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# Certified Reference Materials

# AOCS 0711-A4

Report of the certification process for

Ms1

Canola

Fourth Batch

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ISO 17034:2016
A2LA Certificate 3438.01

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## Abstract

This report describes the preparation and certification of the canola CRM AOCS 0711-A4 produced by AOCS Technical Services in 2017. The CRMs have been prepared according to ISO 17034:2016 and are intended to serve as control material for third party testing of canola for transformation events. The presence of Ms1 in the canola was verified using event-specific, qualitative PCR analysis by Eurofins-GeneScan, New Orleans, LA (an ISO 17025-accredited laboratory). AOCS 0711-A4 is available in 0.5 ml skirted screw-cap self-sealing tubes. The canola Ms1 DNA was provided by BASF Agricultural Solutions US LLC and was extracted from clean leaves. The leaf DNA extract sample shall be stored dry in a sealed container at +4 ˚C in the dark.


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## Acknowledgements

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## Glossary

| | |
|---|---|
| *AOCS* | American Oil Chemists' Society |
| *Conventional Crop* | Crop variety with no history of modern biotechnology and is produced through plant-breeding techniques that rely on selecting and mating parent plants possessing promising traits and repeatedly selecting for superior performance among their offspring |
| *DNA* | Deoxyribonucleic Acid |
| *Detection Limit* | Lowest level at which target DNA can be detected in a sample and be reliably tested by PCR methods. It is typically expressed as a percentage: the ratio of the number of modern biotechnology derived genomes to the number of crop genomes times 100 percent |
| *EC* | European Commission |
| *Genome* | The complete set of genes and DNA associated with an organism |
| *GMO* | Organism that has had genetic sequences modified using molecular-level techniques |
| *ISO* | International Organization for Standardization |
| *PCR* | Polymerase Chain Reaction: technique used to determine whether a sample of plant tissue contains a particular DNA sequence. PCR relies on primer sets that zero in on a particular target DNA sequence and a special DNA-copying enzyme (DNA polymerase) that makes enough copies of the target sequence for identification and measurement |
| *Qualitative PCR* | PCR methods that determine the presence or absence of a specific target DNA sequence at a particular level of detection |
| *Quantitation Limit* | Lowest level at which the amount of target DNA sequence in a sample can be reliably quantified. It is typically expressed as the ration of the number of transgenic genomes to the number of crop genomes times 100 percent |
| *Quantitative PCR* | PCR methods that estimate the relative amount of target DNA sequence in a mixture of DNA molecules |
| *Trait: Ms1* | Glufosinate ammonium herbicide tolerance and male sterility |


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## Introduction

Plant genetic modification is an extension of traditional plant breeding. It allows plant breeders to develop crops with specific traits including insect, disease, and herbicide resistance; processing advantages; and nutritional enhancement. An important component for identifying these new traits is a Certified Reference Material created from leaf, seed, or grain containing the new trait as well as a CRM created from the conventionally bred matrix. The European Commission has mandated that from 18 April 2004, a method for detecting a new event derived from transgenic technology and Certified Reference Material must be available before the EC will consider authorizing acceptance of a new crop derived from transgenic technology. Several nations outside Europe also require grain and ingredients to be labeled above a threshold level before accepting a shipment.

To meet the above regulatory requirements for GMO determination, AOCS 0711-A4 was manufactured from canola according to ISO 10734:2016 and in accordance with EC No 1829/2003. The CRM is available from AOCS.

## Materials and Methods

BASF Agricultural Solutions US LLC prepared the bulk material by taking source leaf material from plants which had been tested individually using several quality standards and was grown from seeds harvested from plants that had themselves passed the same criteria. Plants not meeting the quality standards were removed and destroyed. Leaf material was harvested from the plants which met the quality standards and frozen immediately and stored at -70 °C. The genomic DNA was extracted from leaves of one or more plants according to CTAB-based (Doyle JJ and Doyle JL, 1987) protocol. The integrity and concentration of the genomic DNA was determined by electrophoresis in a 1.0% agarose gel and ethidium bromide-staining and compared to lambda molecular weight standards by digital imaging quantification. The concentration measurement was done in triplicate, repeated in three different gels. No indications for physical degradation were apparent and the DNA migrated at positions higher than 40 Kb.


