





LAB N° 0254 L

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## **TEST REPORT No. 2306081**

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Monopoli, 15/03/2023

Sample reception 13/03/2023

Analysis starting 15/03/2023

date

Dieta Shpk Fshati Kuqan 3001 Elbasan

CUSTOMER

Sample description

Delivered by: TNT 413671045

Sample description: Corn Flouer - Sample Code: 0099

Sampling procedure: By the Customer

Quantity of sample: 200 g each

Return of sample: No

TEST NAME	RESULT	U	U.M.	LOD	LOQ	R %	METHOD	LIMIT VALUE	LEGE ND	FINISHING DATE OF ANALYSIS	SEAT
GLUTEN	ND		mg/kg	3	5		1010-MI-7			15/03/2023	В

Analysis performed at:

B: Via San Donato, 25 - 70043 Monopoli BA)

## Legend

ND: It indicates that the analyte results as "Not Detected" through the analysis performed with the specified method or "<LOD" where it is indicated
U: Extended uncertainty, expressed in the same units of measurement as the result, calculated by using a coverage factor K = 2 (unless otherwise specified) for assuring a confidence
level close to 95%; otherwise, for microbiological tests and for airborne asbestos fibers tests, a confidence interval at the 95% probability level. For microbiological tests, a result derived
measurement uncertainty is reported, estimated according to ISO 19036 as standard uncertainty multiplied by a coverage factor k = 2 and an approximate level of confidence of 95%. The
combined standard uncertainty is assumed to be equal to the intra-laboratory reproducibility standard deviation. Water: for quantitative microbiological tests, the confidence interval of the
result calculated as indicated in the ISO 8199 standard is reported. LOD: Limit of Detection, defined as the lowest concentration of the analyte in a sample that can be detected, but not
quantified, under the specified conditions; expressed in the test report as "ND". LOQ: Limit of Quantitation, the lowest concentration of the analyte in a sample that can be determined, with
acceptable precision and accuracy. R%: Average percentage recovery (it is not used to correct the analytical data on pesticides, metals and mycotoxins). En: Revision "n" of the Test
Report which identifies the Amendment. The amendment replaces and cancels all previous versions of the Test Report.

Methods:

1010-MI-7 = AOAC Ridascreen Gliadin n°120601 2006



The description of the sample and/or any other references to the same are provided by the client. Analytical results shown in this test report refer uniquely to the sample tested. Where the sample has been provided by the customer, the results is referred to the sample as received. A partial reproduction of this document should be authorized by written approval from our Laboratory. If a statement of conformity with a specification or standard is reported, unless the decision norm is already contained in this specification or standard, or except for specific requests of the Customer, the Laboratory adopts as decision rule the direct comparison with the limit without considering measurement uncertainty.

All the documents useful for the traceability of analytical results is kept for four years in our archive.

Test report digitally signed and valid to all intents and purposes of law within the meaning of applicable legislation.

END OF TEST REPORT





