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**Dried milk — Determination of content of
lactic acid and lactates**

Lait sec — Détermination de la teneur en acide lactique et en lactates

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Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

ISO 8069|IDF 69 was prepared by Technical Committee ISO/TC 34, *Food products*, Subcommittee SC 5, *Milk and milk products*, and the International Dairy Federation (IDF). It is being published jointly by ISO and IDF.

This edition of ISO 8069|IDF 69 cancels and replaces ISO 8069:1986, which has been technically revised.

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Foreword

IDF (the International Dairy Federation) is a worldwide federation of the dairy sector with a National Committee in every member country. Every National Committee has the right to be represented on the IDF Standing Committees carrying out the technical work. IDF collaborates with ISO in the development of standard methods of analysis and sampling for milk and milk products.

Draft International Standards adopted by the Action Teams and Standing Committees are circulated to the National Committees for voting. Publication as an International Standard requires approval by at least 50 % of the IDF National Committees casting a vote.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. IDF shall not be held responsible for identifying any or all such patent rights.

ISO 8069|IDF 69 was prepared by the International Dairy Federation (IDF) and Technical Committee ISO/TC 34, *Food products*, Subcommittee SC 5, *Milk and milk products*. It is being published jointly by IDF and ISO.

All work was carried out by the Joint ISO-IDF Action Team on *Lactose and lactate determination*, of the Standing Committee on *Main components of milk*, under the aegis of its project leader, Mr J Romero (US).

This edition of ISO 8069|IDF 69 cancels and replaces IDF 69:1987, which has been technically revised.

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Dried milk — Determination of content of lactic acid and lactates

1 Scope

This International Standard specifies an enzymatic method for the determination of the lactic acid and lactates content of all types of dried milk.

2 Terms and definitions

For the purposes of this document, the following terms and definitions apply.

2.1

lactic acid and lactates content

mass of substances determined by the procedure specified in this International Standard

NOTE It is expressed as milligrams of lactic acid per 100 g of non-fat solids.

3 Principle

A test portion of dried milk is dissolved in warm water. The fat and proteins are precipitated then filtered. The filtrate is treated with the following enzymes and biochemical substances, added simultaneously, but acting in sequence:

- a) L-lactate dehydrogenase (L-LDH) and D-lactate dehydrogenase (D-LDH), in the presence of nicotinamide adenine dinucleotide (NAD), to oxidize lactate to pyruvate and to convert NAD to its reduced form NADH;
- b) glutamate pyruvate transaminase (GPT), in the presence of L-glutamate, to transform pyruvate into L-alanine and to convert L-glutamate to α -ketoglutarate.

The amount of NADH produced is determined by spectrophotometric measurement at a wavelength of 340 nm, and is proportional to the lactic acid and lactates content.

4 Reagents

Use only reagents recognized analytical grade. The water used in the preparation of the enzyme solutions shall be of at least doubly glass-distilled purity and the water used for other purposes shall be glass-distilled or of at least equivalent purity.

4.1 Potassium hexacyanoferrate(II) solution, $c(\text{K}_4[\text{Fe}(\text{CN})_6] \cdot 3\text{H}_2\text{O}) = 35,9 \text{ g/l}$.

Dissolve 35,9 g of potassium hexacyanoferrate(II) trihydrate in water. Dilute with water to 1 000 ml and mix.

4.2 Zinc sulfate solution, $c(\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}) = 71,8 \text{ g/l}$.

Dissolve 71,8 g of zinc sulfate heptahydrate in water. Dilute with water to 1 000 ml and mix.

4.3 Sodium hydroxide solutions

4.3.1 Sodium hydroxide solution I, $c(\text{NaOH}) = 10 \text{ mol/l}$.

Dissolve 400 g of sodium hydroxide in water. Dilute with water to 1 000 ml and mix.

4.3.2 Sodium hydroxide solution II, $c(\text{NaOH}) = 0,1 \text{ mol/l}$.

Dissolve 4,0 g of sodium hydroxide in water. Dilute with water to 1 000 ml and mix.

4.4 Glycerol solution ($\text{C}_3\text{H}_8\text{O}_3$), with a volume fraction of 50 % glycerol.

4.5 Ammonium sulfate solution, $c[(\text{NH}_4)_2\text{SO}_4] = 3,2 \text{ mol/l}$.

Dissolve 422,84 g of ammonium sulfate in water. Dilute with water to 1 000 ml and mix.

