



Call for participation:

The 2019 Global HPV LabNet Proficiency Study for HPV DNA Typing

Accurate and internationally comparable HPV DNA detection and typing methodology is an essential component in research on HPV vaccines and in effective implementation and monitoring. A WHO initiative established a Global HPV LabNet to support the worldwide implementation of HPV vaccines through improved laboratory standardization and quality assurance of HPV testing and typing methods to promote international comparability of results. The major methods for achieving progress towards this goal are developing international biological standards as well as preparing and validating proficiency panels to qualify methods.

We are now seeking international participation in an international HPV DNA testing and typing proficiency study. Laboratories that are or will be involved in HPV surveillance and/or vaccine development are particularly welcome.

Participant laboratories will be asked to perform HPV typing using one or more of their usual assays on the 44 challenges in this panel. This challenge is intended to evaluate assays that type HPV and is not appropriate for assays that detect HPV in general or grouped as high risk/low risk.

Composition of sample material

- 41 tubes containing purified whole genomic plasmids of **HPV 6, 11, 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 68a and 68b** in a background of human cellular DNA. Each sample may contain either single types or a mixture of HPV types at varying concentrations.
- 3 samples containing cell suspensions to allow evaluation of DNA extraction methods.

Participation fee

Participation in the proficiency study is subject to a participation fee per panel consisting of 44 samples: 1000 Euros for commercial entities and 500 Euros for academic entities. Laboratories that have outstanding payments from past Global HPV LabNet proficiency studies will need to clear their debts before their registration is accepted.

Data submission

The International HPV Reference Laboratory in Sweden is organizing this study in collaboration with the Swedish external quality assurance provider Equalis AB, who is responsible for management and distribution. Laboratories that have more than one assay are encouraged to provide results on each assay they commonly use. Data submitted will become the property of the organizers and may be analyzed for publication by the Global HPV LabNet either as an internal document or peer reviewed manuscript. All results will be handled in a coded anonymous fashion, with summaries grouped by method. The code linking data to originating laboratories will be kept confidential. Laboratories that provide data within the required time-frame will receive a copy of their own results and the summary data.

Scientific issues

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Registration

Application forms for the 2019 study can be found at:

<https://www.equalis.se/en/news/global-hpv-labnet-proficiency-study-2019/>

Preliminary dates

19 st of June 2019:	Registration for participation opens.
30 th of August 2019:	Registration for participation closes.
October 2019:	Dispatch of panels begins.

Participation, management and practical issues

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