CALL FOR INFORMATION

The Standards and Trade Development Facility (STDF) is carrying out work to analyse how<u>Good Regulatory Practice (GRP)</u> can be used to improve the quality and effectiveness of sanitary and phytosanitary (SPS) measures in developing countries, in order to ensure health protection and facilitate safe trade. This work is based on the <u>WTO Agreement on the Application of Sanitary and Phytosanitary Measures (SPS Agreement)</u>.

Good regulatory practices are defined as internationally recognized processes, systems, tools and methods to improve the quality of regulations and ensure that regulatory outcomes are effective, transparent, inclusive and sustained (World Bank, 2015). Applying GRP means that SPS measures are effective and cost-efficient so that they achieve the intended outcomes. GRPs can include different processes and tools, such as consultations with the private sector and other stakeholders, Regulatory Impact Assessment (RIA), ex-post review of the implementation of SPS measures, etc.

The SPS Agreement confirms WTO Members' right to adopt SPS measures, based on science, to ensure that food is safe for consumers and to prevent the spread of pests or diseases among animals and plants, while seeking to prevent unnecessary trade disruptions. SPS measures include all relevant laws, decrees, regulations, requirements and procedures including, inter alia, end product criteria; processes and production methods; testing, inspection, certification and approval procedures; quarantine treatments including relevant requirements associated with the transport of animals or plants, or with the materials necessary for their survival during transport; provisions on relevant statistical methods, sampling procedures and methods of risk assessment; and packaging and labelling requirements directly related to food safety (SPS Agreement, Annex A).

Several of the provisions of the SPS Agreement encourage the use of GRPs, including use of international standards (Codex, IPPC, OIE), risk assessment, transparency, advance notifications on draft measures, etc. This survey aims to gather information on if and how SPS agencies in developing countries are applying GRPs to strengthen the development, implementation and review of SPS measures. Information obtained will be compiled in a short document, which will be shared with respondents, as well as the STDF Working Group. The findings will complement regular data collection exercises and work on measuring regulatory performance within OECD countries.

Officials of government ministries/agencies/departments responsible for food safety, animal and plant health in developing countries are kindly requested to complete this survey, before 13 October 2017. In case of questions, please contact STDFSecretariat@wto.org

* 1. Please indicate whether you work for a government agency responsible for:	
Animal health	
Food safety	
Plant health	
Trade / Economy	
Other (please specify)	

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Please complete the following questions from the viewpoint of the SPS area covered by your agency (i.e. food safety, animal or plant health).

* 2. Does your agency consult other government agencies in your country on the development of SPS measures?

This may include consultations on both primary and/or secondary legislation*, as well as diverse types of SPS measures, such as adoption of international (Codex, IPPC, OIE) standards, adoption of appropriate levels of protection (e.g. maximum residue levels for pesticides or veterinary drugs, maximum levels for chemical residues in food or feed), inspection or certification, setting administrative requirements and/or procedures domestically and/or the border, etc.

*Primary or principal legislation generally lays down policies and principles. Secondary legislation (also
known as subsidiary, subordinate or ancillary legislation) generally sets out the details of implementation
(i.e. it gives practical effect to the provisions of primary legislation) and is often collectively referred to by
the term "regulations".
Yes

(i.e. it gives practical effect to the provisions of primary legislation) and is often collectively referred to be the term "regulations".
Yes
○ No
On't Know

* 3. My agency shares draft SPS measures with other relevant parts		
Systematically		
Occasionally		
Rarely		
* 4. My agency shares draft SPS measures with other relevant parts	s of government for their (select as many boxes	as relevant):
Information		
Comment		
Review for coherence with existing legislation		
Review for consistency with other government guidelines		
Other (please describe)		
* 5. My agency consults the following government agencies on the c	development of SPS measures (select as many	boxes as relevant):
Animal health	Public health	
Food safety	Environment	
Plant health	Prime minister's office	
SPS Unit	Finance	
Economy / trade / industry	Justice	
Other (please specify)		
* 6. Guidelines / procedures to consult and/or coordinate with other SPS measures:	parts of government in preparation of	
Exist and are used systematically Are under	r preparation	
Exist and are used occasionally Do not ex	dist	
Exist and are used rarely		
7. Please provide any additional information related to consultation		
SPS measures. For instance, provide specific examples, and if pul relevant guidelines / procedures.	blicly available, provide web links to	
-		

* 8. To what extent, in your view, are international standards (Codex, IPPC, OIE) reflected in SPS measures in your area? This may include full or partial adoption of international standards in primary and/or secondary legislation.
Largely [>70%]
Moderately [40-70%]
Insufficiently [<40%]
On't know
* 9. Is there a regulatory requirement to consider relevant international standards (Codex, IPPC, OIE) in the development of SPS measures?
Yes
○ No
On't Know