4.6 Buffer solution, pH 10.

Dissolve 7,92 g of glycylglycine ($\text{C}_4\text{H}_8\text{N}_2\text{O}_3$) and 1,47 g of L-glutamic acid ($\text{C}_5\text{H}_9\text{NO}_4$) in about 80 ml of water. Adjust the pH to $10,0 \pm 0,1$ at $20 \text{ }^\circ\text{C}$ with sodium hydroxide solution I (4.3.1). Dilute with water to 100 ml and mix.

This solution may be kept for 3 months if stored in a refrigerator at between $0 \text{ }^\circ\text{C}$ and $+5 \text{ }^\circ\text{C}$.

4.7 Nicotinamide adenine dinucleotide solution (NAD).

Dissolve 350 mg of nicotinamide adenine dinucleotide ($\text{C}_{21}\text{H}_{27}\text{N}_7\text{O}_{14}\text{P}_2$) in 10 ml of water.

This solution may be kept for 4 weeks if stored in a refrigerator at between $0 \text{ }^\circ\text{C}$ and $+5 \text{ }^\circ\text{C}$. When the solution is being used, keep the vessel immersed in crushed ice.

4.8 L-Lactate dehydrogenase (L-LDH), from hog muscle suspension.

Dissolve 10 mg of L-lactate dehydrogenase suspension in 1 ml of glycerol solution (4.4). The pH of the obtained suspension should be about 7. The specific activity of the L-lactate dehydrogenase (L-LDH, EC 1.1.1.27) suspension shall be at least 5 500 units/ml at $25 \text{ }^\circ\text{C}$. If not, prepare another L-LDH suspension.

The L-LDH suspension may be kept for 12 months if stored in a refrigerator at between $0 \text{ }^\circ\text{C}$ and $+5 \text{ }^\circ\text{C}$. When the suspension is being used, keep the vessel immersed in crushed ice.

4.9 D-Lactate dehydrogenase (D-LDH), from *Lactobacillus leichmannii* suspension.

Dissolve 5 mg of D-LDH suspension in 1 ml of ammonium sulfate solution (4.5). The pH of the obtained suspension should be about 6. The specific activity of the D-lactate dehydrogenase (D-LDH, EC 1.1.1.28) suspension shall be at least 1 500 units/ml at $25 \text{ }^\circ\text{C}$. If not, prepare another D-LDH suspension.

The D-LDH suspension may be kept for 12 months if stored in a refrigerator at between $0 \text{ }^\circ\text{C}$ and $+5 \text{ }^\circ\text{C}$. When the suspension is being used, keep the vessel immersed in crushed ice.

4.10 Glutamate pyruvate transaminase (GPT), from pig heart suspension.

Dissolve 20 mg of GPT suspension in 1,0 ml of ammonium sulfate solution (4.5). The pH of the obtained suspension should be about 7. The specific activity of the glutamate pyruvate transaminase (GPT, EC 2.6.1.2) suspension shall be at least 1 600 units/ml at $25 \text{ }^\circ\text{C}$. If not, prepare another GPT suspension.

Add 1,0 ml of ammonium sulfate solution (4.5) to the 1 ml suspension with 20 mg of GPT and mix. Centrifuge this 2,0 ml suspension containing 10 mg of GPT/ml at a radial acceleration of 4 000 *g* for 10 min. Transfer 1,0 ml of the clear supernatant liquid and discard the remaining solution and pellet.

The suspension may be kept for 12 months if stored in a refrigerator at between 0 °C and +5 °C. When the suspension is being used, keep the vessel immersed in crushed ice.

4.11 Lithium L-lactate solution

Dissolve 50 mg of lithium L-lactate (C₃H₅O₃Li) in water. Dilute with water to 500 ml and mix.

4.12 Lithium D-lactate solution

Dissolve 50 mg of lithium D-lactate (C₃H₅O₃Li) in water. Dilute with water to 500 ml and mix.

5 Apparatus

Usual laboratory equipment and, in particular, the following.

