





# Luminoskan Ascent®

**User Manual** 

User Manual Rev. 2.1; Dec. 2003, Cat. no. 1507520

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# **Symbols and Markings**

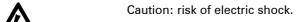
#### SYMBOLS USED IN THE LUMINOSKAN ASCENT

Power ON

Power OFF

Connection to the protective grounding system

#### WARNING MARKINGS USED ON THE INSTRUMENT



Caution: risk of personal injury to the operator or a safety hazard to the surrounding area. See the accompanying documentation.

CE compliance mark

#### WARNING MARKINGS USED IN THE DOCUMENTATION

Caution: risk of electric shock.

Caution: risk of personal injury to the operator or a safety hazard to the surrounding area.

Caution: risk of serious damage to the instrument, other equipment or loss of performance or function in a specific application.

Caution: biohazard risk.





# Luminoskan Ascent, Cat. no. 5300160 and 5300170 (with dispenser) User Manual Rev. 2.1; Dec. 2003, Cat. no. 1507520

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#### 1. Introduction

The Luminoskan Ascent (Fig. 1) designed and made by Thermo Electron is a microplate luminometer which offers versatility and flexibility for even the most demanding luminometric applications. The instrument covers the full range of glow and flash luminometric applications.



Fig. 1. Luminoskan Ascent

Ascent Software controls all the reader functions and provides easy assay optimization, flexible data handling and convenient report formatting. The software for the Luminoskan Ascent is a dedicated software for luminometric applications.

High sensitivity is one of the main benefits of the instrument. The Luminoskan Ascent can also be equipped with filters for luminometric applications.

Up to three reagent dispensers can be fitted on-board, making the reagent addition simple and highly accurate. The ability of the instrument to dispense and measure simultaneously enables the detection of flash luminescence reactions and rapid kinetic applications. For assays requiring temperature control, the instrument has an on-board incubator. Built-in orbital shaking speeds up reaction times and ensures effective mixing.

The robotic integration is simple and effective with the Luminoskan Ascent. The plate carrier allows convenient access for the robotic arm and Ascent Software is easy to integrate with robotic and HIS/LIMS systems. The Luminoskan Ascent is also fully compatible with Thermo Electron robotic plate handling devices, expanding the measurement capacity. For further information, contact your local Thermo representative.

#### 1.1 Intended use

- The Luminoskan Ascent is a high-quality microplate luminometer intended for laboratory research use by professional personnel. The Luminoskan Ascent is used to measure luminescence from suitable 1to 384-well plates mentioned in this manual. It also has incubation, shaking and reagent dispensing capabilities.
- 2. For verification of the entire system, it is recommended that Good Laboratory Practices (GLP) be followed to guarantee reliable analyses.
- 3. Use for self-testing is excluded.

## 1.2 Method descriptions

#### 1.2.1 Luminometric measurement principle

The filter slots 7 and 8 are reserved only for luminometric measurements. Filter slot 7 is empty for measurement and filter slot 8 is blocked to enable measurement of the PMT (photomultiplier tube) dark current. This feature is important to obtain optimal sensitivity. The light path from the measurement well to the first lens is protected by a light shield (Fig. 4.6a and Fig. 4.6b). The light shield is required with 96- and 384-well plates to avoid crosstalk in luminometric measurements. When plates are higher than 15 mm, the light shield must be removed. If the 384-well plate has a height of less than 15 mm, then an adapter must be used below the plate to raise the plate to the proper height. This adapter for low 384-well plates (Fig. 3.6:4) is included in the Luminoskan Ascent instrument and can also be ordered separately.



Caution: A luminometric measurement without a light shield may cause extra crosstalk.



Caution: The light shield must be removed with higher plates.



Note: The emission filter slots 7 and 8 are reserved only for luminometric measurements. Filter slot 7 is empty for measurement and filter slot 8 is blocked to enable measurement of the PMT (photomultiplier tube) dark current.

DO NOT install any filters into slots 7 and 8.

# 2. Instrument Layout

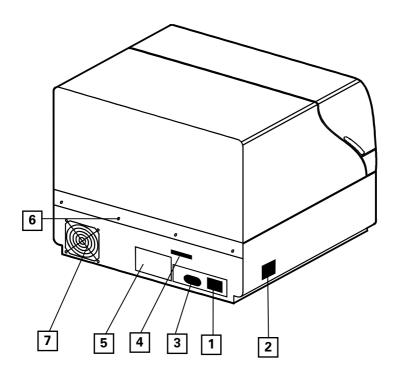
#### 2.1 Front view



- 1 Instrument housing
- 2 Dispenser and optics cover
- 3 Instrument chassis
- 4 Measurement chamber door
- **5** Power, busy and error indicator

Fig. 2.1 Luminoskan Ascent front view

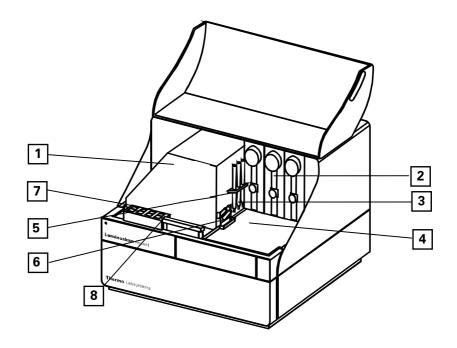
## 2.2 Rear view



- 1 Mains power supply socket
- 2 Power switch (ON/OFF)
- 3 Power fuses
- 4 Serial communication connector for the computer
- 5 Identification plate
- 6 Housing retaining screws
- 7 Cooling-air outlet

Fig. 2.2 Luminoskan Ascent rear view

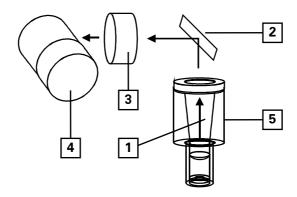
## 2.3 Internal view



- 1 Light cover for the optical unit
- 2 Dispensers (opt.)
- 3 Dispensing head (opt.)
- 4 Leakage tray (opt.)
- **5** Dispensing head holder (opt.)
- 6 Dummy plug
- 7 Control switches
- 8 Cover sensor

Fig. 2.3 Luminoskan Ascent internal view

## 2.4 Optical system



- 1 In the Luminoskan Ascent the light beam is strictly limited by a light shield to avoid crosstalk.
- 2 Mirror
- 3 An optional filter in the filter wheel
- 4 The photomultiplier tube (PMT) detects the light.
- 5 Light shield

Fig. 2.4 Principle of the optical system

#### 2.5 Control switches

The control switch box (Fig. 2.3:7) contains three rocker switches for priming and emptying dispenser tubings, one rocker switch for driving the plate carrier in or out, and a sensor (Fig. 2.3:8) to monitor that the dispenser cover is in the closed position in luminometric measurements. The priming control switches are only functional when the plate carrier is located outside the instrument.

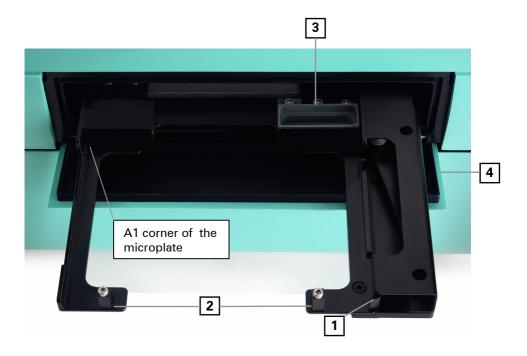
#### 2.6 Incubator

The incubator contains two heating element plates in the measurement chamber, one heating element plate under the microplate and another above it. The incubator only heats, but does not cool.

# 2.7 Dispensers

The optional dispensers, 1 to 3 dispensers from left to right, are located inside the instrument housing under the dispenser cover, as seen in Fig. 2.3. The dispensers consist of modular digital pumps with valves, syringes, tubing and dispensing heads. The dispensing heads have three alternative dispensing positions, one of these dispenses into the well in the measurement position.

## 2.8 Plate carrier



- 1 Positioning lever
- 2 Adjustable stoppers
- 3 Waste strip holder (opt., 4 wells) for the tip priming during the measurement session
- 4 Plate carrier door

Fig. 2.8 Plate carrier

## 3. Installation

## 3.1 Upon delivery

#### 3.1.1 Unpacking

The Luminoskan Ascent is packed in a specially designed shipping carton. Move the unpacked instrument to its site of operation. To prevent condensation, the instrument should be left in its protective plastic wrapping until the ambient temperature has been reached. Unpack the Luminoskan Ascent instrument and accessories carefully with the arrows on the transport package pointing upwards. Open the top of the package and lift the Luminoskan Ascent out of the shipping carton (Fig. 3.1). The following notes and instructions are sent with the instrument and are immediately available when you open the package:

- the packing instructions
- the packing list
- the Thermo Electron Warranty Certificate card
- the performance test reports
- the User Manual.



**Caution:** DO NOT touch or loosen any screws or parts other than those specifically designated in the instructions. Doing so might cause misalignment and will void the instrument warranty.



**Caution:** The Luminoskan Ascent weighs approximately 21 kg (46 lbs.) without dispensers and should be lifted with care. It is recommended that two persons lift the instrument together, taking the proper precautions to avoid injury.

Retain the original packing materials and shipping carton for future transportation. Also retain all the documentation provided with the instrument.

If you relocate your instrument or ship it for service, remember to:

1. Empty the dispenser(s) and remove the tube assembly.

- 2. Remove any loose items from the plate carrier, for example, adapters, plates and priming vessels.
- 3. Remove the power cable as well as the serial cable.
- 4. Replace the transportation lock.

For further information, see Section 8.4 Shipping the instrument (or items).



Fig. 3.1 Luminoskan Ascent

## 3.1.2 Checking delivery for completeness

Check the enclosed packing list against order. If any parts are missing, contact your local Thermo representative or Thermo Electron Oy.

## 3.1.3 Checking for damage during transport

Visually inspect the transport package, the instrument and the accessories for any possible transport damage.

If the carton has been damaged in transit, it is particularly important that you retain it for inspection by the carrier in case there has also been damage to the instrument.

Visually check all interconnections in the basic instrument. Check that there are no loose parts inside the instrument.

If any parts are damaged, contact your local Thermo representative or Thermo Electron Oy.

#### 3.1.4 Environmental requirements and noise

When you set up your Luminoskan Ascent, avoid sites of operation with excess dust, vibrations, strong magnetic fields, direct sunlight, draft, excessive moisture or large temperature fluctuations.

- Make sure the working area is flat, dry, clean and vibration-proof and leave additional room for cables, connections, controlling computer, printer, etc.
- Make sure the ambient air is clean and free of corrosive vapors, smoke and dust.
- Make sure the ambient temperature range is between +10°C (50°F) and +40°C (104°F).
- Make sure relative humidity is between 10% and 90% (noncondensing).

Leave sufficient space (at least 10 cm) at both sides of the instrument and at the back of the unit to allow adequate air circulation. Make space for the controlling computer on one side of the Luminoskan Ascent.

The Luminoskan Ascent does not produce operating noise at a level which could be harmful. No sound level measurements are needed after installation.



**Warning:** DO NOT operate the instrument in an environment where potentially damaging liquids or gases are present.

#### 3.1.5 Things to avoid

DO NOT smoke, eat or drink while using the Luminoskan Ascent. Wash your hands thoroughly after handling test fluids. Observe normal laboratory procedures for handling potentially dangerous samples. Use proper protective clothing. Use disposable gloves. Be sure the working area is well-ventilated.

