



World Organisation
for Animal Health



Substandard and Falsified Veterinary Products Workshop

4 - 6 March 2025. Dar es Salaam, Tanzania.



The
Fleming
Fund



Eswatini Country Experience on Pilot VSAFE

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(VMP NFP)



REGULATION OF VMPs

❑ Animal Disease Act 7, 1965 (as amended) – Principal Act

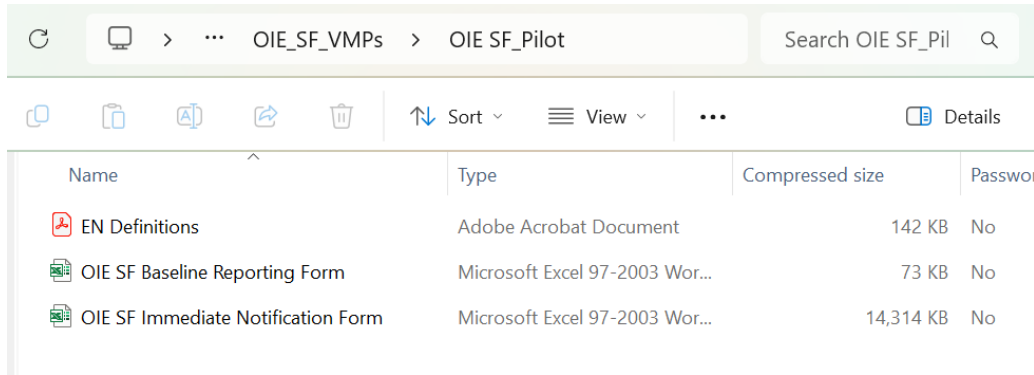
- Animal Disease (Amendment Act), 1990 – restriction of importation, distribution and usage of VMPs & mandating DVLS' oversight authority
- Prohibition of use of Anabolic Hormones, Thyrostatic Substances & Growth Promoters Regulations, 2006
- Regulation & Control of Veterinary Drugs and Medicinal Substances Regulations, 2012; which provide and/or specify for:
 - ✓ Recognition of certain Regulatory Authorities & authorizing the use of VMPs **sourced directly from these Territories**
 - ✓ The listing of authorized VMPs – Compendium of Registered Veterinary Medicinal Substances
 - ✓ The categorization/classification of listed VMPs to regulate retail/distribution and usage
 - ✓ The criteria for authorization of retailers and specifying the category of products they are allowed to import and/or distribute based on their technical capacity
 - ✓ empowering the DVLS **to conduct retailer inspections, and impose sanctions PRN**

SFVP SURVEILLANCE SYSTEM

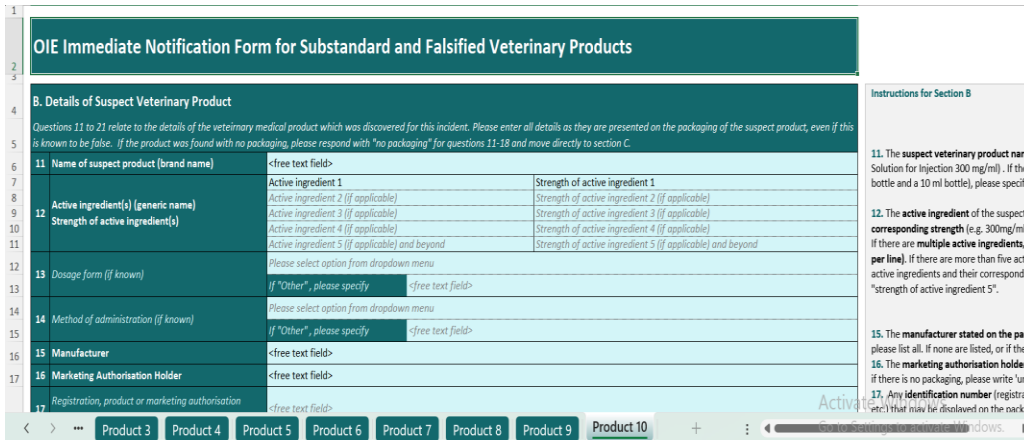
- Organizational Network Structure in relation to VMP Use Monitoring
 - Veterinary Directorate – Headquarters
 - ✓ National Focal Point for VMPs – based at Regional Level
 - Veterinary Field Services – At Regional level
 - ✓ Veterinarians ~ 8 in four regions
 - ✓ Animal Health Inspectors ~ 28 in 28 Subregional offices, across 4 regions
 - ✓ Frontline Veterinary Assistants ~ 240 in the four regions
 - Veterinary Public Health Unit – National VMP residue monitoring
 - Private Veterinarians
 - VMP Retail Industry and Animal Feed Industry Players