- 5.1 **Analytical balance**, capable of weighing to the nearest 1 mg, with a readability of 0,1 mg.
- 5.2 **Glass beaker**, of capacity 50 ml.
- 5.3 **Graduated cylinder**, of capacity 50 ml.
- 5.4 **One-mark volumetric flasks**, of capacity 100 ml.
- 5.5 **Pipettes**, capable of delivering 0,02 ml, 0,05 ml, 0,2 ml, 1,0 ml and 2,0 ml.
- 5.6 **Graduated pipettes**, capable of delivering 5 ml and 10 ml, graduated in 0,1 ml divisions.
- 5.7 **Glass filter funnel**, of diameter about 7 cm.
- 5.8 **Filter paper**, medium fast grade, of diameter about 15 cm, free from lactic acid and lactates.
- 5.9 **Glass rod**.
- 5.10 **Plastic paddles**, capable of mixing the sample-enzyme mixture in the spectrometric cell.
- 5.11 **Spectrophotometer**, capable of measuring at 340 nm, equipped with cells of optical path length 1 cm.
- 5.12 **ParafilmTM**¹⁾.

6 Sampling

A representative sample should have been sent to the laboratory. It should not have been damaged or changed during transport or storage.

Sampling is not part of the method specified in this International Standard. A recommended sampling method is given in ISO 707 | IDF 50.

Store the sample in such a way that deterioration and change in composition are prevented.

1) ParafilmTM is an example of a product available commercially. This information is given for the convenience of users of this International Standard and does not constitute an endorsement by ISO or IDF of this product.

7 Preparation

7.1 Preparation of test sample

Transfer the test sample to a container with capacity about twice the volume of the sample and provided with an airtight lid. Close the container immediately. Mix the sample thoroughly by repeatedly shaking and inverting the container.

During preparation, avoid exposure of the test sample to the atmosphere in order to minimize adsorption of water.

7.2 Test portion

Weigh, to the nearest 1 mg, 1,0 g of the test sample in a 50 ml glass beaker (5.2).

7.3 Blank test

Carry out a blank test by proceeding as specified in 7.4 and 8.2, using all reagents but omitting the test portion.

7.4 Preparation of solution and deproteinization

7.4.1 Dissolve the test portion (7.2) in about 20 ml of water preheated to between 40 °C and 50 °C, while stirring with the glass rod (5.9) or suitable means. Transfer the contents of the glass beaker quantitatively to a 100 ml one-mark volumetric flask (5.4) by rinsing the beaker with water. Cool the contents of the flask to about 20 °C.

7.4.2 Add to the solution (7.4.1), in the following order, 5,0 ml of potassium hexacyanoferrate(II) solution (4.1), 5,0 ml of zinc sulfate solution (4.2) and 10,0 ml of sodium hydroxide solution II (4.3.2), swirling thoroughly after each addition. Dilute with water to the 100 ml mark. Mix thoroughly and allow the mixture to stand at room temperature for 30 min.

7.4.3 Filter through a filter paper (5.8), discarding the first fraction of the filtrate.

Use of a centrifuge is a suitable alternative to filtration.

8 Procedure

CAUTION — Avoid contamination, especially with perspiration.

8.1 Test to check the activity of reagents

8.1.1 Whenever a new batch of reagents (4.6 to 4.10 inclusive) is prepared, or when such reagents have been kept in a refrigerator without being used for more than 2 weeks, or when restarting analytical work after a period of analytical inactivity, or whenever other conditions may justify it, perform the following test for the recovery of lactates.

8.1.2 Pipette 10 ml of lithium L-lactate solution (4.11) into each of two 100 ml one-mark volumetric flasks (5.4). Pipette 10 ml of lithium D-lactate solution (4.12) into each of two other 100 ml one-mark volumetric flasks (5.4). Determine the L-lactic acid and lactates content and the D-lactic acid and lactates content of the solutions in the two pairs of 100 ml flasks, proceeding as specified in 7.4.2, 7.4.3 and 8.2.

8.1.3 Calculate the lithium lactate content, w_L , expressed in milligrams per litre, using one of the following equations:

a) for the L-lactate solution:

$$w_L = 341 \times A$$

b) for the D-lactate solution:

$$w_L = 346 \times A$$

where

A is the numerical value of the absorbance at 340 nm, calculated in accordance with 8.2.1 and 8.2.2;

341 is the numerical value of the factor after substituting the molecular mass of lithium L-lactate ($M_r = 96,1$) and the final volume ($V_1 = 2,24$ ml) in 9.1 when L-lactate recoveries are evaluated;

346 is the numerical value of the factor after substituting the molecular mass of lithium D-lactate ($M_r = 96,1$) and the final volume ($V_1 = 2,27$ ml) in 9.1 when D-lactate recoveries are evaluated.