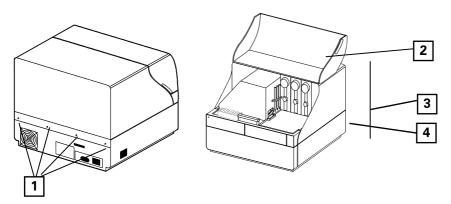
Never spill fluids in or on the equipment.

#### 3.1.6 Technical prerequisites

Place the instrument on a normal laboratory bench close to the mains power supply socket. The net weight of the unit is approx. 21 kg (46 lbs.).

The instrument operates at voltages of 100 – 240 Vac. The frequency range is 50/60 Hz.

## 3.2 Releasing the transportation lock

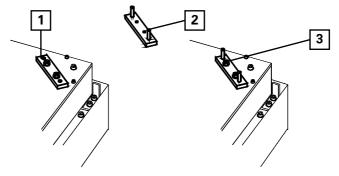


- 1 Cover retaining screws, 4 pieces
- 2 Dispenser and optics cover
- 3 Rear of the cover
- 4 Point to where cover lifted

Fig. 3.2a Removing the instrument cover

- 1. Remove the four cover retaining screws (Fig. 3.2a:1).
- 2. Open the dispenser and optics cover (Fig. 3.2a:2).
- 3. Lift the rear of the cover at first about 3 cm (Fig. 3.2a:3).
- 4. Lift the cover aside (Fig. 3.2a:4).
- 5. Undo the two screws (Fig. 3.2b:1) at the right rear corner of the measurement chamber.
- 6. Turn the locking piece upside down (Fig. 3.2b:2).

- 7. Fit the locking piece back with the fitting screws (Fig. 3.2b:3).
- 8. Refit the cover by first fixing the front corners.



- 1 Connected locking piece with two screws
- 2 Disconnected locking piece turned upside down
- 3 Reconnected locking piece

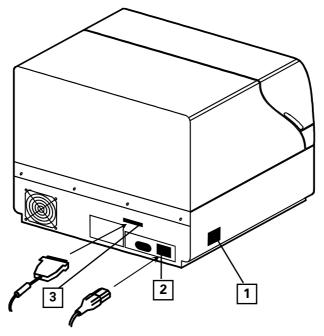
Fig. 3.2b Removing the transportation lock



Caution: If the locking piece is not fitted into its place, light may enter the measurement chamber and affect the results.

## 3.3 Power and computer connections

- 1. Ensure that the power switch (Fig. 3.3:1) is in the OFF position.
- 2. Connect the mains supply cable to the mains power supply socket (Fig. 3.3:2). The instrument box contains two different mains supply cables with North American and European types of plugs. Select the correct type used in your laboratory. If any other type of mains supply cable is needed, use only cables certified by the local authorities.
- 3. Connect the instrument to a correctly installed line power outlet that has a protective conductor also called ground or earth.
- 4. Connect the serial cable to the serial connector (Fig. 3.3:3) and secure it with the locking screws. Connect the other end similarly to the controlling computer.



- 1 Power switch (ON/OFF)
- 2 Mains power supply socket
- 3 Serial connector

Fig. 3.3 Power and computer connections



**Warning:** Always make sure the power switch on the instrument is in the OFF position and remove the mains power supply cable from the back of the instrument prior to any installation or relocation of the instrument.



**Warning:** Never operate your instrument from a power outlet that has no ground connection.

## 3.4 Installation of dispensers

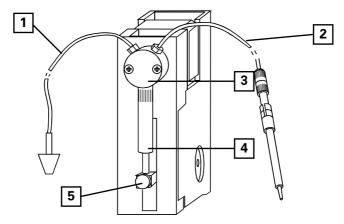
The optional dispensers 1 to 3 are installed in number order from left to right. The complete dispensing assemblies are packed with the accessories. The aspirate tubing (Fig. 3.4:1) is factory installed into the left hole of the valve. Ensure that the aspirate tubing is finger-tight. If necessary, turn the fitting another quarter to half turn using a 7.9 mm (5/16 in.) wrench. The aspirate tubing is used to fill the syringe with reagent. When using the dispensers, make sure that the aspiration tube end is completely submerged in the reservoir and there is a sufficient volume of the reagent in the reservoir (for all primings and the actual dispensing).

Fit the complete dispensing tube assembly (Fig. 3.4:2) into the right hole of the valve and tighten it finger-tight. Then turn the fitting another quarter to half turn using a 7.9 mm (5/16 in.) wrench. The dispensing tube is used to dispense reagent from the syringe into a microplate. Place the dispensing heads in the dispensing head holder on the left-hand side of the dispensers.

First push the plunger manually upwards into the upper position before tightening the plunger lock screw (Fig. 3.4:5). Ensure that the plunger lock screw is sufficiently tightened. Note that the plunger can be extremely firm.

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<sup>&</sup>lt;sup>1</sup> Instructions concerning the pump are reproduced from CAVRO XP3000 Modular Digital Pump Operators Manual made by Cavro Scientific Instruments, Inc., USA, 1994.



- **1** Aspirate tube assembly (incl. tubing and end weight)
- 2 Complete dispensing tube assembly
- 3 3-port valve
- 4 Dispensing syringe (1.0 ml) and plunger
- 5 Plunger lock screw

Fig. 3.4 Automatic dispenser unit

## 3.5 Installation of the drop plate

The instrument is supplied with a special drop plate (Fig. 3.5a). The drop plate is used to protect the instrument from damage caused by accidental dispensing without any microplate. If the user forgets to place a microplate onto the plate carrier but has the drop plate in place, the reagent will be dispensed into the drop plate, not inside the instrument. The drop plate is placed like an adapter into the plate carrier (Fig. 3.5b) and the microplate is placed onto the drop plate. The holding capacity of the drop plate is 19 ml of liquid.



**Note**: The drop plate cannot be used for bottom reading.



Fig. 3.5a Drop plate



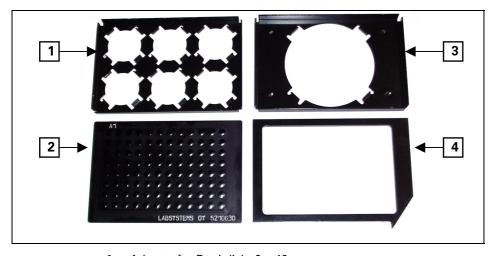
Fig. 3.5b Drop plate placed into the plate carrier

## 3.6 Plate adapters

Reading certain plate types with the Luminoskan Ascent instrument requires special plate adapters (Fig. 3.6). These plate adapters are required for the following reasons:

- 1. Raising the 384-well plate with a height of less than 15 mm to the proper height for luminometric measurement.
- 2. If the 384-well plate height is less than 13.5 mm, an adapter must be used to raise the plate for dispensing.
- Positioning of other sample vessel types than microplates (for example, Petri dishes or Terasaki plates) onto the plate carrier for measurement and dispensing.
- 4. Supporting flexible PCR plates and tubes on the plate carrier for measurement and dispensing.

When using any adapter, measure the height of the plate and the adapter together and ensure that this value is correctly entered into the Ascent Software plate template parameter file. Refer to the Ascent Software User's Guide for further information.



- 1 Adapter for Petri dish, 6 x 40 mm
- 2 Adapter for 96-well PCR plates
- 3 Adapter for Petri dish, 93 mm
- 4 Adapter for low 384-well plates

Fig. 3.6 Example of plate adapters

**Note:** If the orientation of the plate adapter is important, the correct orientation is printed on the adapter or the adapter has been designed in such a way that only one orientation is possible.

When the adapter is needed, place it into the plate carrier. Then place the sample vessel(s) on the corresponding adapter or a drop plate, select the correct plate template (marked "with adapter") from Ascent Software and run the instrument normally.



Note: Always remember to remove the plate adapter before using the instrument with any other plate types.

## 3.7 Adjusting the plate carrier

The plate carrier of the Luminoskan Ascent has been designed for plates with different footprints and to be robot compatible. The plate carrier has two adjusting knobs with two different orientations. These knobs have been set for standard plate types at the factory. The setting is also valid for the robotic 384-adapter but readjustment of the knobs will be needed with the plate types requiring special adapters (see Section 3.6). Fig. 3.7 shows how the knobs should be adjusted for special plate adapters. Both knobs should be rotated 90° clockwise after which all special adapters fit into the plate carrier.

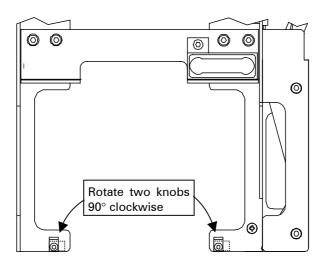


Fig. 3.7 Two plate carrier knobs rotated 90° clockwise for special plate adapters

## 3.8 Operational check

Switch the instrument ON. The instrument automatically performs a complete set of initialization tests or adjustments.

When the initialization tests and adjustments have been completed successfully, the indicator LED turns from yellow to green.

If anything fails in the initialization tests or adjustments, the indicator LED will turn red. In this case, switch the instrument OFF and ON again. If this failure is repeated, contact authorized technical service.

The instrument is ready for operation.

The instrument also performs different kinds of additional runtime hardware tests before each measurement run, like automatic blanking.

Because of the relative nature of luminometry, we recommend you use known samples or controls to verify proper instrument operation.

# 4. Operation

The Luminoskan Ascent is fully computer-controlled. Thermo Electron Ascent Software controls the reader functions and provides complete data handling and report formatting. For further details on how to use the software, refer to the Ascent Software User's Guide.



**Warning:** DO NOT use any other software than Thermo Electron software designed for this instrument. Use only valid combinations of the PC software and the instrument's internal FEPROM.



**Warning:** The Luminoskan Ascent instrument does not verify the logic flow of the received commands.

The instrument is equipped with a power switch (ON/OFF) and a three-color LED indicator. When the instrument is switched ON, the color indicates the state of the instrument:

The instrument is ready and waiting for a command.

	,
Orange	The instrument is busy, executing a command.
Red	The instrument has found an error, the error message is sent
	to the computer and the computer has not acknowledged it.

## 4.1 Switching on

When switched on, the instrument performs the initialization tests. When all the initialization tests and adjustments have been performed successfully, the indicator LED turns green and the instrument is ready to receive computer commands. The recommended warm-up time is 15 minutes, but the instrument will perform commands immediately after the initialization period.

## 4.2 Loading the microplate

The microplate is loaded onto the plate carrier of the instrument for measurement. The plate carrier is able to handle microplates of different sizes, therefore the free space in the plate carrier is clearly larger than the standard 96-well plate. The positioning lever (Fig. 4.2:1) in the plate carrier (Fig. 4.2) will automatically position the plate correctly into the upper left corner of the carrier when the plate is driven in.

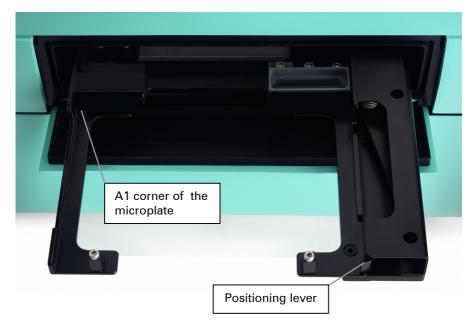


Fig. 4.2 Plate carrier

#### 4.3 Luminometric measurement

Luminometric reading is optimized for 96- and 384-well plates and top reading. To obtain the best sensitivity and the lowest crosstalk, a light shield (Fig. 4.6a) between the optics and the plate must be used. In luminometric measurements the dispenser cover must be closed.