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BASF Agricultural Solutions US LLC delivered 2 mg of Ms1 canola to AOCS. Five (5) working samples of DNA, 10 μg each, were prepared from the composite and sent to Eurofins-GeneScan, New Orleans, LA (an ISO 17025-accredited laboratory) for event-specific, qualitative PCR analysis to screen for the presence of the intended event, Ms1. This testing was for presence confirmation as well as homogeneity purposes.

The leaf used to manufacture the Ms1 materials was shown to contain the Ms1 event as well as the absence of P35S, BXN, 2mEPSPS, cp4EPSPS , cp4EPSPS, NPTII, Barstar, and PAT sequences using PCR protocols at BASF Agricultural Solutions US LLC. The Ms1 canola leaf DNA was packaged by SGS-Midwest Seed Services in sterile, 0.5 ml skirted screw-cap self-sealing tubes in aliquots of 10 μg.

AOCS used the Random Number Generator function of Microsoft Excel to select samples for verification of gene presence, homogeneity, and to rule out degradation during packaging. Sample numbers AOCS 0711-A4: 10, 55, 59, 83, and 151 were sent to Eurofins-GeneScan, New Orleans, LA (an ISO 17025-accredited laboratory) for event-specific, qualitative PCR analysis to screen for Ms1 presence in the samples.

## Stability

Stability of these CRMs has been listed as 1 year from the certification date. The materials were sealed and stored in the dark at 4 °C, therefore not exposed to air and are expected to be stable for longer than the estimated expiration date. The stability of the leaf DNA extract material will be reevaluated annually. If the samples still test positive for the presence of the trait, the certificates will be extended.

## Results and Discussion

### Sample Homogeneity

The PCR data for the Ms1 homogeneity samples is presented in Table 1.


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| Table 1. Results of the homogeneity testing performed by Eurofins-GeneScan, New Orleans, LA (an ISO 17025-accredited laboratory) on the Ms1 bulk material 0711-A4 provided by BASF Agricultural Solutions US LLC | |
|---|---|
| **Sample** | **Ms1 Presence** |
| Homogeneity Sample 1 | Positive |
| Homogeneity Sample 2 | Positive |
| Homogeneity Sample 3 | Positive |
| Homogeneity Sample 4 | Positive |
| Homogeneity Sample 5 | Positive |

**Prepared Sample Verification**

After the bulk material was packaged, five (5) samples were identified by the Microsoft Excel Random Number Generator and sent to Eurofins-GeneScan, New Orleans, LA (an ISO 17025-accredited laboratory) for event-specific, qualitative PCR analysis. These results are presented in Table 2. This data confirms the presence of the Ms1 gene after the packaging of AOCS 0711-A4. These results are consistent with the homogeneity data presented in Table 1.

| Table 2. Results for the verification of AOCS 0711-A4 Ms1 canola material as tested by Eurofins-GeneScan, New Orleans, LA (an ISO 17025-accredited laboratory) with Ms1 event-specific, qualitative PCR analysis. | |
|---|---|
| **Sample** | **Ms1 Presence** |
| AOCS 0711-A4 10 | Positive |
| AOCS 0711-A4 55 | Positive |
| AOCS 0711-A4 59 | Positive |
| AOCS 0711-A4 83 | Positive |
| AOCS 0711-A4 151 | Positive |


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## References

Center for Environmental Risk Assessment GM Database
http://www.cera-gmc.org/?action=gm_crop_database

Eurofins-GeneScan; 2219 Lakeshore Drive, Suite 400, Metairie, LA 70122;Telephone: +1 504 297 4330 Toll Free: +1 866 535 2730 Fax: +1 504 297 4335 http://www.gmotesting.com

International Seed Testing Association, International Rules of Seed Testing: Seed Science and Technology Rules, 2012

ISO 17025:2005 and ISO 17025:2017, General Requirements for the Competence of Testing and Calibration Laboratories

ISO 174034:2016 (E) General Requirements for the Competence of Reference Material Producers

Regulation (EC) No 1829/2003 of the European Parliament and of the Council of 22 September 2003 on genetically modified food and feed; https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX%3A32003R1829&from=en

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