

SPECTRA 8P



USER MANUAL

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Ver.: 1.00

1. DESCRIPTION

1.1 TECHNICAL FEATURES

SPECTRA 8P is a photometer used for biochemical tests.

Light source	Digitally controlled Halogen lamp (12V - 20Watt)
Wavelength	6 standard interference filters: 340nm, 405nm, 505nm, 546nm, 578nm, 630nm plus 2 optional.
Photometric Range	0 - 3.0 O.D.
Flow cell system	Flow cell with 10mm light path, interchangeable with disposable macro, semi- micro, or special optical glass cuvettes.
Delay time	4 sec.
Reaction volume	300 μL per test min.
Temperature control	Peltier elements, between 15°C - 45°C.
Incubator	8 positions.
Display	Graphic, 240x128 pixel.
Printer	Thermal graphic, 24 characters/line.
Communications	USB port.
Dimensions	L:28 W:36 H:22 cm
Power supply	100-240 VAC 50/60 Hz.

1.2 PRINCIPLE OF OPERATION

SPECTRA 8P is a photometer that measures the concentration of a substance in a liquid via optical principles. Through the implementation of methods designed for the particular substance it is possible to reproduce chemical reactions that turn into a solute capable of absorbing light of a specific wavelength. The physical law that determines the concentration according to absorption of light is known as the Lambert-Beer law:

$$C = k * \log(\frac{l_0}{l_1})$$



Dove:

- I_0 è the intensity of light incident.
- I₁ è the intensity of refracted light.
- **k** is a constant that depends on physical parameters such as temperature T, the optical path ℓ , the wavelength of incident light λ .
- **C** is the concentration of solute.

The quantity $\log(\frac{l_0}{l_2})$ is called as *absorbance (ABS)* or *optical density (OD)*.

It is proportional to the concentration by k. SPECTRA 8P directly measure the absorbance and calculate the factor k using reagents and calibrators. This makes it possible to determine the concentration C.

2. INSTALLATION

The installation of the instrument must be performed by a qualified technician. In any case, the following precautions should always be performed:

- Place the instrument into a dry and clean site.
- Adjust the instrument so that it is slightly tilted towards the operator
- Keeping clean the instrument, especially the front suction tube to avoid contamination between samples and between examinations.
- Use only the power cord supplied or one of equal or characteristics CE marked, CSA and UL. It should be noted that the instrument has internal self-fuse .
- Make sure the power supply is connected to an electrical ground .
- Connect the waste connector on rear side to a bottle using the tube.
- Check O.D. linearity by using periodically standard calibrators.
- If the measured absorbance is greater than 3.000 ABS, repeat the reading or dilute the sample
- Check periodically the regular calibration of the peristaltic pump.
- Keep clean and dry the surfaces of the flow cell.
- Wait at least 15 minutes after switching the instrument to make sure it has reached thermal equilibrium.
- Do not expose the instrument to heat and keep it away from direct sunlight.







3. USER

3.1 PRINCIPAL MENU

The main screen SPECTRA 8P has five items :

SPECTRA 8P

1-EXECUTE METHOD 2-MODIFY METHOD 3-READ ABS 4-SETTINGS 5-FLOW CELL WASH

Using the keyboard you can access all functions of the menu by pressing the button corresponding to the number specified on the left. The possible functions are:

- 1. **EXECUTE METHOD**: It allows to perform a method previously stored in the instrument. After selecting the method (see 3.2) the party suffered from the execution.
- 2. **MODIFY METHOD**: Allows you to change or insert a new method in the instrument. After selecting the method (see 3.2) the part now with the change of the same.
- 3. READ ABS: Allows you to read the absorbance in manual mode. This can be done in two ways: if you

press the button ${\bf 3}$ it is possible to access the reading through cuvette , if you press the button

SWITCH it is possible to access the reading through flow cell.

- 4. **SETTINGS**: Provides access to menu settings.
- 5. FLOW CELL WASH: It allows to perform a washing of the cell flow.

3.2 STRING INPUT

The input of a string is done by inserting a character at a time. There are two ways to make the character input:

- Using button \leftarrow and \rightarrow : Each time you press one of the keys you see the next or previous

character. To go to the following letter must press **ENTER**. To confirm the input press

ENTER two times or press **F1**.