Reading 1- to 48-well plates as well as bottom reading can cause increase in background signal and effect sensitivity in luminometric measurements, because the light shield cannot be used and most of these plates are transparent, which enables light leakage from one well to another.

A measurement function has several phases:

- The anode voltage of the PMT is set according to the selected value of the software.
- The filter slot 8 (= block) is held in place until the plate door is closed.
   The empty position/selected filter is then driven to the measurement positions. The filter slot 8 is also used for the blanking procedure to compensate for the possible PMT drift.
- 3. In the blanking procedure the instrument reads the PMT dark signal. If the bottom value drifts, the results are compensated. To obtain the most accurate results, the measurement time of the dark must be as long as the measurement time of the sample. The measurement time of the dark is divided into two parts. The first half is measured before the sample measurement and the second half after the sample measurement. The time used for the instrument blanking procedure can be selected from Ascent Software (refer to the Overview part of the Ascent Software User's Guide).



**Note:** Ensure that there are no items preventing the closure of the plate carrier door when measurement is started. DO NOT open the plate carrier door during measurement.



**Note:** DO NOT install any filters into the filter wheel positions 7 and 8.

#### 4.3.1 Luminometric scaling

The measured results are expressed as Relative Light Units (RLU). Scaling is a way to convert readings to show desired values.

- 1. Prepare a scaling reference solution and a blank solution and pipette several wells of blanks and the known concentration.
- 2. Measure the wells using the correct filter or an open slot.
- Calculate the average values of the measured reference values and blank values. The blank values should be very small compared to the reference values.
- 4. Calculate the scaling factor using calculated average values:

Factor = Known reference / (Measured reference - Measured blank)

#### Example

- The known concentration is 500 pmol/well.
- The calculated average of the measured reference wells is 1825 RLU.
- The calculated average of the measured blank wells is 0.2 RLU.
- Factor = 500 / (1825 0.2) = 0.274
- After scaling the measured result is: 0.274 x 1825 RLU = 500 pmol/well.

You can find the scaling entry in Ascent Software. The menu selection is **Setup | Filters**. Select the corresponding filter or None and key in the scaling factor. The measured results are expressed as Relative Light Units (RLU).

#### 4.4 Other functions

#### 4.4.1 Orbital shaking

The track movement system can perform the shaking action. The speed is adjustable from 60 to 1200 rpm (revolutions per minute) and the diameter of the orbital movement is adjustable from 1 to 50 mm.

Some combinations of speed and diameter would cause too high g-forces inside the well area resulting in spills inside the measurement chamber. Therefore, only certain combinations are available.

The following tables (Table 4.4.1a – Table 4.4.1e) show the recommended and the not recommended but available and unavailable speed and diameter combinations with different plate types. These tables are based on the liquid used being of low viscosity like water and the volumes being appropriate (the wells are not full). The gray area in the tables show the recommended (Recommended), the black area the not recommended (Possible), and the white area the prevented speed and diameter combinations (Not allowed).

n 1 3 5 7 9 11 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Diameter (mm)

Table 4.4.1a Shaking speeds for 48- (or more) well plates

Table 4.4.1b Shaking speeds for 24-well plates

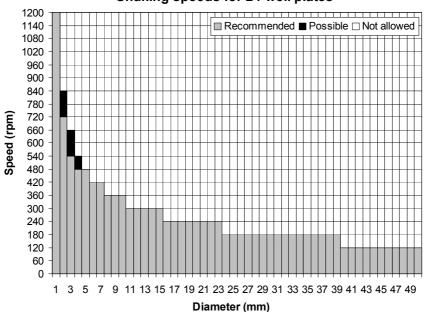


Table 4.4.1c Shaking speeds for 12-well plates

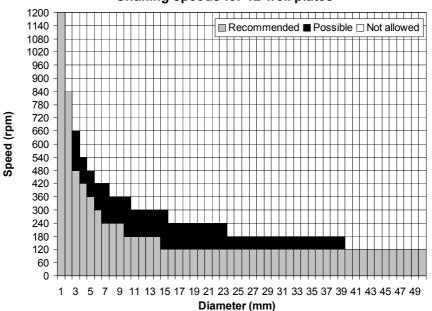


Table 4.4.1d Shaking speeds for 6-well plates

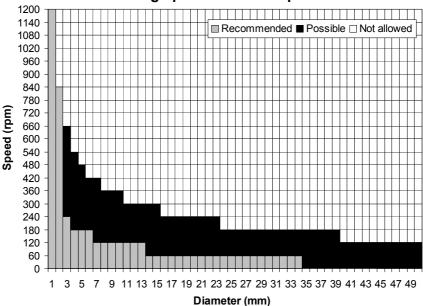
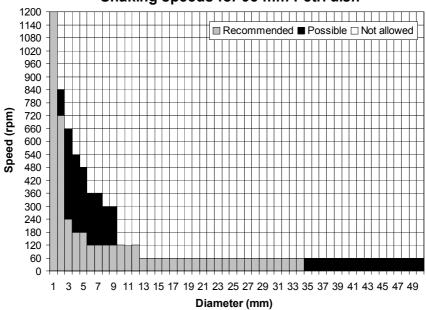


Table 4.4.1e Shaking speeds for 95 mm Petri dish



#### 4.4.2 Incubator

The incubator contains two heating element plates in the measurement chamber, one heating element plate under the microplate and another above it. Both heating element plates are temperature-controlled. The upper plate is slightly warmer than the lower plate to avoid condensation on the plate lid. The measurement chamber is large to adopt different plate formats and therefore extensive evaporation may cause some variations in the temperatures between the wells. Consequently, when using incubations with extended periods of time in the instrument, a plate lid is recommended.

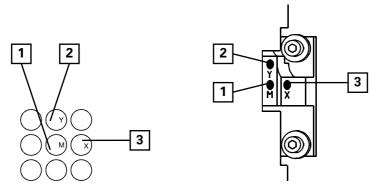
If the incubation period is long without any other important functions, the Luminoskan Ascent automatically changes the place of the plate within the measurement chamber to minimize temperature differences between the wells.

#### 4.4.3 Dispensers

The optional dispensers, 1 to 3 dispensers numbered from left to right, are located inside the instrument housing under the dispenser cover (Fig. 2.1:2). The dispensing heads have three alternative dispensing positions, one of these dispenses into the well in the measurement position (M). The positions are optimized for a 96-well plate (Fig. 4.4.3a). When the dispensers are not in use, the dispensing heads may be stored in the dispensing head holder, but the tip holes to the measurement chamber must be closed with dummy plugs.



**Note:** Ensure that the tip position in the instrument corresponds to that defined in Ascent Software.



- 1 *M position*: Dispensing head directed in the measurement well position. Dispensing into the well during the measurement step is possible with this position.
- Y position: Dispensing head directed in the well next to the measurement position in the Y direction. When this position is used for dispensing, an extra plate movement is carried out before the measurement step causing minor time delays.
- 3 *X position*: Dispensing head directed in the well next to the measurement position in the X direction. When this position is used for dispensing, an extra plate movement is carried out before the measurement step causing minor time delays.

Fig. 4.4.3a Dispenser tip positioning optimized for a 96-well plate

Before inserting the dispensing head into a dispensing position, prime the syringe and tubing to an external priming vessel. The instrument has no internal priming vessel for priming the syringe and tubing. You can find the priming instructions in the Ascent Software User's Guide. The minimum priming volume needed is 700  $\mu$ l and the recommended volume is 2700  $\mu$ l.

The Luminoskan Ascent also has control switches for priming the dispenser tubing. With these switches, priming can be carried out alternatively by using an external priming vessel or by using an empty microplate on the plate carrier as a priming vessel or a drop plate. When priming is performed with the dispensing heads installed into the M, X or Y positions, place the empty priming vessel into the plate carrier. Initiate priming by pressing the corresponding switch when the plate carrier is located outside the unit. Priming is carried out as long as the corresponding switch is pressed.



Note: The priming control switches are full ctional only when the plate carrier is located outside the instrument. DO NOT use priming vessels higher than the actual place intended to be used in the assay. Notice the dispensing height adjustment.

Priming with Ascent Software must always be performed with the dispensing heads removed from the M, X or Y dispensing positions.



**Note:** Never use liquids that can cause any precipitation, clotting or contain any mechanical particles with the automatic dispensers.

The instrument has a Prime Tip feature. If this function is selected in the **Dispense** or **Dispense** And **Measure** steps in Ascent Software, the dispenser dispenses about  $5~\mu$ l reagent into the tip priming vessels every time the instrument fills the syringe. This makes the volume of the first well equal to that of the others. We recommend you use the tip priming feature when the dispensing volumes are small, for example,  $5-20~\mu$ l. Notice that the **Execute by** command causes the syringe to fill.

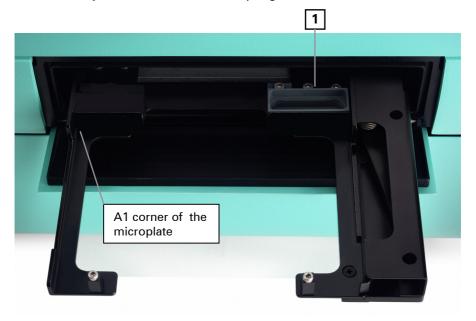


Fig. 4.4.3b Tip priming vessel (1) is a piece of a breakable strip

The recommended tip priming vessel consists of four wells of a breakable 96-well plate strip (Fig. 4.4.3b). There is a holder for the tip priming vessel in the right rear corner of the plate carrier. The four-well piece of a strip should be exchanged after about 300 tip primings.

You may need to adjust the dispenser speed. The default setting is for a water-based liquid. You can find the adjustments and selections in Ascent Software.

When dispensing is started, the liquid volume in the well should be less than half of the total well volume (for example, the volume should be less than 200  $\mu$ l in a typical 96-well plate).

#### 4.4.4 Dispenser head height adjustment

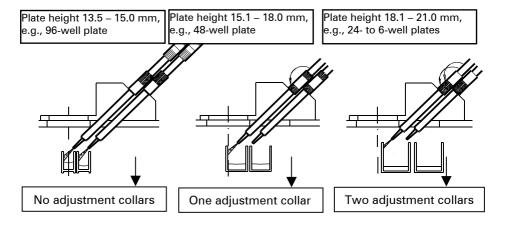


Fig. 4.4.4a Dispenser head height

The correct dispenser head height (Fig. 4.4.4a) is very important to avoid contaminating neighboring wells. The correct tip height is also important to prevent damaging the tip or the plate.



**Note:** Ensure that the dispenser tips are always correctly inserted sufficiently deep into their slots.

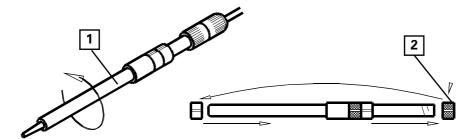
The plate height is one of the template parameters in Ascent Software and it is defined as the height of the uncovered well from the bottom of the plate, not the inside height of the well. The selected dispenser head height, the used plate and the template selected in Ascent Software must match to avoid problems. To see the heights of plates in Ascent Software, select **Setup** and edit **Plate Templates**.

