Outcomes of the online questionnaire on the quality of veterinary products and pharmacovigilance

6th cycle regional webinar for OIE Focal Points for Veterinary Products,

Middle East
1. Does your country currently have a designated governmental body for registration of veterinary products, or is there work in progress to develop it?
If you have designated governmental body for registration of veterinary products; do you have a database or list for all authorized/registered veterinary medicinal product
Is the list of registered products publicly accessible?
Do you have pharmacovigilance (PHV) legislation implemented in your country, and a functioning pharmacovigilance system? Do you have a PHV guideline(s) that are used in the post-marketing activities?
What role(s), if any, do you think the OIE should play in defining the minimum requirements for a pharmacovigilance system for veterinary medicinal products?
Do you think that an OIE document describing how to set up a basic pharmacovigilance system would be beneficial for your country?
Do you consider that a pharmacovigilance system could be set up at a Regional -Sub-Regional level?
Conclusion

1- Most of the countries have/ in progress, a designated governmental body for registration of VP, and a database publically accessible.

2- Most of the countries have a PHV legislation and a functioning system with guidelines used in post marketing activities. The key elements of this system include strict control and reporting of sales in clinics, pharmacies and wholesalers.

3- Countries in the region are in favor that OIE should prepare a relevant chapter, in the codes/manuals, on the minimum requirements, for setting standards to a basic PHV system, which could be set, occasionally at regional/sub-regional level.
THANK YOU