- Using numerical keypad: For each key has an associated set of characters. Pressing the same button repeatedly to display these characters in sequence. Pressing a button other than the one preceding it automatically switches to the next character. To confirm the input press

ENTER two times or press **F1**.

It's possible to use the two way of input simultaneously and independently the same as always active. To delete a character press **DEL** to terminate without any input, press **ESC**.

3.3 NUMBER INPUT

Enter a number via the numeric keys. To confirm press **ENTER**, to delete press **DEL** or to terminate press **ESC**.

3.4 ASPIRATION THROUGH FLOW CELL

The aspiration of the liquid through the cell flow follows these steps:

- **LIQUID ASPIRATION:** The instrument alerts the operator via beeper to immerse the suction tube into the liquid. The open end of the tube should always aspirate the liquid at the bottom of the cuvette. Diagram shows the example of an aspiration from cuvette



Once the tube is immersed in liquid, press button **SWITCH** to start aspiration.

- **LIQUID WASHING:** At the end of the aspiration of the liquid (based on the settings) the instrument alerts the operator via buzzer and wait three seconds to allow the removal of the tube from the liquid
- AIR GAP ASPIRATION: Permits, if enabled, to create a gap of air in aspiration tube

3.5 WASHING OF FLOW CELL

The washing of the cell flow follows these steps:

- **REQUEST OF WASHING LIQUID:** The tool alerts the operator via beeper to immerse the suction tube in the washing liquid. The same requirements used for suction. Once the tube immersed in the liquid,

press the button **SWITCH** to start aspiration.

- **DISCHARGE OF LIQUID:** After aspiration of the liquid (based on the settings) the instrument alerts the operator via buzzer and wait three seconds to allow the operator to remove the tube from the liquid..
- **PAUSE WASHING:** if washing is started manually from the main menu the instrument performs a break (according to settings). During the execution of a washing method is performed without any break
- **WASHING:** The instrument washes the liquid.

3.6 SELECTING A METHOD

The functions 1.EXECUTE METHOD and 2.MODIFY METHOD from principal menù request to perform or modify a method through SELECT METHOD menù:

SPECTRA 8P	
SELECT METHOD:	
1000-SLOT0	
2001-SLOT1	
B002-SLOT2	
4003-SLOT3	
5004-SLOT4	
− ← PREV	NEXT \rightarrow

The instrument has 250 memory slots each capable of containing a method and numbered from 0 to 249. The

list of methods is organized to five pages each and the methods by arrows \leftarrow and \rightarrow you can move from page to page. The selection of the method may occur in two ways :

- QUICK SELECTION: The shortcut through the keyboard pressing the highlighted number (da 1 a 5) to the left of each item.
- NORMAL SELECTION: The normal selection is made directly from the keyboard by entering the number of slots in which the method is stored by pressing **ENTER** and entering the number corresponding to the method.

Pressing the button **PRINT** it's possible print a list of previously stored methods. It's'advisable to print the list in order to quickly locate the slot where is stored the method to select. To enter a new method it is necessary to find a free memory slot and select it.

To return to the main menu press **ESC**.

4. ABS READING

4.1 READING THROUGH CUVETTE

To measure the absorbance of a solute contained in the cuvette from the main menu by selecting the READ ABS. In this way do not use the cell flow. The screen is:

SPECTRA 8P
ABS: 0.000
GFILTER: 340nm G-CELL TEMPERATURE 37.0°C G-TIMER: 0 sec
F1-ZERO F2-ABS

Le possibili opzioni sono:

- 1. FILTER: Repeatedly pressing the button **1**, you can select the wavelength at which to read. Each time you change the filter you need to read again the zero reference of absorbance.
- 2. **CELL TEMPERATURE**: Pressing **2**, you can change the temperature of the cell by entering a new reading. Along the same line you can read the current temperature of the cell.
- 3. TIMER: The instrument allows the setting in seconds of a timer. Pressing **3**, it's possible entering a number of seconds and soon after the release starts the countdown. At the end of the counting instrument alerts the operator via beeper. Pressing any button turns off the buzzer. Using other functions stop the countdown.
- 4. F1-ZERO: Pressing **F1** the instrument reads the absorbance and imposed as the zero reference and

shows **0.000**.