Some 384-well plates are lower than the standard 96-well plates. If the plate height is less than 13.5 mm, an adapter must be used to raise the plate. Measure the height of the plate and the adapter together and enter this value to the plate template.



Caution: Plate manufacturers may change the dimensions of plates without any prior notice or change in order numbers. Check the dimensions when you start using plates from a new box.

The dispenser head height is adjusted with red adjustment collars by moving them from either side of the fixed stopper collar (Fig. 4.4.4b).



- 1 Dispenser head tube
- 2 Red adjustment collars

Fig. 4.4.4b Changing the position of the red adjustment collars

- 1. Remove the dispenser head tube (Fig. 4.4.4b:1) from the brass tube lock.
- 2. The red adjustment collars (Fig. 4.4.4b:2) should be moved from one side of the fixed collar to the other to select the correct dispenser head height.
- 3. Fit the dispenser head tube back into place.

#### 4.4.5 Chemical resistance of the dispenser <sup>1</sup>

The following table (Table 4.4.5) is intended to provide guidelines for compatibility with materials used in the fluid path of the dispensers. Compatibility information is based on charts provided by the material manufacturer. Cavro recommends that each laboratory determines compatibility for their respective applications.



Caution: Failure to determine compatibility of chemicals used in individual applications with the XP 3000, may result in damage to the dispenser and/or test results.

Plastic materials used in dispensers:

Polysulfone: Cross Flow Manifold Assembly in the aspirate syringe

Teflon (PTFE, TFE, FEP): tubing; valve plug, and seal

Kel F: valve body

Polypropylene: fittings for tubing, and dispensing tip



**Note**: Also take into account the chemical resistance of microplates.

#### Classification in the table:

No data available

0 No effect – excellent

1 Minor effect – good

2 Moderate effect – fair

3 Severe effect – not recommended

<sup>&</sup>lt;sup>1</sup> Instructions concerning the pump are reproduced from CAVRO XP 3000 Modular Digital Pump Operators Manual made by Cavro Scientific Instruments, Inc., USA.

Table 4.4.5 Compatibility chart of materials suitable with the dispenser

Solvent I	Polysulfone	Teflon	Kel F	Polypropylene
Acetaldehyde	_	0	0	0
Acetates	_	_	0	0
Acetic Acid	0	0	0	0
Acetic Anhydride	_	_	0	_
Acetone	3	0	0	0
Acetyl Bromide	_	0	_	_
Ammonia	0	0	-	0
Ammonium Acetate	_	0	_	_
Ammonium Hydroxide	-	0	0	0
Ammonium Phosphate	_	-	0	0
Ammonium Sulfate	-	_	0	0
Amyl Acetate	_	0	_	3
Aniline	_	0	0	0
Benzene	3	0	3	*
Benzyl Alcohol	_	0	0	0
Boric Acid	_	0	0	0
Bromide	_	0	0	*
Butyl Alcohol	2	0	0	1
Butyl Acetate	3	0	-	*
Carbon Sulfide	_	0	-	*
Carbon Tetrachloride	0	0	1	3
Chloroacetic Acid	_	0	0	_
Chlorine	_	0	1	3
Chlorobenzene	3	-	-	3
Chloroform	3	0	-	3
Chromic Acid	3	0	0	_
Cresol		0		*
Cyclohexane	0	0	_	3
Dimethyl Sulfoxide (DN	/ISO) 0	0	0	0
Ethers	_	0	_	**
Ethyl Acetate	3	0	_	0
Ethyl Alcohol	0	0	_	0
Ethyl Chromide		0	1	3

Continued

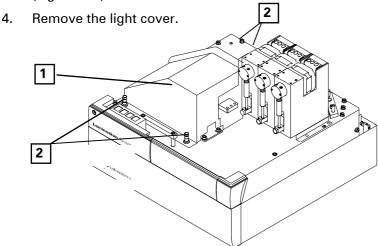
Solvent	Polysulfone	Teflon	Kel F	Polypropylene
Formaldehyde	0	0	0	0
Formic Acid	_	0	0	0
Freon	2	0	2	0
Gasoline	2	0	0	3
Glycerine	0	0	0	0
Hydrochloric Acid	0	0	0	0
Hydrochloric Acid (	conc.) 0	0	0	0
Hydrofluoric Acid	2	0	0	*
Hydrogen Peroxide	_	0	0	0
Hydrogen Peroxide	(conc.) –	0	0	0
Hydrogen Sulfide	_	0	0	0
Kerosene	2	0	0	0
Methyl Ethyl Ketone	e (MEK) 3	0	_	0
Methyl Alcohol	0	0	_	0
Methylene Chloride	3	0	0	3
Naphtha	0	0	1	0
Nitric Acid	0	0	0	0
Nitric Acid (conc.)	3	0	0	_
Nitrobenzene	_	0	_	**
Phenol	_	0	_	0
Pyridine	3	0	_	_
Silver Nitrate	_	0	_	0
Soap Solutions	_	0	_	0
Stearic Acid	_	0	_	*
Sulfuric Acid	0	0	0	0
Sulfuric Acid (conc.	) 3	0	0	_
Sulfurous Acid	_	0	0	0
Tannic Acid	_	0	0	0
Tannin Extracts	_	_	_	_
Tartaric Acid	_	0	_	_
Toluene	3	0	1	**
Trichloroethylene	3	0	3	3
Turpentine	2	0	0	**
Water	0	0	0	0
Xylene	3	0	0	*

<sup>\*</sup> Polypropylene – satisfactory to 22°C (72°F), \*\* Polypropylene – satisfactory to 49°C (120°F)

# 4.5 Changing the measurement direction

The measurement direction can be changed by moving the whole optical unit from above the measurement chamber to below the measurement chamber, or vice versa.

- 1. Switch off the instrument and disconnect the mains power supply cable (Fig. 3.3:2). There are anode voltages inside the mains power supply box and the optical unit if the power is on.
- 2. Remove the instrument cover as described in Section 3.2 Releasing the transportation lock.
- 3. Undo the four finger nuts (Fig. 4.5a:2) to release the light cover (Fig. 4.5a:1).



- 1 Light cover
- 2 Finger nuts, 4 pieces

Fig. 4.5a Removing the light cover of the optical unit

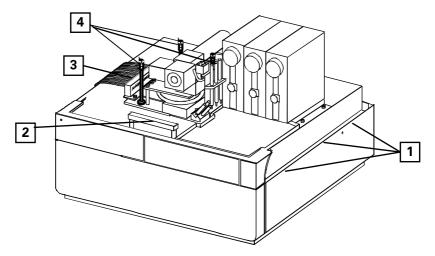
5. Undo the six screws, three screws on each side of the instrument (Fig. 4.5b:1), fixing the measurement chamber to the chassis.

6. Lift the measurement chamber into the upper position from the handle (Fig. 4.5b:2). There are hinges in the rear and a gas spring (Fig. 4.5c:3) holds the chamber in the upper position.



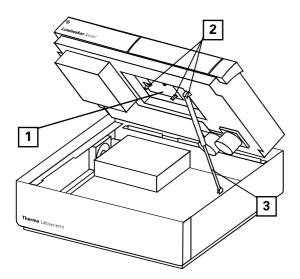
Caution: The measurement chamber is very heavy. DO NOT leave your fingers between the measurement chamber and the bottom case.

7. Remove the three short finger nuts holding the cover (Fig. 4.5c:2). Then remove the cover plate (Fig. 4.5c:1) from the opposite position of the optical unit under the measurement chamber.



- 1 Six screws, three screws on each side of the instrument
- 2 Handle
- 3 Flat cable
- 4 Long finger nuts, 3 pieces

Fig. 4.5b Removing the optical unit



- 1 Cover plate
- 2 Short finger nuts, 3 pieces
- 3 Gas spring

Fig. 4.5c Cover plate

- 8. Lower the measurement chamber back into the down position from the handle.
- 9. Unplug the flat cable (Fig. 4.5b:3) from the optical unit. DO NOT remove any other parts from the optical unit.
- 10. Undo the three long finger nuts (Fig. 4.5b:4) and remove the optical unit.
- 11. Lift the measurement chamber into the upper position from the handle.
- 12. Place the optical unit into the position where you removed the cover plate (Fig. 4.5c:1). Fix it with the long finger nuts and plug in the cable connectors and the flat cable. No adjustments are needed.
- 13. Lower the measurement chamber back into the down position.
- 14. Place the cover plate instead of the removed optical unit and fix it with the short finger nuts.
- 15. Fit the six screws back to hold the measurement chamber and replace the instrument cover. The unfitted cover may increase stray light and harm the measurement.
- 16. To change the measurement direction, the optical unit and the cover plate are interchangeable.

# 4.6 Installing or removing the light shield

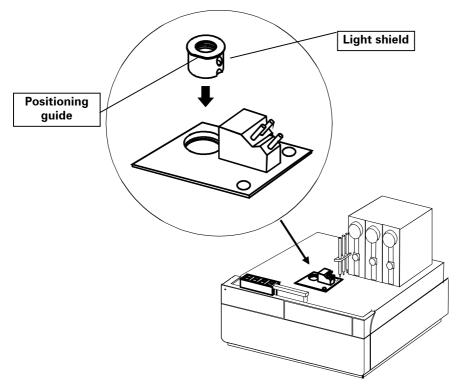


Fig. 4.6a Installing the light shield

- 1. Switch off the instrument and disconnect the mains power supply cable (Fig. 3.3:2).
- 2. Remove the instrument cover as described in Section 3.2 Releasing the transportation lock.
- 3. Undo the four finger nuts (Fig. 4.5a:2) and remove the light cover (Fig. 4.5a:1).
- 4. Undo the three long finger nuts (Fig. 4.5b:4) and remove the optical unit or, if the optical unit is below the measurement chamber, remove the cover plate (Fig. 4.5c:1).

5. When removing the light shield, use your little finger to lift the light shield up. When installing the light shield, ensure that the positioning guide is placed towards the front of the instrument against the corresponding guide in the holder (Fig. 4.6a and Fig. 4.6b). If the light shield is not positioned correctly, the optical unit does not slot down into its correct place but stays swinging.



Caution: Mispositioning of the light shield will cause discrepancies in the measurement results or may cause the instrument not to work.

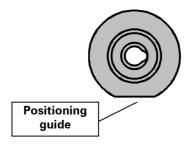


Fig. 4.6b Top view of the light shield

6. Fit the optical unit, light cover and the instrument cover back into their places.

# 4.7 Installing the filters

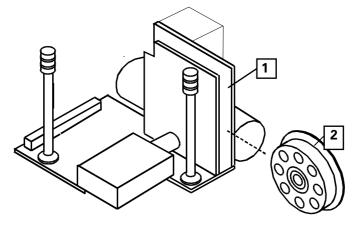
- 1. Switch off the instrument and disconnect the mains power supply cable (Fig. 3.3).
- 2. Remove the instrument covers as described in Section 3.2 Releasing the transportation lock points 1 4.
- 3. If the optical unit is above the measurement chamber, undo the four finger nuts (Fig. 4.5a:2) and remove the light cover (Fig. 4.5a:1). You can find more detailed instructions in Section 4.5 Changing the measurement direction.
- 4. If the optical unit is below the measurement chamber, undo the six screws (Fig. 4.5b:1) fixing the measurement chamber to the chassis.