EXPERIENCE ON SFVP GLOBAL INFORMATION & ALERT SYSTEM PILOT PROJECT

• Pilot Phase 1



Name	Type	Compressed size	Password
EN Definitions	Adobe Acrobat Document	142 KB	No
OIE SF Baseline Reporting Form	Microsoft Excel 97-2003 Wor...	73 KB	No
OIE SF Immediate Notification Form	Microsoft Excel 97-2003 Wor...	14,314 KB	No



OIE Immediate Notification Form for Substandard and Falsified Veterinary Products

B. Details of Suspect Veterinary Product

Questions 11 to 21 relate to the details of the veterinary medical product which was discovered for this incident. Please enter all details as they are presented on the packaging of the suspect product, even if this is known to be false. If the product was found with no packaging, please respond with "no packaging" for questions 11-18 and move directly to section C.

11. Name of suspect product (brand name)	<free text field>	
12. Active ingredient(s) (generic name) Strength of active ingredient(s)	Active ingredient 1	Strength of active ingredient 1
	Active ingredient 2 (if applicable)	Strength of active ingredient 2 (if applicable)
	Active ingredient 3 (if applicable)	Strength of active ingredient 3 (if applicable)
	Active ingredient 4 (if applicable)	Strength of active ingredient 4 (if applicable)
	Active ingredient 5 (if applicable) and beyond	Strength of active ingredient 5 (if applicable) and beyond
13. Dosage form (if known)	Please select option from dropdown menu If "Other", please specify <free text field>	
14. Method of administration (if known)	Please select option from dropdown menu If "Other", please specify <free text field>	
15. Manufacturer	<free text field>	
16. Marketing Authorisation Holder	<free text field>	
17. Registration, product or marketing authorisation	<free text field>	

Instructions for Section B

11. The suspect veterinary product name Solution for Injection 300 mg/ml). If the bottle and a 10 ml bottle, please specify.

