Changes to legislation: This version of this Decision was derived from EUR-Lex on IP completion day (31 December 2020 11:00 p.m.). It has not been amended by the UK since then. Find out more about legislation originating from the EU as published on legislation.gov.uk. (See end of Document for details)

ANNEX IV

MODEL OF ANALYTICAL REPORT

Note: please compile an annex form for each sample tested

Parameter to be reported	Information provided
Name and address of the test laboratory ^a	
Test report identification code ^a	<<000>>
Laboratory sample identification code ^a	<<000>>
Size of laboratory sample ^a	X kg
In case of sample division: Number and size of analytical samples	X analytical samples of Y g
Number and size of test portions analysed ^a	X test portions of Y mg
Total DNA amount analysed ^a	X ng/PCR
DNA sequence(s) tested for ^a :	For each of the following provide reference to the method used and the average Ct number obtained Rice marker: 35S promoter: NOS terminator: CryIAb/CryIAc:
Other sequence(s) tested for:	Validation status: (e.g. inter-laboratory validated, in-house validated [please indicate according to which standard, guideline]) Description of DNA sequences detected (reference + target genes): Specificity of the method (screening, construct-specific or event-specific): Absolute Limit of Detection (copy number): Practical Limit of Detection (LOD related to the sample analysed), if determined:
Description of positive controls for target DNA, and reference materials ^a	Source and nature of the positive control and reference materials (e.g. plasmid, genomic DNA, CRM)
Information on the positive control ^a	Please indicate the amount (in ng DNA) of positive control analysed and the average Ct number obtained
Comments	
a Obligatory fields	1

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