Lift the measurement chamber into the upper position. There are hinges in the rear and a gas spring (Fig. 4.5c:3) holds the chamber in the upper position.



Caution: The measurement chamber is very heavy. DO NOT leave your fingers between the measurement chamber and the bottom case.

- 5. Unplug the flat cable (Fig. 4.5b:3) from the optical unit. Undo the three long finger nuts (Fig. 4.5b:4) and remove the optical unit and place it on a table.
- 6. The filter wheel (Fig. 4.7a:2) can be removed by undoing the fitting screw (Fig. 4.7a:1). DO NOT touch the filter surfaces.
- 7. Select the first free filter slot. DO NOT install any filters into the filter wheel positions 7 and 8.



- 1 Fitting screw(s) holding the filter wheel in place
- 2 Filter wheel

Fig. 4.7a Filter wheels in the optical unit

- 8. Undo the fitting screws (Fig. 4.7b:1).
- 9. Remove the spring wheel (Fig. 4.7b:2).
- 10. Remove the spacer collar (Fig. 4.7b:3).

- 11. Remove the dummy filter or previously installed filter used (Fig. 4.7b:4). Install a new filter so that the small arrow on the filter rim points away from the filter wheel. The arrow shows the direction of the light flow. DO NOT install any filters into the filter wheel positions 7 and 8.
- 12. Fit the spacer collar, spring wheel and the fitting screws. Fit the filter wheel back into place.

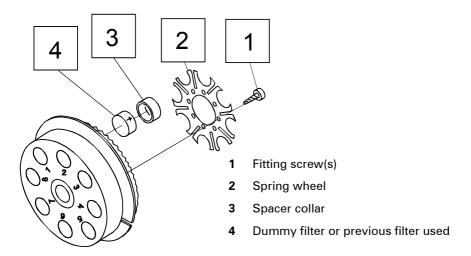


Fig. 4.7b Filters in a filter wheel

13. Replace the optical unit and connect the flat cable. Lower the measurement chamber and replace the instrument cover.

#### 4.8 Shutdown

- Remove any microplates left on the plate carrier or breakable strips from the tip priming vessel. Dispose of all microplates and strips as biohazardous waste.
- Flush the pump(s) out thoroughly with distilled water. Empty all the tubings.
- Place the dispensing tips into the tip holder (Fig. 2.3:5).
- Switch off the Luminoskan Ascent by pressing the power switch on the left-hand side of the instrument into the OFF position.
- Wipe the plate carrier surface and the external surfaces of the instrument with a soft cloth or tissue paper moistened with distilled water or a mild detergent solution.
- Push the plate carrier manually in.
- If you have spilt infectious agents on the plate carrier, disinfect with 70% pure ethanol in distilled water or some other disinfectant. See Section 8.2 Decontamination procedure.
- If you are not using the Luminoskan Ascent for an extended period of time, always clean the external surfaces of the instrument.
- Finally put the dust cover on.

### 5. Maintenance



**Note:** Follow normal laboratory safety procedures with regard to biohazardous, infectious, radiologic or toxic materials when maintaining the instrument.

# 5.1 Routine cleaning of the instrument

For reliable operation keep the instrument free of dust and spills of liquids. We recommend that you clean the case of the instrument periodically. A soft cloth dampened in mild detergent is sufficient. We recommend that you service the instrument at least yearly.

If you believe liquid has entered the luminometer, switch the instrument off and contact your local Thermo representative or Thermo Electron or for technical service (see Sections 8.1 Service request protocol and 8.4 Shipping the instrument (or items). If any surfaces have been contaminated with biohazardous material, a sterilizing solution must be used.

The prescribed decontamination procedure (see Section 8.2 Decontamination procedure), or similar routine, must be performed before returning the instrument to the supplier for service or repair. All instruments must be accompanied by a completed and signed Certificate of Decontamination securely attached to the exterior of the packaging (see Section 8.3 Certificate of Decontamination).

# 5.2 Cleaning the optical system

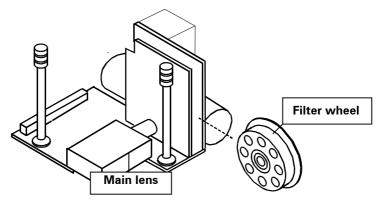


Fig. 5.2 Cleaning the optical unit

- Switch off the instrument and disconnect the mains power supply cable. Locate the optical unit and remove it according to the instructions in Sections 3.2 Releasing the transportation lock and 4.5 Changing the measurement direction.
- 2. Clean the main lens (Fig. 5.2), the possible filters and the light shield (Fig. 4.6a) with a cloth dampened with 96% pure ethanol and afterwards with a lint-free cloth or a lens tissue.



DO NOT use any other liquids to clean the optical unit. Avoid any harsh treatment.

3. Replace all the removed parts and reconnect the instrument to the mains power supply.

#### 5.2.1 Visual filter check

The useful life of a filter depends on environmental factors, such as dust, humidity and temperature. Filters have a one year warranty.

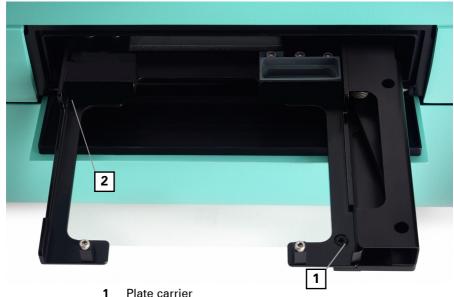
Carry out the visual check in the following way:

Visually check the filter(s) by holding it (them) against an even light source. If the color of the filter is even, then the filter is suitable for use. On the

other hand, if the filter appears to be mottled or discolored, discard the filter since it is either damaged or defective.

The best alternative is to measure the filters with a spectrophotometer.

#### Cleaning the plate carrier 5.3



- Plate carrier
- A1 corner of the microplate

Fig. 5.3 Plate carrier

Clean the plate carrier (Fig. 5.3) with a cloth dampened with distilled water. In case of any spills of infectious agents, clean the plate carrier with disposable towels dampened with a disinfectant solution containing 2% glutaraldehyde. Always use disposable gloves. Place the disposable towels and gloves in a biohazardous waste container.

# 5.4 Replacing the fuses

- 1. Switch off the instrument and disconnect the mains power supply cable (Fig. 5.4:1). Locate the fuse holders (Fig. 5.4:2) at the rear of the instrument.
- 2. Open the fuse holder to expose the fuse (Fig. 5.4:3) using a screwdriver.
- 3. Replace the faulty fuse with the spare provided with the accessories or with the same certified type.
- 4. Reconnect the instrument to the mains power supply. If the fuse blows again, contact your local Thermo representative for technical service.

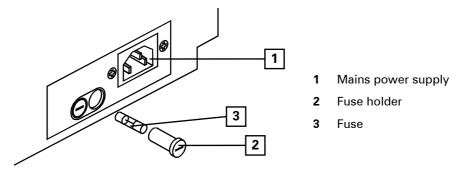


Fig. 5.4 Replacing the fuses

# 5.5 Routine maintenance of optional dispensers and main lens <sup>1</sup>

To obtain optimum performance and maximum useful life from the dispensers, it is important that the recommended cleaning maintenance instructions are followed.

Luminometry is a very sensitive detection technology. Therefore, take special care to avoid any contamination of any parts of the dispenser tubings and follow all GLP (Good Laboratory Practice) recommendations.

#### 5.5.1 Basic maintenance

- 1. The basic maintenance procedure should be performed regularly to ensure proper dispenser operation.
- Rinse the dispenser tubings out thoroughly with distilled water after each use.
- 3. DO NOT allow the dispensers to run dry for more than a few cycles.
- 4. Inspect the dispensers for leaks and rectify any problems immediately.
- 5. Wipe up all spills on and around the dispensers immediately.

#### 5.5.2 Extended maintenance

Clean the fluid path thoroughly using one of the procedures outlined below. There are three ways that the dispensers may be cleaned:

- Weak detergent
- 10% bleach
- · Weak acid and base

# 5.5.2.1 Weak detergent or 10% bleach

Remove the dispensing heads from the dispensing positions and DO NOT let any cleaning fluids enter the measurement chamber. Use external containers.

1. Prime the dispensers with a weak detergent or 10% bleach solution and leave it in the dispensers with the syringes full for 30 minutes.

<sup>&</sup>lt;sup>1</sup> Instructions concerning the pump are reproduced from CAVRO XP 3000 Modular Digital Pump Operators Manual made by Cavro Scientific Instruments, Inc., USA.

- 2. After the 30-minute period, remove the aspirate tubing from the detergent or bleach solution and remove all the fluid from the syringes and tubing into a waste container.
- 3. Flush the dispenser a minimum of 10 cycles with distilled water.

#### 5.5.2.2 Weak acid and base in sequence

Remove the dispensing heads from the dispensing positions and DO NOT let any cleaning fluids enter the measurement chamber. Use external containers.

1. Prime the dispensers with 0.1 M NaOH and leave the solution in the dispensers for 10 minutes with the syringes full.



**Note**: DO NOT spill any 0.1 M NaOH onto any instrument surfaces to avoid damage of the instrument. If needed, use suitable protection covering.

- 2. Flush the dispensers with distilled water.
- 3. Prime the dispensers with 0.1 M HCl and leave the solution in the dispensers for 10 minutes with the syringes full.
- 4. After the 10-minute period, remove the aspirate tubing from the 0.1 M HCl solution and remove all the fluid from the syringes and tubing into a waste container.
- 5. Flush the dispensers a minimum of 10 cycles with distilled water.

# 5.5.2.3 Cleaning the main lens

It is recommended that you check the main lens of the optical unit weekly.

- 1. Remove the optical unit if it is above the measurement chamber, or the cover plate if the optical unit is below the measurement chamber, as described in Section 4.5 Changing the measurement direction.
- 2. If needed, clean the main lens according to the instructions in Sections 5.2 Cleaning the optical system.

### 5.6 Periodic maintenance

There are three parts which require periodic maintenance: tubing; syringe seals, and valves. If they become worn out, the symptoms are:

- Poor precision and accuracy
- A variable or moving air gap
- Leakage
- Drops and spills

The frequency of replacement will depend on the duty cycle, fluids used and instrument maintenance.

If any of these symptoms occur and it is not obvious which component is causing the problem, the easiest and most economical way is to replace one component at a time in the following order: (1) dispensing or aspirate tubings and/or dispensing tip, and (2) syringe.

#### 5.6.1 Replacing the dispenser tubings

- 1. To remove either the dispensing tube or the aspirate tube assembly from the valve, gently loosen the fittings either manually or using a 7.9 mm (5/16 in.) wrench. Unscrew the fittings and remove the tubing.
- We recommend you replace the complete assemblies always when replacement is necessary. Alternatively, only some parts of the assemblies can also be replaced. If only the dispensing tubing is changed, first remove the complete dispensing tube assembly from the dispenser unit.

