Submission of method proposal for validation to ENGL SMV advisory group

*For each proposed method, please fill a separate form and send it by e-mail to engl-secretariat@jrc.ec.europa.eu. Where information is not available, please indicate with "**NA**". Where a point is considered not relevant, please indicate with "**IR**". When multiple options are offered for the answer, please mark with an "X" the option that better describes your method.*

1 Name of proposer and references

Name of proposer and institution:	
Address:	
Phone/fax:	
E-mail:	

2 General description of the proposed method

To be proposed, according to the criteria discussed within the AG SMV, the method should close an analytical gap and /or offer and added value in comparison with the current stock of recommended methods found in the GMOMETHODS database (http://gmo-crl.jrc.ec.europa.eu/gmomethods/). The criteria should also consider cost, speed, feasibility of implementation into the European system, and availability of positive material globally.

Type of method:

Simplex:	☐
Multiplex:	☐
High-throughput:	☐
Real-time:	☐
Other:	☐
If other specify:	
Qualitative:	☐
Quantitative:	☐
GMO-specific:	☐
Event-specific:	☐
Construct-specific:	☐
Element-specific:	☐
Taxon-specific:	☐
Plant-specific:	☐
Donor organism-specific:	☐

Genetic target(s) of method (please, describe the full name of the target(s) below:

GMO:

Taxon:

Sequences of primers and probes used (5-3'):

GMO detection

Primer forward:
Primer reverse:
Probe:
Detection chemistry:
Other primers:

Taxon detection

Primer forward:
Primer reverse:
Probe:
Detection chemistry:
Other primers:

Name of the method (if any):

Reference to peer-review publications or other method publication/report (please if possible send this document together with the form):

3 Justification for proposing the method

Is the proposed method closing an analytical gap*?

Yes: ☐
No: ☐

If yes which one?

**This criterion should allow method to cover:*

- *EU authorised GM events (EC/1829/2003) or under the 'Low Level Presence' (LLP) legislation (EC/619/2011)*
- *Unauthorised known GMOs*

Is the proposed method offering an added value (cost/time, efficiency etc.) in comparison with the methods proposed in the GMOMETHODS database (e.g. multiplex)?

Yes: ☐
No: ☐

If yes please specify which one:

Is the proposed method complying with ENGL methods acceptance criteria as specified in the document "Definition of minimum performance requirements for analytical methods of GMO testing" (http://gmo-crl.jrc.ec.europa.eu/guidancedocs.htm)?

Yes: ☐
No: ☐

If not what are the criteria not assessed?

If not what are the criteria assessed and not complying?

Is reference material available for this method?

Yes: ☐
No: ☐

If yes please select one of the following cases:

Reference material AOCS: ☐
IRMM: ☐
Other type of reference material: ☐
If other please specify:

Can the method be easily implemented on real-time PCR platform?

Yes on any type of real-time PCR machine: ☐
Yes, but specific type of real-time PCR machines have to be used: ☐
Please specify:
No, other type of machines are necessary: ☐
Please specify:

4 Other relevant information (if any)