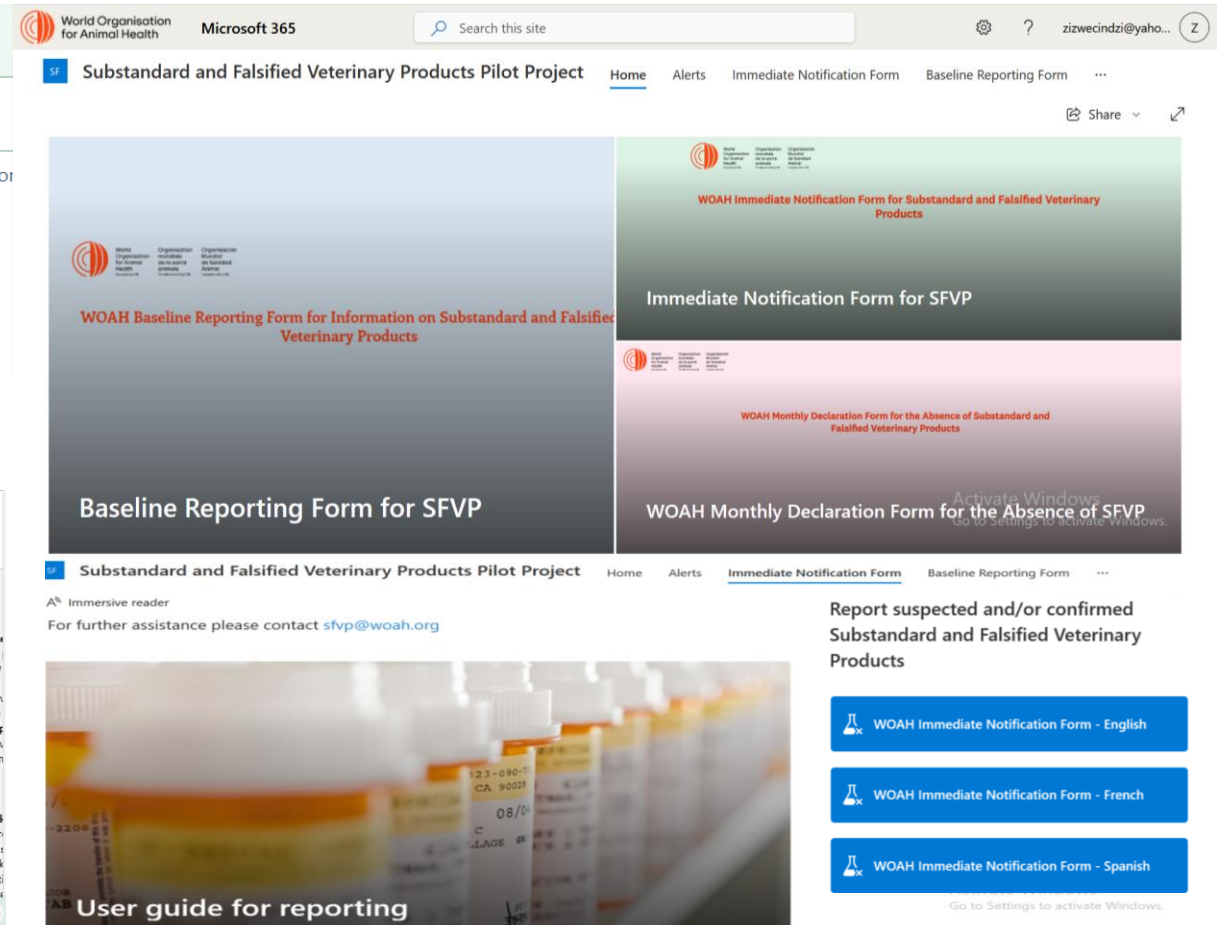
12. The active ingredient of the suspect corresponding strength (e.g. 300mg/ml). If there are multiple active ingredients, per line. If there are more than five active ingredients and their corresponding strength of active ingredient 5".

15. The manufacturer stated on the pack please list all. If none are listed, or if then

16. The marketing authorisation holder: if there is no packaging, please write "unk"

17. Any identification number (registration) that may be disclosed on the pack

• Pilot Phase 2 to VSAFE



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Substandard and Falsified Veterinary Products Pilot Project Home Alerts Immediate Notification Form Baseline Reporting Form

Share

WOAH Baseline Reporting Form for Information on Substandard and Falsified Veterinary Products

Baseline Reporting Form for SFVP

WOAH Immediate Notification Form for Substandard and Falsified Veterinary Products

Immediate Notification Form for SFVP

WOAH Monthly Declaration Form for the Absence of Substandard and Falsified Veterinary Products