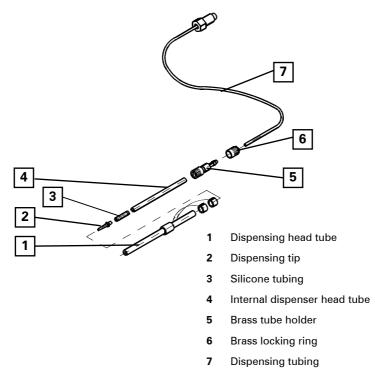
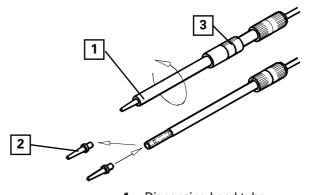


Fig. 5.6.1 Structure of the complete dispensing tube assembly

- 3. Remove the dispensing head tube (Fig. 5.6.1:1) from the brass tube holder (Fig. 5.6.1:5).
- 4. Loosen the brass locking ring (Fig. 5.6.1:6).
- 5. Remove the dispensing tip (Fig. 5.6.1:2) and the connecting piece of silicone tubing (Fig. 5.6.1:3).
- 6. Remove the internal dispenser head tube (Fig. 5.6.1:4), the brass tube holder and the brass locking ring.
- 7. Insert the new dispensing tubing into the brass tube holder/locking ring and the internal dispenser head tube.
- 8. Connect the dispensing tip with the connecting piece of silicone tubing into the new dispensing tubing.

- 9. Fit the dispensing tip and tubing into the dispenser head tube. When you push the dispensing tubing gently towards the tip, the internal dispenser head tube should be visible about 1 mm. This can be adjusted through the length of the connecting piece of silicone tubing.
- First screw the brass tube holder onto the dispenser head tube.
   Gently tighten the tubing and ensure that the tube end stays in the silicone tubing. Then tighten the brass locking ring.
- 11. To fit a new tubing, insert the fitting into the valve and tighten it finger-tight. Using a 7.9 mm (5/16 in.) wrench, turn the fitting another quarter to half turn.

#### 5.6.2 Replacing the dispensing tip

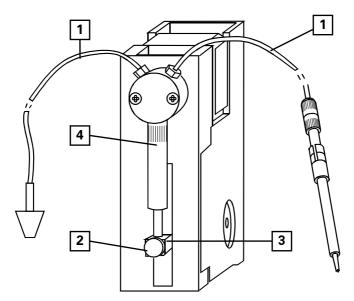


- 1 Dispensing head tube
- 2 Dispensing tip
- 3 Brass tube holder

Fig. 5.6.2 Replacing the dispensing tip

- 1. Remove the dispensing head tube (Fig. 5.6.2:1) from the brass tube holder (Fig. 5.6.2:3).
- 2. Replace the dispensing tip (Fig. 5.6.2:2) connected with a small piece of silicone tubing in the dispensing tube.
- 3. Replace the dispensing head tube.

#### 5.6.3 Replacing the dispenser syringe



- 1 Aspirate and complete dispensing tube assemblies
- 2 Plunger lock screw
- 3 Plunger holder arm
- 4 Dispenser syringe (1.0 ml) and plunger

Fig. 5.6.3 Replacing the dispenser syringe

- 1. Remove the liquid from the dispenser syringe (Fig. 5.6.3:4) and from the tubings.
- 2. Switch off the power from the instrument.
- 3. Push the plunger manually into the upper position.
- Loosen the plunger lock screw (Fig. 5.6.3:2) approximately three full turns.
- 5. Pull the plunger holder arm (Fig. 5.6.3:3) firmly down.
- 6. Unscrew the syringe from the valve.
- 7. To fit the new dispenser syringe, screw the syringe into the valve, pull the syringe plunger down to the plunger holder arm and screw it into place. Make sure the plunger lock screw is securely tightened.

# 6. Troubleshooting Guide



**Note:** DO NOT use the instrument if it appears that it does not function properly.

# 6.1 Troubleshooting

Pro )lem	Cai se	Actic 1
Instrument does not turn on correctly	No power connection	Check the power cable and the fuses
No connection between instrument and PC	Incorrect software type	Check that the software is installed for the correct instrument model
	Loose serial cable connections	Check the cable connections
	Software in simulation mode	Switch the connection to the instrument
Too high background	No liquid in the well	Always use some liquid in the blank wells
	Unclean plate	Use a disposable plate only once
	Contaminated reagents	Replace the reagents used
	Microplate material	Check if the plate manufacturer or material has changed
	Contaminated tubing	Replace/clean the tubing
	Phosphorescence from the plastic	Use only plastics designed for luminometry without phosphorescence
Too low/high signal	Changed scaling factor	Check the scaling factor used
	Microplate material	Check if the plate manufacturer or material has changed <i>Continued</i>

Pro vlem	Ca: se	Actic 1
Even, low signal level for the whole plate	Bottom reading of non- transparent plate	Use top reading
	Bottom reading when drop plate in its place	Remove the drop plate
Too large deviation between replicates	Unclean plate	Use a disposable plate only once
	Dust or dirt in the wells	Keep the plates protected before use
	Foaming in the sample	Use a lower dispensing or shaking speed
		Use correct pipetting techniques (reverse)
Illogical results	Plate inserted with incorrect orientation	Rotate the plate with the A1 well facing towards the upper left corner
Too low or variable	Inadequate priming	Reprime the dispenser
volume from dispenser		Use tip priming during the session
	Incorrect dispensing height	Check that the dispenser heads are positioned at the correct height
	Incorrect dispensing height	Check that the dispenser head positions in the instrument and in the software settings are identical
		Use plate adapters with low 384-well plates
	Loose syringe, plunger or tubings	Check all the connections in the dispenser
	Aspiration tube not in the liquid	Ensure that there is a sufficient volume of
		Continued
		Ensure that the aspiration

Pro vlem	Cai se	Action
		tube end weight is at the bottom of the vial and attach it firmly, if necessary
Liquid droplets outside the wells after	Incorrect plate template used	Check the plate type used in the software
dispensing	Too high total volume used	Use smaller volumes
	Lid used with the plate	DO NOT use plate lids with the dispensers
	Damaged dispensing tip	Replace the tip
	Incorrect installation of the dispensing head	Check that the dispensing head is inserted deep enough
	Incorrectly adjusted dispenser head assembly	Contact authorized technical service
Noise from the instrument	Light shield not removed when using high plates	Remove the light shield
	Lid left on the plate	Remove the light shield or the lid
	Dispenser head installed to the incorrect height	Check the dispenser/plate height
	Filter wheel too tight	Loosen the filter wheel screw
	Transportation lock not removed	Remove the transportation lock
	Plate adapter for low 384-well plates left under a 96-well plate	Remove the adapter when using 96-well plates
	aasi a oo iron piato	Continued

Pro >lem	Cai se	Actic
Plate carrier moves incorrectly	Foreign object, e.g., a tip priming strip, obstructs the plate carrier movement	Ensure that there are no loose objects inside the instrument
"Too high background drift" message	Dirty measurement window	Clean the measurement window
Accidental dispensing into the instrument		Contact authorized technical service

# 6.2 Error messages

Error messages issued by the instrument are listed in the Reference part of the Ascent Software User's Guide in the chapter on warning and error messages.

# 7. Frequently asked questions (FAQ) about the Luminoskan Ascent

# Q: Can Luminoskan Ascent be upgraded for fluorometric applications?

**A**: Luminoskan Ascent is a dedicated luminometer with no possibility to add the fluorometric option. It is not possible to add the luminometric option to the Fluoroskan Ascent. Only the Fluoroskan Ascent FL can measure both fluorescence and luminescence.

# Q: Are Petri dish adapters standard accessories that come with the Luminoskan Ascent?

**A**: Petri dish adapters are not standard accessories that come with the reader and thus they should be ordered separately.

# Q: When is the adapter for low 384-well plates needed?

A: If the plate height is less than 13.5 mm, an adapter must be used to raise the plate. The adapter for low 384-well plates is needed when you <u>dispense</u> and/or <u>measure</u> into low 384-well plates.

# Q: How is the scaling factor in Ascent Software used?

**A**: The scaling factor is a constant that multiplies all the readings from the instrument. It is defined separately for each filter pair and beam size (fluorometry) or filter position (luminometry). The scaling factor can be used to convert the instrument reading to a desired level or to adjust several instruments to give similar signal values.

# Q: What plate colors can be used for luminescence?

A: White plates are generally used for luminometric measurements to enhance the signal and to avoid possible crosstalk from high signals in neighboring wells.

# Q: Can dispensing be performed into 864-well plates?

A: No, dispensing into 864-well plates is not possible.

# Q: When is the light shield needed with the Luminoskan Ascent?

A: The light shield should be used with 96- and 384-well plates. If the plates are higher than 15 mm, the light shield must be removed.

- Q: How many filters can be installed into the Luminoskan Ascent?
- A: A maximum of six filters into the filter wheel positions 1 6.
- Q: What plate types can be measured with the Luminoskan Ascent?

**A**: The Luminoskan Ascent can measure plate formats from 1 to 864. Customized plates with maximum dimensions of 90 mm x 143 mm x 25 mm can be used.

# Q: Is it possible to measure 96-well PCR plates with the Luminoskan Ascent?

A: Yes, and you can use the Luminoskan Ascent for measuring both PCR plates and 0.2 ml PCR tubes but you need a special adapter. This adapter is suitable, for example, for Hybaid Omniplates, Robbins Cycloplates and Axygen PCR microplates or tubes from Greiner or Sarstedt.

# Q: What kind of plates are best to use with the Luminoskan Ascent when you have to reduce the crosstalk as much as possible and minimize the background?

A: There are clearly two options. You should use either white Microlite 1+ plates (Cat. no. 7571) or white universal binding plates (Cat. no. 9502887).

# Q: What kind of plate adapters are available for the Luminoskan Ascent?

**A**: The adapters listed below are available. Other adapters are available on request.

5210310	Adapter for Petri dish 2 x ø 59 mm
5210300	Adapter for Costar Petri dish ø 95 mm
5210380	Adapter for Falcon/Greiner Petri dish ø 93 mm
5210330	Adapter for Petri dish 6 x ø 40 mm
5210320	Adapter for low 384-well plates
5210390	Robotic 384-adapter
5210340	Adapter for Terasaki plates
5210630	Adapter for PCR tubes and 96-well PCR plates

### 8. Instrument Service

# 8.1 Service request protocol

If the Luminoskan Ascent requires service, contact your local Thermo representative or Thermo's service department. DO NOT under any circumstances send the instrument for service without any prior contact. It is imperative to indicate the fault and nature of the required service. This will ensure a faster return of the instrument to the customer.

The Thermo representative or distributor takes care of sending Thermo's service department a Feedback Form (Complaint-Order), which contains a more detailed description of the fault, symptom or condition. Give all the necessary information to the distributor, who will fill in and forward the Feedback Form to Thermo's service department.

Check Section 8.4 Shipping the instrument (or items). You will find instructions on how to proceed before shipping the instrument for service.

Check that any necessary decontamination procedure has been carried out before packing. See Sections 8.2 Decontamination procedure and 8.3 Certificate on Decontamination. Ensure that the Certificate of Decontamination is sent with the instrument.

The technical service department will keep you up to date with the progress of service and provide you with any further details you might need, for example, on maintenance, serviceability, troubleshooting and replacement.

# 8.2 Decontamination procedure

Decontamination should be performed in accordance with normal laboratory procedures. Any decontamination instructions provided with the reagents used should be followed.

A decontamination procedure is only recommended when infectious substances have been in direct contact with any part(s) of the instrument.

If there is any risk of contamination with biohazardous material, the procedure recommended below or some other corresponding decontamination procedure must be performed.