* 11. During the process of developing SPS measures (i.e. prior to their formal adoption and entry into
force), does your government assess the risks to human, animal or plant life or health?
Yes
○ No
On't Know

	d/or plant life or health are assessed, e.g. when new legislation ard or existing risk assessment is prepared, new health risks	
Systematically		
Occasionally		
Rarely		
* 13. Are risk assessment principles and gui used in your area?	delines developed by the relevant international organizations	
Yes, fully	○ No	
Yes, to a moderate extent	On't Know	
Yes, to a limited extent only		
* 14. Who is responsible for assessing the ri	isks to human, animal and/or plant life?	
My agency		
Another government agency or other specialized unit (please specify)		
* 15. Written guidelines to assess the risks t		
Exist and are used systematically	Are under preparation	
Exist and are used occasionally	On not exist	
Exist and are rarely used		
	ion on how risks to human, animal and plant health are assessed	
relevant guidelines or methodologies.	n specific examples. If publicly available, provide web links to	
	n specific examples. If publicly available, provide web links to	
	n specific examples. If publicly available, provide web links to	
	n specific examples. If publicly available, provide web links to	
	n specific examples. If publicly available, provide web links to	
	n specific examples. If publicly available, provide web links to	
	n specific examples. If publicly available, provide web links to	
	n specific examples. If publicly available, provide web links to	

* 17. During the process of developing SPS measures (i.e. prior to their formal adoption and entry into force), does your government assess the expected impact on trade in order to ensure that SPS measures are not more trade-restrictive than required to achieve the appropriate level of sanitary or phytosanitary protection?			
Yes			
○ No			
Opn't Know			

40. The second of CDC		
18. The potential impacts of SPS measure	s on trade are assessed:	
Systematically		
Occasionally		
Rarely		
19. Who is responsible for assessing the p	otential impacts of SPS measures on trade?	
My agency		
Another government agency or other specialized	ed unit (please specify)	
² 20. Written guidelines / methodologies to a	ssess the potential impacts of SPS measures on t	rade:
Exist and are used systematically	Are under preparation	
Exist and are used occasionally	On not exist	
Exist and are rarely used		

22. Does your agency provide information and consult stakeholders (e.g. private sector, industry groups, consumer groups, general public) in your country or beyond on SPS measures in its area? For instance, consultations may take place through provision of information on new legislation, public hearings, and distribution of draft regulations for comments, etc.
Yes
○ No
On't Know

* 23.	Public consultation takes place:		
	Systematically		
	Occasionally		
	Rarely		
* 24.	My agency consults the following <u>domestic</u> stakeholders (select as many boxes as relevant):		
	Importers/Exporters/Traders		
	Industry associations		
	Primary producers		
	Consumer organizations		
	Civil society organizations		
	Other (please specify)		
25	My agency consults the following <u>foreign</u> stakeholders (select as many boxes as relevant):		
	Other governments		
	Importers/Exporters/Traders/Industry		
	Consumer organizations		
	Civil society organizations		
	Other (please specify)		
* 26.	Guidelines/procedures to consult stakeholders in your sector:		
\bigcirc	Exist and are used systematically		
	Exist and are used occasionally		
\bigcirc	Exist and are rarely used		
* 27.	* 27. Comments are received from stakeholders following consultations on draft SPS measures:		
	Systematically		
	Occasionally		
	Rarely		
\bigcirc	Never		

* 28. Comments received from stakeholders on draft SPS measures are considered by my agency:	
Systematically	
Occasionally	
Rarely	
Never	
* 29. Responses to comments received from stakeholders on draft SPS measures are published:	
Systematically	
Occasionally	
Rarely	
Never	
instance addressing methods, challenges, results, with examples if possible. If publicly available, provide web links to relevant guidelines or procedures.	

* 31. After an SPS measure in your area enters into force, is a review or evaluation carried out to assess
how the measure (individually or as a group) is being implemented, and whether it is achieving the intended objective?
Yes
○ No
On't Know

$\ensuremath{^{\star}}$ 32. The implementation of SPS measures is	reviewed and evaluated:	
Systematically (i.e. after a certain number of years	s)	
Occasionally		
Rarely		
* 33. Who is responsible for reviewing or evaluation the intended objectives?	uating the implementation of SPS measures in your area and wheth	er they are achieving
My agency		
Another government agency or specialized unit (p	please specify)	
		I
·	or evaluate the implementation of SPS measures:	
Exist and are used systematically	Are under preparation	
Exist and are used occasionally	On not exist	
Exist and are rarely used		
* 35. The findings of work to review and/or eva published:	aluate the implementation of SPS measures are shared or	
Systematically		
Occasionally		
Rarely		
Never		
* 36. How does your agency follow-up on work measures?	to review and/or evaluate the implementation of SPS	
27. Diagon provide any additional information	a halaw Far instance, describe have CDC massacrass and	
reviewed and indicate if stakeholders can red	n below. For instance, describe how SPS measures are quest a review (with examples of the findings of any such vaxuallable, provide web links to relevant guidelines or	
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8. Please indicat	te your country of	f work		\neg		
	ke to receive a sl			dings of this sur	vey, as well as ot	her news
om the STDF, pl	ease provide you	ır name and em	ail address.			

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THANK YOU VERY MUCH FOR YOUR FEEDBACK.

If you would like any additional information on this \underline{survey} or the STDF, please visit $\underline{www.standardsfacility.org}$ or contact STDFSecretariat@wto.org