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## PROFICIENCY TESTING CENTRE GENERAL PROTOCOL

# Introduction

The Proficiency Testing Centre (PTC), Pesticide Management Division (PMD) of National Institute of Plant Health Management (NIPHM), Hyderabad, India has been accredited in accordance with the standard ISO/IEC 17043:2010 (Conformity Assessment-General Requirements for Proficiency Testing) as Proficiency Testing Provider in the Field of Chemical Testing, by National Accreditation Board for Testing and Calibration Laboratories, an autonomous body under the Department of Science and Technology, India. The certificate number is P-0015, valid until 23<sup>rd</sup> May, 2018.

This protocol contains general procedures for Proficiency Testing (PT) Schemes organized by the Proficiency Testing Centre for *pesticide residues in Water, Fruits, Vegetables, Cereals and Pulses and Pesticide Analysis in Pesticide Technical and Pesticide Formulation samples.* 

# These Proficiency Testing schemes are open for all accredited laboratories and laboratories going for accreditation.

The aim of these PTs is to obtain information regarding the quality, accuracy and comparability of the data generated by Pesticide Residues Testing and Pesticide Formulation Testing Laboratories.

Participating laboratories will be provided with an assessment of their analytical performance and the reliability of their data – compared to the other participating laboratories.

It is mandatory to all the Pesticide Residue Testing Laboratories working under central sector scheme "*Monitoring of Pesticide Residues at National Level*" and involved in analysis of above samples as per the plan decided by the MPRNL scheme to participate in PT schemes on pesticide residues.

As per the directives of Government of India, It is mandatory for all regulatory State Pesticide Testing Laboratory (PTL), Regional Pesticide Testing Laboratory (RPTL) and Central Insecticide Laboratory (CIL) working in India to participate in PT schemes.

## Confidentiality

All information supplied by a participant to the PTC is treated as confidential and will not be shared to others (interested parties / regulatory body) without written consent from participants.

Each participant will be given unique ID (Laboratory code) for individual PT round. This will





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be shared to only respective participant at the time of dispatch of report.

# Communication

The official language used in NIPHM-PTs is English. Communication between participating laboratories during the test on matters concerning this PT exercise is not permitted. Proficiency Testing Centre has procedures to enable the participants to appeal against the evaluation of their performance in a proficiency testing scheme.

## **Announcement / Invitation Letter**

The announcement of the individual PT will be issued at least 2 months before the PT Item is distributed to the laboratories.

The announcement will be published on the NIPHM portal and additionally distributed via email to the interested laboratory (which has already submitted the details to NIPHM).

The announcement will contain an invitation letter, details on how to participate and where to find additionally-related documents, as well as some preliminary information on the specific protocol such as the tentative calendar, the name of the commodity expected to be used, and the tentative Target Pesticide List.

## Mode of Payment

Participant can pay the fees through Demand Draft in the name of National Institute of Plant Health Management Payable at Hyderabad.

or through Online ;

1.	Name of the Beneficiary	: NIPHM REVENUE ACCOUNT
2.	Name of the Bank	: State Bank of India
3.	Branch	: Budvel Branch, Teachers Colony, Budvel,
		Rajendranagar, Hyderabad - 500030, Telangana
4.	IFSC	: SBIN0012818
5.	Bank A/C No.	: 32917658917

## **Target Analyte / Product List**

This list contains Product and or analytes to be analyzed, is available with Plan of PT scheme on NIPHM Website.

## **Instruction to Participants**

For each PT, an instruction to participant will be sent along with the PT Item. This instruction will contain general instruction including handling and storage of PT items, formats used for acknowledgement of receipt of PT item, format for reporting the result, Contact details of PT Coordinator.





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This instruction also contains willingness of participatory laboratory regarding sharing the PT results with interested parties.

# **General procedures for reporting results**

Laboratories are responsible for reporting their results to the PTC within the stipulated deadlines. Do not keep any column blank (unattended) either you write result or write as not detected / below detection level / not analyzed / not participated or equivalent wording. The results should be submitted in two significant decimal places. Each laboratory must report only one result for each of the analytes detected in the Test Items, using the analytical procedure(s) that they would routinely use for each compound for monitoring purposes. Use appropriate units of measurement as specified in the instruction sheet.

In case of pesticide residue analysis for fruits, vegetables, cereals and pulses, one test item is intentionally treated with pesticides (Spike) and one is not (Blank). Both test items have to be analyzed by the laboratories and any pesticide detected in them shall be reported. In case of water only one water sample (Spike) will be given to participants.

In case of pesticide formulation analysis, individual samples are sent to individual laboratory with tentative range. Participant laboratories require reporting appropriate result. In some cases PT items may prepared synthetically in the laboratory in the range where normally the formulations are not available in the market.

# Methodology information

All laboratories are requested to provide information on the analytical method(s) they have used.

# **Estimation of Assigned value:**

In order to minimize the influence of out-lying results on the statistical evaluation, the assigned value is estimated using robust statistics as described in ISO 13528:2015 [Consensus Value from Participants].

## **Estimation of Standard Deviation for Proficiency Assessment (SDPA):**

In case of Residue Analysis, standard deviation for Proficiency Assessment will be calculated using Fit-For Purpose Relative Standard Deviation (FFP-RSD) approach.

The FFP-RSD is set at 25% of Assigned value based on the experience.

In case of Pesticide Formulation Analysis, the value of Standard Deviation for Proficiency Assessment ( $\sigma_{pt}$ ) determines on the basis of predictive models of the appropriate form of the Horwitz equation as described in ISO 13528:2015.





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However, NIPHM-PT organizer also reserves the right to employ other approaches on a caseby-case basis considering analytical difficulties and experience gained from previous proficiency tests.

# **Evaluation of Participant Performance**

**A) z-scores**: It has been calculated for all the participants using the assigned value and standard deviation stated above using following formula.

# B) Z' scores

If the criteria for acceptability of assigned value in terms of standard uncertainty of assigned value is not passing i.e. u(Xpt), > 0.3  $\sigma pt$ , , then uncertainty of assigned value is taken into account while estimating the performance score in terms of z' score.

## **Publication of results**

The PTC-PMD-NIPHM will publish a preliminary report, containing participant results and tentative z-score values for all parameter in the test sample, within 2 months from the deadline for result submission.

The Final Report will be published after the PTC-Panel has discussed the results. The final report may be published up to 3 months after the deadline for results submission.

## Feedback

After the distribution of the final report, participating laboratories will be given the opportunity to give their feedback to the organizer and make suggestions for future improvements.

## Disclaimer

The PTC-Panel of NIPHM retains the right to change any parts of this PTC – General Protocol based on new scientific or technical information. Any changes will be communicated in due course.





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# Laboratory Rights

After the Final Report has been sent, the laboratories will have the right to communicate the nonconformity of their result evaluation in written form. Any detected errors in the preliminary report should also be reported to the organizer. The organizer, assisted by the Scientific Committee, will decide upon any re-evaluation and will give a corresponding explanation.

\*\*\*\*\*\*\*\* End \*\*\*\*\*\*\*\*