We strongly recommend that the complete decontamination procedure is performed before relocating the instrument from one laboratory to another.

#### **Example of disinfectants**

- Formaldehyde solution 10%
- Pure ethanol 70% (in distilled water)
- Virkon solution 1 3%
- Glutaraldehyde solution 4%
- Chloramine T



Caution: Always use disposable gloves and protective clothing and operate in a well-ventilated area.



Caution: DO NOT use denatured ethanol.

- Prepare the disinfectant: 200 ml 10% formaldehyde solution or 200 ml 4% glutaraldehyde solution (or another agent recommended by your safety officer).
- 2. Empty the plate carrier.
- 3. Switch off the power and disconnect the mains power supply cable and the computer cable.
- 4. Disinfect the outside of the instrument using a cloth dampened with 70% pure ethanol in distilled water.
- 5. Place the instrument in a large plastic bag. Ensure that all the lids are open.
- 6. Place a cloth soaked in the prepared solution into the bag. Ensure that the cloth does not make contact with the instrument.
- 7. Close the bag firmly and leave the instrument in the bag for at least 24 hours.
- 8. Remove the instrument from the bag.
- 9. Clean the instrument using a mild detergent.

- 10. Remove any stains using 70% pure ethanol in distilled water.
- 11. After performing this decontamination procedure, label the instrument with a signed and dated Certificate of Decontamination.

#### 8.3 Certificate of decontamination

The decontamination procedure is required prior to shipping the instrument to Thermo Electron Oy, for example, for repair. If, for any reason, the instrument is shipped back to Thermo Electron Oy, it must be accompanied by a dated and signed Certificate of Decontamination which must be attached to the outside of the package containing the instrument. See Section 8.2 Decontamination procedure.

Failure to confirm decontamination will incur additional labor charges or at worst the items will be returned for proper cleaning.

Before returning any instrument(s) or items, ensure that they are fully decontaminated. Confirm A or B status (see below).

## Certificate of Decontamination

Name:	
Address:	
Tel./Fax:	
Instrument:	Serial no.:
	eturned items have not been contaminated by body fluids adioactive materials or any other hazardous materials.
	urned items have been decontaminated and can be handled ersonnel to health hazards.
Materials used in the ur	nit: Chemicals + Biological • Radioactive
Specific information about contaminants:	
Decontamination procedure:	
Date and place:	
Signature:	
Name (block capitals):	
*) The signature of a f been used with radioac	Radiation Safety Officer is also required when the unit has tive materials.
This unit is certified by	the undersigned to be free of radioactive contamination.
Date and place:	
Signature:	
Name (block capitals):	

### 8.4 Shipping the instrument (or items)

When you ship the instrument for service remember to:

- 1. Empty the dispenser(s) and remove the tube assembly.
- 2. Remove any loose items from the plate carrier, for example, adapters, plates and priming vessels.
- 3. Remove the power cable as well as the serial cable.
- 4. Decontaminate the instrument beforehand.
- 5. Fit the transportation lock.
- 6. Pack the instrument according to the enclosed packing instructions.
- Use the original packaging to ensure that no damage will occur to the instrument during shipping. Any damage will incur additional labor charges.
- 8. Inform about the use of hazardous materials.
- Enclose a dated and signed Certificate of Decontamination (see Section 8.3 Certificate of Decontamination) both inside and attached to the outside of the box in which you will return your instrument or other items.
- 10. Indicate the fault after you have been in touch with your local Thermo representative or Thermo's service department.
- 11. Enclose the return authorization number (RGA) given by the Thermo representative.

See Section 14.1 General specifications for details on storage and transportation temperatures.

### 8.5 Service contracts

We strongly recommend that you maintain and service the instrument every twelve months on a contract basis by the manufacturer's trained service engineers. This will ensure that the product is properly maintained and gives trouble-free service. Contact authorized technical service for further details.

## 9. Disposal of the Instrument and Materials

### 9.1 Disposal of the instrument

For the disposal of the instrument do the following:



- Decontaminate the instrument prior to disposal. See Sections 8.2 Decontamination procedure and 8.3 Certificate of Decontamination.
- Dispose of the instrument according to the legislation stipulated by the local authorities concerning take-back of electronic equipment and waste. The proposals for the procedures vary by country.
- Regarding the original packaging and packing materials, use the recycling operators known to you.
- For further information, contact your local Thermo representative.

### 9.2 Disposal of materials

Refer to local regulations for the disposal of infectious material.



The samples can be potentially infectious. Dispose of all used microplates, PCR tubes, disposable gloves, syringes, disposable tips, etc., as biohazardous waste.

# 10. Ordering Information

### 10.1 Product code numbers

## **Microplate Luminometers**

Item	Cat. no.
Luminoskan Ascent	5300160
Luminoskan Ascent with dispenser	5300170
Additional dispenser option (factory installation)	5210230

#### **Accessories**

ltem .	Cat. no.
1st Dispenser kit (user installation)	2805621
Additional dispenser kit (user installation)	2805630
Plate carrier kit	2806140
Adapter for Costar Petri dish Ø 95 mm	5210300
Adapter for Falcon/Greiner Petri dish Ø 93 mm	5210380
Adapter for Petri dish 2 x Ø 59 mm	5210310
Adapter for dispensing into a low 384-well plate	5210320
Adapter for Petri dish 6 x Ø 40 mm	5210330
Adapter for Terasaki plate	5210340
Drop plate	2805880
Light shield	1006470
Dispensing tip	1047661
Complete dispensing tube assembly (Fig. 3.4.2 and Fig. 5.6.1)	24071580
Aspirate tube	24071510
Dispensing tubing (Fig. 5.6.1:7)	24071500
Dispensing syringe (1.0 ml) and plunger (Fig. 3.4:4)	24071490
Syringe seal	2805600
3-port valve	2805610
Leakage tray (Fig. 2.3:4)	10478030
End weight	1002380
Red adjustment collar (Fig. 4.4.4b:2)	1002820
Silicone tube (for connecting pieces)	0412290
RS-232C interface cable D9 Female/D25 Female	2305290
RS-232C interface cable D25 Female/D25 Female	2303760
Fuse 3.5 A/250 V UL 198G Time Delay (10 pcs min. order)	1210950
Dust cover	1610460
Luminoskan Ascent User Manual	1507520
Ascent Software User's Guide	1507530CD
Thermo Electron Ascent Software	5185450CD

## **Thermo Electron Microtiter plates**

Cat. no.	Description	Bottom	Qty
Thermo Micr	otiter Plates for Luminescence		
7416	Microlite 1 Plate	Flat	50/box
7417	Microlite 2 Plate	Flat	50/box
7418	Microlite TCT Plate, sterile	Flat	50/box
7571	Microlite 1+ Plate	Flat	50/box
7572	Microlite 2+ Plate	Flat	50/box
7521	Microlite 1+ Plate	U	50/box
7522	Microlite 2+ Plate	U	50/box
9502887	White 96 Well Plate UB	Flat	50/box
95029580	White 96 Well Plate EB	Flat	50/box
95029770	White 96 Well Plate Sterile	Flat	40/box
Cat. no.	Description	Bottom	Qty
	otiter Plates for Luminescence, Strips and As		Qty
7421	Microlite 1 1x12 Strip assembled	Flat	100/box
7403	Microlite 1 1x12 Strip	Flat	320/box
7400	Microlite 2 1x12 Strip assembled	Flat	100/box
7410	Microlite 2 1x12 Strip	Flat	320/box
7561	Microlite 1+ 1x12 Strip assembled	Flat	100/box
7566	Microlite 1+ 1x12 Strip	Flat	320/box
7562	Microlite 2+ 1x12 Strip assembled	Flat	100/box
7567	Microlite 2+ 1x12 Strip	Flat	320/box
95029510	White Strip 1x8 assembled UB	Flat	50/box
95029530	White Strip 1x12 assembled UB	Flat	50/box
95029660	White Breakable Strip 1x8 Assembled UB	Flat	50/box
Cat. no.	Description		Qty
	otiter 384 Well Plates		Qty
95040010	White 384 Round Well Plate		50/box
95040010	White 384 Round Well Plate Sterile with lid		40/box
33040230	write 304 noutly well riate Sterile With IIa		HU/DUX

### Thermo Electron Microtiter plates Cont.

Cat. no.	Description	Bottom	Qty
Thermo Mic	rotiter Streptavidin-Coated Plates		
95029273	White BioBind Strip 1x8 Assembled	Flat	5/box
95029303	White BioBind Br. Strip 1x8 Assembled	Flat	5/box
Cat. no.	Description		Qty
Thermo Mic	rotiter Plate Accessories		
5500 <sup>1)</sup>	Universal Polystyrene Lid, Sterile		100/box
5550 <sup>1)</sup>	Styrene Individually Wrapped Lid, Sterile		50/box
9503210 <sup>2)</sup>	Microplate Lid, 96-well		15/box
9503220 <sup>2)</sup>	Microplate Lid		15/box
6305	Vinyl Lid for 1x12 strip assembly		100/box
6604 <sup>1)</sup>	Holder for 1x12 strip assembly		10/box
6000 <sup>1)</sup>	Workstation 1x12 strip assembly		1/box

<sup>1)</sup> Accessories for ex-Dynex products

<sup>&</sup>lt;sup>2)</sup> Accessories for ex-Thermo products

## 10.2 List of recommended spare parts

Cat. no.	<i>ltem</i>	1–2 unit(s)/year	10 units/year
24071580	Complete dispensing tube	1	5
24071500	assembly Dispensing tubing	2	8
1047661	Dispensing tip	4	20
2805690	Aspirate tube assembly	1	5
24071490	Dispenser syringe (1.0 ml) and plunger	1	5
1210950	Fuse 3.5 A/250 V UL 198G Time Delay	1	5
1006470	Light shield	1	3
1002820	Red adjustment collar	2	10

## 10.3 Ordering filters

When ordering luminometric filters, please contact Thermo Electron Oy. E-mail address: <a href="mailto:info.microplateinstruments@thermo.com">info.microplateinstruments@thermo.com</a>.

## 11. Glossary and Abbreviations

ATP Adenosine triphosphate, a biological molecule

that is commonly used as a reference chemical

for luminometric sensitivity.

**Chassis** The framework of the instrument.

Chemiluminescence A compound that emits light following a

chemical reaction is said to be

chemiluminescent.

Crosstalk Interfering signal from neighboring wells.

**Decade** Order of magnitude. A logarithmic value that is

used for presentation of dynamic range.

**Decontamination** Removal or neutralization of radiologic,

bacteriological, chemical or other

contamination.

**Disinfection** The destruction of pathogenic bacteria, usually

with an antiseptic chemical or disinfectant.

**Dynamic range** Dynamic range refers to the range of signals an

instrument can read, from the minimum to the maximum detectable. For example, dynamic range of seven decades means that the difference between the lowest and highest

signals that can be measured is 107.

**Error message** Indication that an error has been detected.

**Initialization** Initialization tests are so-called self-tests, which

are carried out prior to operation to ascertain that the necessary instrument adjustments have

been carried out.

**LED** Light-emitting diode.

Luminescence Emission of light (other than from thermal

energy causes) such as bioluminescence.

Luminometer An instrument used for measuring the intensity

of luminescent radiation.

**Luminometric label** 

(Luminophore)

A substance which emits light at room temperature. A group of atoms that can make a

compound luminescent.