8.1.4 Taking into consideration the purity of the lithium L-lactate and lithium D-lactate used to prepare the solutions, the recovery of lithium L- or D-lactate from any of the flasks (8.1.2) shall be within the range $100 \% \pm 5 \%$. If the recoveries are not within this range, check the reagents, the operating technique, the accuracy of the pipettes and the condition of the spectrophotometer. Take the required action to obtain appropriate results. Repeat the test until satisfactory results are obtained.

8.2 Determination

8.2.1 Transfer using the required pipette (5.5) into the 1 cm cell of the spectrophotometer (5.11) according to the scheme in Table 1.

Table 1 — Procedure scheme

Pipette into spectrophotometer cells	Blank	D-Lactate standard	L-Lactate standard	Sample
Distilled water	1,000 ml	—	—	—
Standard (8.1.2)	—	1,000 ml	1,000 ml	—
Sample filtrate (7.4.3)	—	—	—	1,000 ml
Buffer solution, pH 10 (4.6)	1,000 ml	1,000 ml	1,000 ml	1,000 ml
NAD solution (4.7)	0,200 ml	0,200 ml	0,200 ml	0,200 ml
GPT suspension (4.10)	0,020 ml	0,020 ml	0,020 ml	0,020 ml
Mix the contents of the cell using a plastic paddle (5.10) or cover the 1 cm cell with parafilm (5.12) and invert several times. After mixing, leave the cell and its contents for 5 min before measuring the absorbance (A_{b0} and A_{s0}) against water at a wavelength of 340 nm.				
L-LDH suspension (4.8)	0,020 ml	—	0,020 ml	0,020 ml
D-LDH suspension (4.9)	0,050 ml	0,050 ml	—	0,050 ml
Exactly 45 min after mixing, measure the absorbance of the test solution (A_{b45} and A_{s45}) against water at a wavelength of 340 nm.				
Leave the cell again and after exactly 60 min after mixing measure the absorbance of the test solution (A_{b60} and A_{s60}) again against water at a wavelength of 340 nm.				

The L- or D-lactic acid and lactates content may be determined separately by adding either L-LDH (4.8) or D-LDH (4.9).

When only L-lactic acid and lactate are measured, the absorbance may be measured after 30 min and 45 min, respectively, after mixing.

8.2.2 Calculate the net absorbance value, A , to be used in the calculation (9.1) by using the following equation:

$$A = [(A_{s60} - A_{s0}) - 4(A_{s60} - A_{s45})] - [(A_{b60} - A_{b0}) - 4(A_{b60} - A_{b45})] \quad (1)$$

where

A_{s60} is the numerical value of the absorbance of the test solution measured after 60 min in 8.2.1;

A_{s0} is the numerical value of the absorbance of the test solution measured in 8.2.1;

A_{s45} is the numerical value of the absorbance of the test solution measured after 45 min in 8.2.1;

A_{b60} is the numerical value of the absorbance of the blank test solution measured after 60 min in 8.2.1;

A_{b0} is the numerical value of the absorbance of the blank test solution measured in 8.2.1;

A_{b45} is the numerical value of the absorbance of the blank test solution measured after 45 min in 8.2.1.

A slowly proceeding side reaction may occasionally occur. The contribution to the absorbance caused by this side reaction may be eliminated by extrapolation to the absorbance at time zero.

When only L-lactic acid and lactate are measured (see 8.2.1), the absorbance may be measured after 30 min and 45 min, respectively. In this case, the equation is changed as follows:

$$A = [(A_{s45} - A_{s0}) - 3(A_{s45} - A_{s30})] - [(A_{b45} - A_{b0}) - 3(A_{b45} - A_{b30})] \quad (2)$$

where

A_{s30} is the numerical value of the absorbance of the test solution measured after 30 min in 8.2.1;

A_{b30} is the numerical value of the absorbance of the blank test solution measured after 30 min in 8.2.1.

8.2.3 If the increase in absorbance calculated according to 8.2.2 exceeds 0,500 units, repeat the procedures specified in 8.2.1 through 8.2.3, using an appropriate aqueous dilution of the filtrate from both the test portion (7.4.3) and the blank test solution (7.3).

