatory institution, the Ministry of Health, and at the European Medicines Agency (EMA). It will aim at the integrated development of the following knowledge and skills:

1. reconstruct and interpret the national and supranational reference legal framework for the government policy sector of the veterinary medicinal product, including the secondary rank standards and the technical/application instructions that necessarily integrate the regulatory framework ("specifications" techniques, application guidelines, etc.) - <u>first phase of the project;</u>

2. participate in the government, in the organization of public administrations (both at the national level -Ministry of Health - and at the regional and local level - Zooprophylactic institutes/ULSS companies) through the implementation of innovative strategies strongly oriented towards users and the effectiveness of actions put in place, as well as the valorization of resources - <u>second phase of the project</u>;

3. support institutional planning also through the experimentation of innovative tools of the different governance models in a comparative key between policy sectors, between European, state and substate government levels and between national cases, which adequately take into account the opportunities offered by new information and communication technologies (ICT) - <u>third phase of the project</u>.

Hosting group(s) for the period abroad (tentative list, may change): European Medicines Agency- The Committee for Veterinary Medicinal Products (CVMP)