



LACTOSE

water content determination

Lactose is a milk sugar made of rennet whey. In its production, after separating the whey, you receive the so-called whey permeate that is thickened through evaporation and demineralized. This process is followed by crystallization which results in emergence of lactose crystals, with about 10% of water content, that are dried for example in dryers with a fluidized bed. After this stage of the engineering process, the water content in lactose does not exceed 0.5%. This value is controlled by way of drying control samples, e.g. with the use of MA/R or MA/X2 moisture analyzers manufactured by Radwag. Lactose is an ingredient in numerous products of the bakery (bread), confectionery (cakes, ice-creams), dairy (yogurts, kefir, cheese), pharmaceutical (drugs, supplements) as well as meat and food concentrate (fillers) industries. The water content analysis is therefore essential not only for engineering processes but also to obtain a final product of high quality.



The application note includes basic information for validation of the lactose drying method with the use of MA/R and MA/X2 moisture analyzers series by Radwag Wagi Elektroniczne. The application note may be the basis for elaborating own drying method with special regard to distinctive features of the product in question.



Lactose – water content determination

The method with the use of IR radiation

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TERMS

ACCURACY of determining water / dry matter content is the difference between the result of the water / dry matter content received in the moisture analyzer method and the result of the water / dry matter content received while drying the same sample through a reference method.

PRECISION is a degree of compliance between independent results of the test, received in specific conditions. The measure of precision is a standard deviation from a series of several measurements.

REFERENCE METHOD

The reference method parameters are usually specified in standards or other discipline-specific documents as the so-called guides. If such documents are unavailable, the drying temperature that does not cause the sample to change colors is used. Such an approach applies to previously dehydrated products and raw products.

SAMPLE PREPARATION

Before testing, the sample must be stored in a tightly sealed container. Before testing, mix the sample.

ACCESSORIES

Laboratory dryer, glass weighing vessels with a lid, AS 220.X2 analytical balance, laboratory spoon

METHOD DESCRIPTION

Place the sample with a mass of ca. 5 g in pre-dried glass weighing vessels. Specify the real mass of the sample in question with the use of the balance whose weighing accuracy is 0.1 mg (AS 220.X2). Put weighing vessels with the sample and lids in the temperature-controlled laboratory dryer. Dry sample at the temperature of 102°C for 3 hours. After this period, remove vessels and put into the desiccator until they cool down and weigh afterwards. Place samples in the laboratory dryer again and keep on drying them for 30 minutes. Cool them down and weigh again. Repeat the procedure until you obtain a stable sample mass or record the sample mass growth after drying.

RESULTS

Sample name	LACTOSE
Water content (%)	0.21
Standard deviation (%)	0.01

LACTOSE – WATER CONTENT ANALYSIS WITH THE MOISTURE ANALYZER

The water content testing with the use of the moisture analyzer (IR radiation) entails two phenomena: convection and radiation. The sample temperature rises from outer layers to the bottom of the sample. The temperature gradient in the sample structure minimizes through optimization of the thickness of the dried sample and drying temperature. Too high drying temperature may lead to surface burning of the sample, which is hard to diagnose when the sample color is dark.

SAMPLE PREPARATION

Before analyzing, store samples in sealed containers. Mix the sample before collection for testing.

ACCESSORIES

MA/R or MA/X2 moisture analyzer, laboratory spoon, disposable aluminum weighing pans.

METHOD DESCRIPTION

Set drying parameters presented below. Collect the sample with a mass of ca. 2 g and distribute a thin layer of the sample throughout the weighing pan. Lock the drying chamber manually or automatically.

DRYING PARAMETERS / RESULTS

Sample name	LACTOSE
Drying profile	Standard
Drying temperature	90°C
Sample mass (g)	~ 1.5 ÷ 2
End of analysis	Auto 1
Water content (%)	0.24
Standard deviation (%)	0.04
Analysis time \bar{x} (min)	1

ACCURACY OF THE MA/R ÷ MA/X2 METHOD

Sample name	LACTOSE
Water content (%) – Ref.	0.21 ± 0.01
Water content (%) – MA R/X2	0.24 ± 0.04
Analysis accuracy (%)	0.03

RESERVATION

The method in question has been verified by the Research Laboratory, yet the results do not include factors arising from diversity of tested samples, operators' personal skills as well as measuring capability used by moisture analyzer users. For this reason Radwag shall not be held responsible for drying parameters but they can be used to elaborate own drying method.