9 Calculation and expression of results

9.1 Calculation

Calculate the lactic acid and lactates content, w_L , expressed as milligrams of lactic acid per 100 g of non-fat solids, by using the following equation:

$$w_L = \left(\frac{A \cdot M_r}{k \cdot l \cdot m} \right) \times \left(\frac{V_1 \cdot V_4 \cdot V_5}{V_2 \cdot V_3} \right) \times \left(\frac{100}{w_s} \right) \times 10^5 \quad (3)$$

where

A is the numerical value of the absorbance at 340 nm, calculated in accordance with 8.2.2;

M_r is the relative molecular mass of lactic acid ($M_r = 90,1$);

k is the molar absorption coefficient of NADH at 340 nm, i.e. $6,3 \times 10^6 \text{ cm}^2/\text{mol}$;

l is the optical path length of the spectrophotometric cells ($l = 1$ cm), in centimetres;

m is the mass of the test portion (7.3), in grams;

V_1 is the total volume of liquid in the spectrophotometric cell (see 8.2.1), in millilitres:

— when both L- and D-lactic acid and lactates are determined $V_1 = 2,29$ ml,

— when only L-lactic acid and lactates are determined $V_1 = 2,24$ ml,

— when only D-lactic acid and lactates are determined $V_1 = 2,27$ ml;

V_2 is the volume of the filtrate (see 7.4.3) in the spectrophotometric cell (see 8.2.1), in millilitres;

V_3 is the volume of the filtrate (see 7.4.3) taken for the dilution (see 8.2.3), if appropriate;

V_4 is the volume of the obtained solution in 7.4.2 (i.e. $V_4 = 100$ ml), in millilitres;

V_5 is the volume to which the test solution was diluted (see 8.2.3), if appropriate, in millilitres;

w_s is the non-fat solids content of the sample, expressed as mass fraction in percent.

NOTE The determination of fat is not part of the method specified in this International Standard. A recommended method for the determination of fat in dry milk is given in ISO 1736.

9.2 Expression of results

Express the test results in whole figures.

10 Precision

10.1 Interlaboratory test

The values for the repeatability and reproducibility were derived from the result of an interlaboratory test carried out in accordance with ISO 5725²⁾.

Details of the interlaboratory test on the precision of the method have been published (see Reference [5]). The values derived from this interlaboratory test may not be applicable to concentration ranges and matrices other than those given.

10.2 Repeatability

The absolute difference between two independent single test results, obtained using the same method on identical test material in the same laboratory by the same operator using the same equipment within a short interval of time, will in not more than 5 % of cases be greater than the following:

- a) for an arithmetic mean of lactic acid and lactates content ≤ 60 mg per 100 g of non-fat solids: 10 mg/100 g;
- b) for an arithmetic mean of lactic acid and lactates content > 60 mg per 100 g of non-fat solids: 15 % (relative) of the arithmetic mean.

2) ISO 5725:1986, *Precision of test methods — Determination of repeatability and reproducibility for a standard test method by inter-laboratory tests* (now withdrawn) was used to obtain the precision data.

10.3 Reproducibility

The absolute difference between two single test results, obtained using the same method on identical test material in different laboratories with different operators using different equipment, will in not more than 5 % of cases be greater than the following:

- a) for an arithmetic mean of lactic acid and lactates content ≤ 100 mg per 100 g of non-fat solids: 15 mg/100 g;
- b) for an arithmetic mean of lactic acid and lactates content > 100 mg per 100 g of non-fat solids: 20 % (relative) of the arithmetic mean.

11 Test report

The test report shall specify:

- a) all information necessary for the complete identification of the sample;
- b) the sampling method used, if known;
- c) the test method used, with reference to this International Standard;
- d) all operational details not specified in this International Standard, or regarded as optional, together with details of any incidents which may have influenced the test result(s);
- e) the test result(s) obtained and, if the repeatability has been checked, the final quoted result obtained.

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Annex A (normative)

Good laboratory practice (GLP) rules for the performance of enzymatic analyses

A.1 Introduction

Good Laboratory Practice rules for enzymatic analyses are less well known than those for other chemical analyses. Attention needs to be directed to such rules in order to obtain results with a satisfactory accuracy and precision. Prior to analyses, read the GLP rules described below.

