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Questions for Discussion on Authorisation Systems in the Region

Focal Points for Veterinary Products – Middle East

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Questions for Middle East Member Countries...

- 1. Do you have a joint or stand alone national competent Authority (human and vet or human-vet) in your country?
- 2. Do you have a legislation for authorisation/registration of veterinary medicines including antibiotics, antiparasitics and vaccines, diagnostic tests?
- 3. Do you have the possibility to take legal actions when the products on the market is not authorised/registered?



Questions for Middle East Member Countries...

- 4. Who delivers marketing authorisation in you country, what is the model or structure?
- 5. Do you have a national procedure for authorizing/registering veterinary medicinal products including antibiotics, antiparasitics and vaccines, diagnostic tests? Please describe briefly.
- 6. Do you recognize authorisation registration done by other country(ies)? If yes, what is the base for that?



Questions for Middle East Member Countries...

- 7. Do your countries have a "multinational" procedure? (more than one country preparing the evaluation together, which could lead one marketing authorisation which is valid to those countries who made the evaluation).
- 8. Do you recognize registration or authorisation delivered by sub-regional level?



Questions for Middle East Member Countries...

- 9. What is the base in your country to issuing a marketing authorisation or register a products?
 - Evaluation of the quality, safety and efficacy aspect of the products based on the dossier provided by the applicants? Something else?

Thank you for your contribution







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Thank you for your attention

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