5. F2-ABS: Pressing F2 the instrument reads absorbance and displays the result. Before reading the absorbance of a sample is always necessary to make zero ABS. If this was not done, the instrument indicates it both through the screen and with the buzzer. If you change the wavelength, the zero is canceled and the instrument requires re-reading.

To print reading result, press **PRINT**. To return to the main menu press **ESC**.

4.2 READING THROUGH FLOW CELL

To measure the absorbance of a solute through aspirating peristaltic pump, from the main menu by pressing

the button **SWITCH**. The screen is

SPECTRA 8P		
ABS:	0.000	
	340nm	
B-TIMER:	0 sec	
4-WASH FLO F1-ZE	W CELL RO	

The options are:

- 1. FILTER: Repeatedly pressing the button **1**, you can select the wavelength at which to read. Each time you change the filter you need to read again the zero reference of absorbance
- 2. **CELL TEMPERATURE**: Pressing **2**, you can change the temperature of the cell by entering a new reading. Along the same line you can read the current temperature of the cell.
- 3. TIMER: The instrument allows the setting in seconds a timer. Pressing **3**, it's possible entering a number of seconds and soon after the release party the countdown. At the end of the counting instrument alerts the operator via beeper. Pressing any button turns off the buzzer. Using other functions stop the countdown.
- 4. WASH FLOW CELL: Pressing button **4** will start the process of washing of the cell flow.
- 5. F1-ZERO: Pressing **F1** the instrument reads the absorbance and imposed as the zero reference the

last reading ABS and shows **0.000.**

6. **SWITCH**: Pressing the SWITCH button starts the procedure of aspiration of the sample after which the instrument automatically reads the absorbance and displays. Finally, the flow cell is empitied

To print the ABS reading press **PRINT**. To return to main menu press **ESC**.

5. METHOD MODIFICATION

Select, from the main menu, the function MODIFY METHOD and select the method to change or enter a new one by selecting an empty slot. In both cases, the procedure for release of the parameters is the same: the insertion of a new method is, in fact, consists of a modification of default method that each memory slot already contains. For this reason we will refer only to editing of a new method.

The editing of the parameters is done for pages. Each page contains a number of parameters and the sequence of pages is fixed. Within each page it's possible to move from one parameter to another using the arrow keys

 \leftarrow and \rightarrow and the selected parameter is shown on video by two side arrows \leftarrow e \rightarrow . Pressing the

button **ENTER** it's possible to modify selected parameter. When the operator has completed the changes,

you can go to the next page of parameters by pressing **ESC**. The last page is the one that permits to save

changes. If the operator decides to save, selects **YES**, otherwise, if decides not to save, chooses **NO** and in this case, all changes will be lost

EDITING METHOD #: 0		
NAME: «	<i>(</i>	
TYPE:	END POINT	
ZERO:	WATER	
UNITS:	mg/dL	
TEMP:	37.0 °C	
CALIB:	K-FACTOR	

Pressing the button **DEL** to delete method or pressing **PRINT** to print method.

Depending on the method set some options and some parameters may be present or not. Below is the table of the parameters set for each type of programmable method:

PARAMETER	ENDPOINT	FIXED TIME	KINETIC	NOTE
NAME	YES	YES	YES	Method name
ТҮРЕ	END POINT	FIXED TIME	KINETIC	Method type
ZERO	WATER	WATER	WATER	Zero type
	REAGENT	REAGENT	REAGENT	
	SAMPLE			
UNITS	See table	See table	See table	Unit of measure
				used
TEMP[°C]	YES	YES	YES	Temperature
				execution method
CALIB	K-FACTOR	K-FACTOR	K-FACTOR	Type of calculation
	STANDARD	STANDARD	STANDARD	of the factor k
DECIMALS	0-3	0 - 3	0 - 3	Number of decimal
				digits used in
				displaying and
				printing
SAMPLE[µL]	YES	YES	YES	Sample volume
				specified by the
				method
REAGENT(1)[µL]	YES	YES	YES	Reagent volume
				specified by the
				method
REAGENT2[µL]	NO	NO	YES	Second reagent
				volume specified by
				the method
FILTER(1)	YES	YES	YES	Wavelength of the
				light incident
FILTER2	YES	NO	NO	Wavelength of the
				light incident
				methods for two-
				colored methods