A.2 Reagents

A.1.1 Use only enzymes of the prescribed grade (specific activity, concentration, contaminants with enzymatic activities, solvent).

A.1.2 Use only coenzymes of the prescribed grade (purity grade, salt or acid form, contaminants).

A.1.3 All reagents other than the enzymes and the coenzymes shall be of analytical grade.

A.1.4 The water for the preparation of the enzyme solutions and the other reagents shall be doubly glass-distilled.

A.1.5 The water for the preparation of the sample solutions shall be glass-distilled or deionized.

A.1.6 Store reagents and enzyme suspensions/solutions according to the instructions (usually between 2 °C and 8 °C).

A.1.7 Do not freeze enzyme suspensions.

A.1.8 When the expiry date of a reagent has been exceeded, either discard the reagent or check the efficiency of this reagent by examining standard solutions with varying amounts of analyte. The obtained absorbances shall be proportional to the concentrations.

A.1.9 Buffer solutions taken from the refrigerator shall be warmed up to room temperature before being added to the assay mixture.

A.3 Photometric and spectrophotometric cells

A.3.1 Use glass or plastic cells with an optical path length of 1 cm.

NOTE Plastic cells have the following advantages over glass cells:

- a) they are cheap (disposable);
- b) greater numbers of analyses are possible;
- c) within one batch, plastic cells agree very well with regard to the absorbance.

A.3.2 Whenever a new batch of cells is put into use, check their optical path length against that of a precision cells (e.g. quartz cell), as follows.

Fill the precision cell and plastic cells with water and measure the absorbance (A_1) of each cell against water. Fill the cells, after rinsing, with a solution of NADH (approximately 0,15 mg/ml) and again measure the absorbance (A_2) against water. For both the precision cell and the plastic cells, calculate $A_2 - A_1$. If the difference ($A_2 - A_1$) between the two cell types exceeds 0,5 % of the net absorbance measurement for the precision cell, calculate the average percentage difference and take this into account for the path length, l , in Equation (3).

A.3.3 Always use clean and unscratched cells. Dry or clean the optical sides of the cells with a soft tissue only.

A.3.4 It is advisable not to measure the absorbance of the sample test cells against that of the blank test cell, since no information will be obtained about the order of magnitude of the absorbance of the blank test itself. Measure the absorbance of both the sample and the blank test cell against water and calculate the difference.

A.3.5 Do not measure the absorbance of a sample or blank test cell against an empty cell (because of light diffusion).

A.3.6 Mix the contents of the cell with a plastic paddle or by sealing the cell with parafilm and gently swirling.

A.3.7 Remove air bubbles from the walls of the cells using a paddle. Avoid scratching the optical side of the cell.

A.3.8 Always use the same kind of cells for the measurement of the sample test and blank test.

A.3.9 Always place glass or quartz cells in the same position in the cell holder. For this purpose, mark one optical side of the cell.

A.4 Photometers and spectrophotometers

A.4.1 Use a spectrophotometer (bandwidth ≤ 10 nm), a filter photometer provided with an interfering filter (bandwidth ≤ 10 nm), or a spectrum line photometer equipped with a mercury vapour lamp. Measurements carried out using a spectrophotometer or filter photometer shall be made at the absorption maximum of NADH or NADPH, i.e. 340 nm. Those carried out using a spectral line photometer with a mercury vapour lamp shall be made at 365 nm or 334 nm.

NOTE The molar absorption coefficients of NADH and NADPH measured at 334 nm, 340 nm and 365 nm are as follows:

- a) NADH and NADPH at 334 nm (Hg): $6,18 \times 10^6$ cm²/mol;
- b) NADH and NADPH at 340 nm: $6,3 \times 10^6$ cm²/mol;
- c) NADPH at 365 nm (Hg): $3,5 \times 10^6$ cm²/mol;
- d) NADH at 365 nm: $3,4 \times 10^6$ cm²/mol.

A.4.2 A linear relationship up to an absorbance of 2,0 shall exist between the absorbance and the concentration of NADH or NADPH. Check the linearity as follows:

- a) pipette 2,00 ml of distilled water into a cell and measure the absorbance A_0 against water;
- b) pipette 0,10 ml of NADH solution (0,5 mg/ml) into the cell; mix the contents of the cell and measure the absorbance A_1 .