Photomultiplier tube

(PMT)

A photoelectric cell that converts light into

electric current and amplifies the current.

RH Relative humidity.

**RLU** or rlu Relative Luminescence/Luminometric/Light

Units. The arbitrary units in which luminescence

intensity is reported.

Revolutions per minute. rpm

Scaling The measured values are expressed as RLU.

Scaling is a way to convert readings to show

desired values.

**Self-tests** Initialization tests and adjustments that the

instrument performs prior to operation as well

as autocalibration.

#### 12. Literature

Fluorescent and Luminescent Probes for Biological Activity. A Practical Guide to Technology for Quantitative Real-Time Analysis (1999). Mason W. T., (ed.). Second edition, Biological Techniques Series, Academic Press.

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Chemiluminescence: Principles and Applications in Biology and Medicine (1988). Campbell, A. K.. Horwood, Chichester, U.K.; 608 pages.

**Nonradioactive labeling and detection of biomolecules** (1992). Kessler C. (ed.). Springer-Verlag, Berlin; 436 pages.

**Bioluminescence Methods and Protocols** (1998). LaRossa R. A. (ed.). Methods in Molecular Biology, Vol. 102. Humana Press., Totowa, New Jersey; 310 pages.

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**Bioluminescence & Chemiluminescence** (2001). Case J. F., Herring P. J., Robison B. H., Haddock S. H. D., Kricka L. J., Stanley P. E. (eds.). World Scientific Publishing Co. Pte. Ltd., Singapore; 517 pages.

## 13. Warranty Certificate

Thermo Electron Microplate Instrumentation Business products are fully guaranteed against defective parts and materials, including defects caused by poor workmanship, for a period of one year from the date of delivery.

Thermo will repair or replace defective parts or materials during the term of warranty at no extra charge for materials and labor provided that the products were used and maintained in accordance with Thermo's instructions. The warranty is invalid if products have been misused or abused.

For the warranty to be effective, the product must have been purchased either directly from Thermo or from an authorized Thermo distributor. The guarantee is not transferable to a third party without prior written approval from Thermo.

This guarantee is subject to the following exclusions:

- Any defects caused by normal wear and tear.
- Defects caused by fire, lightning, flood, earthquake, explosion, sabotage, war, riot, or any other occurrence of the type listed above.
- Refurbished products that are subject to different warranty conditions.

THIS WARRANTY IS IN LIEU OF ALL OTHER EXPRESSED OR IMPLIED WARRANTIES, INCLUDING BUT NOT LIMITED TO ANY IMPLIED WARRANTIES OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. The seller is not liable for any loss or damage arising out of or in connection with the use of the product or other indirect damages.

These warranty terms and conditions can be obtained from your local Thermo dealer.

This card acts as a warranty certificate.

## 13.1 Warranty limitations

The following items are not included in the warranty:

- Consumables
- Software programs
- Fuses.

## 14. Specifications

Thermo Electron reserves the right to change any specifications without prior notice as part of our continuous product development program.

## 14.1 General specifications

Weigh	Basic unit 21 kg (46 lbs.).
	3 optional dispensers add 3.5 kg to the weight.
Overal dimensions	420 mm (16.5 in.) (W) x 420 mm (16.5 in.) (D) x
	340 mm (13.4 in.) (H), options included.
Operating conditions	+10°C - +40°C, RH 90% max.
(indoo use)	Tested according to IEC 60068-2-1 test Ab, (Cold).
(	IEC 60068-2-2 test Bb, (Dry heat).
	IEC 60068-2-3 test Ca, (Damp heat).
Transportation conditions	-40°C – +70°C, packed in transportation packaging.
	Tested according to IEC 60068-2-1 test Ab, (Cold).
	IEC 60068-2-2 test Bb, (Dry heat).
Storag : conditions	-25°C – +50°C, packed in transportation packaging.
Storag / conditions	Tested according to IEC 60068-2-1 test Ab, (Cold).
	IEC 60068-2-2 test Bb, (Dry heat).
	120 00000 2 2 toot BB, (BTy Hout).
Mains ∋ower supply	100 – 240 Vac, 50/60 Hz, nominal
Fuses	2 x 3.5 A/250 V,
	UL 198G Time Delay, 5 x 20 mm.
	·
Power consumption	200 VA max., 32 VA standby.
Compi ter interface	Serial RS-232C port. Baud rate 9600. Character
	format 1 start bit, 8 data bits, 1 stop bit, no parity.
	Flow control XON/XOFF.

Continued

## General specifications Cont.

Detect <sub>'</sub> r	Photomultiplier tube.
Filters or luminometric measu ements	Filters can be used; maximum 6 filters.
Plate t pes	1- to 384-well plates. Can also be programmed for nonstandard configurations. Maximum dimensions 90 mm x 134 mm x 25 mm.
Shakeı	Built-in orbital shaker with adjustable speed and diameter.
Incuba or	Temperature range from RT (25°C) +3°C to 45°C, when the ambient temperature is 25°C. The temperature is selected using Ascent Software.
Dispen iers	$1-3$ optional dispensers. Syringe volumes $1000~\mu$ l. Dead volume $600~\mu$ l. Metal-free fluid path. Exchangeable valve, syringe, tubing and tip. Autoclavable tubing and tip.

### 14.2 Safety specifications

The Luminoskan Ascent fulfills the following requirements:

EN 61010-1:1993 + A2:1995/IEC 61010-1:1990 + A1:1992 + A2:1995 Installation Category (Overvoltage Category) II; Pollution Degree 2 EN 61010-2-010:1994 + A2:1996/IEC 61010-2-010:1992 + A1:1996 CSA C22.2 No. 1010.1 M1992

The safety specifications are also met under the following environmental conditions in addition to or in excess of those stated in the operating conditions:

Altitude up to 2000 m Temperature  $+5^{\circ}\text{C} - +40^{\circ}\text{C}$ 

**Mains supply**  $\pm$  10% (if larger than specified above)

fluctuations

Installation category II according to IEC 60664-1 (see Note 1)

(overvoltage category)

Pollution degree 2 according to IEC 60664-1 (see Note 2)

#### **Notes**

- 1) The *installation category* (overvoltage category) defines the level of transient overvoltage which the instrument is designed to withstand safely. It depends on the nature of the electricity supply and its overvoltage protection means. For example, in CAT II which is the category used for instruments in installations supplied from a supply comparable to public mains, such as hospital and research laboratories and most industrial laboratories, the expected transient overvoltage is 2500 V for a 230 V supply and 1500 V for a 120 V supply.
- 2) The *pollution degree* describes the amount of conductive pollution present in the operating environment. Pollution degree 2 assumes that normally only nonconductive pollution, such as dust, occurs with the exception of occasional conductivity caused by condensation.

Both of these affect the size of the electrical dimensioning within the instrument.

### 14.3 In conformity with the requirements

#### The Luminoskan Ascent bears the following markings:

Type 392 100 – 240 Vac 50/60 Hz, 200 VA CE mark CSA monogram UL monogram

#### The Luminoskan Ascent conforms to the following requirements:

73/23/EEC (Low Voltage Directive) 89/336/EEC (Electromagnetic Compatibility Directive, EMC) FCC Part 15, Subpart B/Class B

#### Safety performance:

EN 61010-1:1993 + A2:1995/IEC 61010-1:1990 + A1:1992 + A2:1995

#### **EMC** performance:

EN 50081-1:1992 Generic emission standard. Residential,

commercial and light industry.

EN 50082-1:1997 Generic immunity standard. Residential,

Class A

commercial and light industry.

EN 61326-1:1997 + A1:1998 Product family standard.

#### Test standards EN 55022:1998

Performance limits
Class B. 150 kHz – 1 GHz

EN 61000-3-2:1995 + A1:1998

+ A2:1998 + A13:1997 +

A14:2000

EN 61000-3-3:1995

ANSI C63.4:1992 Class B, 450 kHz – 1 GHz; 30 MHz – 1000 MHz

EN 61000-4-2:1995 + A1:1998 EN 61000-4-3:1996 + A1:1998 4 kV CD, 8 kV AD, Criteria B 3 V/m, 80 MHz – 2 GHz, Criteria A

EN 61000-4-4:1995

1 kV, Criteria B

EN 61000-4-5:1995

2 kV line to ground, 1 kV line to line, Criteria B

EN 61000-4-6:1996 3 V<sub>rme</sub>, 150 kHz – 80 MHz, Criteria A

EN 61000-4-8:1994 3 A/m, Criteria A

EN 61000-4-11:1994 30%/10 ms, Criteria B

60%/100 ms, Criteria C > 95%/5 s, Criteria C 100%/20 ms, Criteria B

# 14.4 Performance specifications

Warm- ıp time	< 15 min to rated accuracy.
Measu ing speed	Depends on the plate type and the measurement type. The minimum kinetic interval time is 15 s for a 96-well plate (from well A1 back to the same well A1).
Lumin metric spectral range	270 – 670 nm
Lumin metric measu ement range	Up to approx. 5000 Relative Light Units (RLU)
Lumin metric sensitivity	1 fmol ATP/well (typical) in a white Thermo Electron 96-well strip plate (Thermo Electron ATP monitoring kit). Limit = $(2 \times SD_{Blank}) / I_{Ref-Blank} \times C_{Ref}$
Lumin metric dynamic range	> 9 decades over whole gain setting area
Shakeı	Orbital method, speed 60 – 1200 rpm, Ø 1 – 50 mm
Incuba or	Heaters: Warm-up time from RT (25°C) to 37°C, 15 min. Liquid in the well: Ambient temperature 23°C, covered 96-well plate, 200 µl/well. Temperature accuracy: Mean temperature of the wells ± 1°C Uniformity: ≤ 1°C
	Warm-up speed 1 h from 23°C to 37°C on an average 90% of the set value and the ambient temperature.
Dispen ;ers	Dispensing volume 1 – 1000 $\mu$ l in 1 $\mu$ l increments. Accuracy $\pm$ 3 $\mu$ l avg. for volumes 5 $\mu$ l and above Precision 5 – 19 $\mu$ l < 5% 20 – 1000 $\mu$ l < 2% Minimum dispensing speed 25 s, 96-well plate, 5 $\mu$ l/well, tested with distilled water.

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## Appendix A. Brief User's Guide

- Unpack the instrument (p. 19).
- Check delivery for completeness (p. 20).
- Check for damage during transport (p. 20).
- Place the instrument according to environmental requirements (p. 21).
- Release the transportation lock (p. 22).
- Install the dispenser tubings if equipped with dispensers (p. 25).
- Plug in the instrument (p. 23).
- Switch the instrument ON and ensure that the green light is lit (p. 31).
- Install Ascent Software (p. 31).
- Create an Ascent Software session (see Ascent Software User's Guide: Procedure Desktop | Session | New).
- We recommend you perform a dummy run first with an empty plate when you have created a new session and then a run with known samples or controls.
- Load the microplate with prepared samples, blanks and controls.
   Place the microplate onto the plate carrier of the instrument so that the A1 well is located in the upper left corner of the plate carrier (p. 32).
- Start the session by pressing the START button on the Procedure Desktop toolbar of Ascent Software.
- Follow the instructions provided by the software.
- Switch the instrument OFF after routine operation (p. 53).
- Maintain the Luminoskan Ascent instrument on a regular basis (p. 55).

## **Appendix B. Addresses**

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