WOAH Monthly Declaration Form for the Absence of SFVP

Report suspected and/or confirmed Substandard and Falsified Veterinary Products

WOAH Immediate Notification Form - English

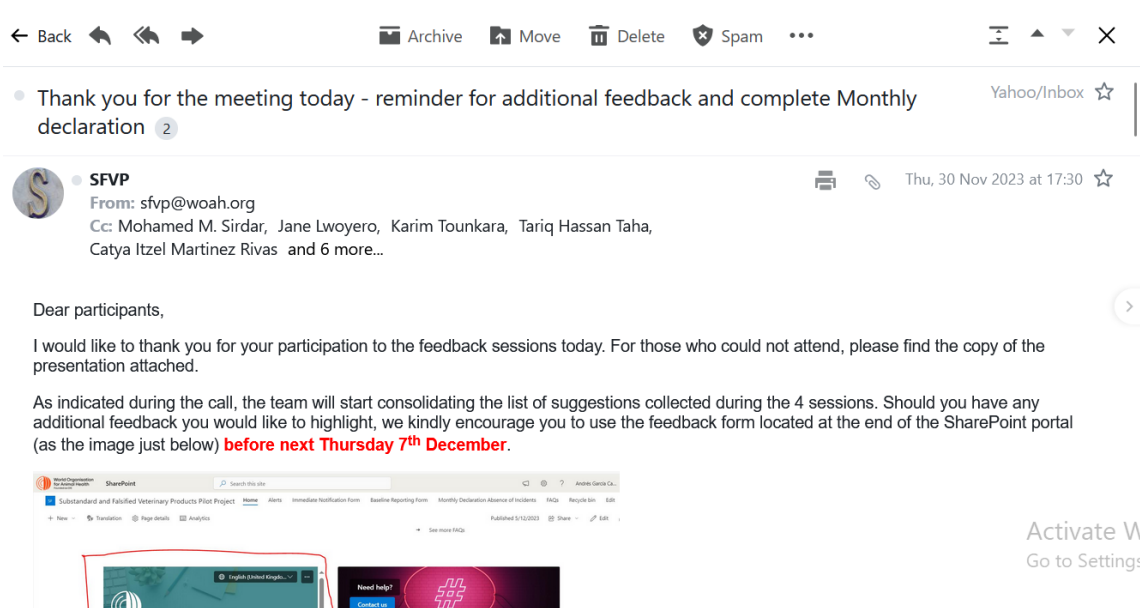
WOAH Immediate Notification Form - French

WOAH Immediate Notification Form - Spanish

Go to Settings to activate Windows.

EXPERIENCE ON SFVP GLOBAL INFORMATION & ALERT SYSTEM PILOT PROJECT

- An all inclusive consultative, collaborative and transparent process



CONSOLIDATED FEEDBACK RECEIVED DURING THE VIRTUAL MEETINGS FROM PARTICIPANTS OF PILOT PHASE 2 OF THE GLOBAL INFORMATION & ALERT SYSTEM FOR SUBSTANDARD AND FALSIFIED VETERINARY PRODUCTS

Executive Summary

On 29th and 30th Nov, WOA – SFVP Team organised four 2-hour meetings with participants of the pilot phase 2 for the Global Information & Alert System for Substandard and Falsified veterinary products. The sessions were conducted in the three WOAH official languages at different time zones in order to accommodate the largest number of participants. Sixty percent of members participated in the meeting, sharing their experiences and feedback of the current pilot. They also shared how they would like the system to evolve in the future. However, some internet difficulties impeded the attendance for some other Member participants hence all Members received the presentation and were asked to provide feedback by using the Form included at the end of the SharePoint platform developed for the pilot phase 2.

- Due consideration for Member States' specific needs

EXPERIENCES ON VSAFE PORTAL

- ❑ An excellent digital platform for reporting incidents of SFVPs, including **absence of incidents**
 - User friendly and hassle free
 - Non time consuming; Baseline & Immediate ~ 10-15min, Monthly ~ 1min
 - Set reporting timelines, with email reminders from WOAHS SFVP Team
 - Tabulated definitions and classification criteria for SFVPs
 - User Guides and easy communication platform with WOAHS SFVP Team
 - Provides region specific alert system on SFVPs incidents
 - Provides good and important links to Publicly Available VMP Databases & other Organisations

- ❑ Possible Future Considerations
 - Elaboration on links to other related resources, e.g. Relevant Laboratory Support Services

Bravo WOAHSFVP Team!!



Thank You