PARAMETER	ENDPOINT	FIXED TIME	KINETIC	NOTE
WEIGHT2	0-1	NO	NO	Weight of the
				second wavelength
K-FACTOR	YES	YES	YES	Factor K
INTERCEPT	SI	NO	NO	Offset
STD NUMBER	1-8	1	1	Number of
				standards or
				calibrators
STD#n	VEC	VEC only STD#1	VEC . only STD#1	Concentration of
310#11	TES	TES - UTILY STU#1	TES - Unity STU#1	the standard n
FITTING CURVE	LINEAR REG	NO	NO	Type of
	LINEAR P-W			approximation for
				multistandard
				methods
READ TIME[sec]	NO	YES	NO	Time reading for
				Fixed-Time
DIRECTION	NO	INCR	INCR	Expected direction
		DECR	DECR	of the reaction
START dABS	NO	YES	YES	Expected direction
				edge of initial
				absorbance
INCUBATION(1)[sec]	NO	YES	YES	Incubation time
INCUBATION2[sec]	NO	NO	SI	Incubation time of
				second reagent
				(only if
				REAGENT2≠0)
RFAGENT BIAS	NO	VES	NO	Select YES to
				include in
				calculating the
				variation of the
				reagent
READINGS #	NO	NO	YES	Number of readings
				to be performed

PARAMETER	ENDPOINT	FIXED TIME	KINETIC	NOTE
DELTA TIME[sec]	NO	NO	YES	Time between
				reading and the
				subsequent
	VEC	VEC	VEC	
	YES	YES	YES	Maximum expected
				concentration
NORMAL MIN	YES	YES	YES	Minimum expected
				concentration
LINEARITY MAX	YES	YES	YES	Concentration
				above which the
				method is no longer
				reliable
REAG. MAX ABS	YES	YES	YES	Maximum expected
				absorbance of the
				reagent
	VEC	VEC	VEC	Minimum expected
REAG. WIIN ADS	TES	TES	TES	absorbance of the
				absorbance of the
				reagent
FIT MIN	NO	NO	0-1	Minimum FIT for
				interpolation curve

6. METHOD EXECUTION

To execute a method choose, from the main menu, the function EXECUTE METHOD and select the method requested. The execution of a method shall be in the form of assisted procedure. Initially it runs a series of preliminary operations common to every type of method (described below). According to the print settings selected, the instrument can print the method before execution.

6.1 REQUEST OF FLOW CELL

The first request relates to the flow cell. If the operator intends to work with the flow cell, he selects the YES button **F1** or **SWITCH**, otherwise the button **F2**. Pressing **ESC** aborts execution. If the operator has chosen to work with cell flow, **SWITCH** button is activated in order to continue execution of the method. Otherwise **SWITCH** is turned off. Depending on the use of flow cell, the instrument will run volume reagent or sample aspirations and washing cycles.



6.2 TEMPERATURE CHECK

The instrument cecks the temperature that is required by the method and expects the cell reading only if the temperature is correct. Alternatively, the operator can skip checking by ignoring the error.



6.3 LAMP CHECK LAMPADA AND BLANK WATER READING

Initially, the instrument takes the reading of a sample of water. This reading is used to check the lamp and, as requested, as zero reference. If the lamp check fails the operator can run it again, leaving the execution or continue ignoring the error message

CHECK LAMP		
VOLTAGE:	3.000 V	
GOOD LAMP CALIBRATION		

6.3 REAGENT CHECK

According to the settings for the method execution, the instrument may request or not the reading of absorbance of working reagent. If in the method running it was specified as white reagent then the instrument performs the reading of absorbance, otherwise the reading is done only if the parameters of maximum and minimum absorbance of the reagent are specified in the method. The absorbance reading is displayed on the screen.



6.4 STANDARD READING

If the method execution is specified with the calibration standard, the instrument requires reading in sequence from the standard number 1. After reading, the operator can accept it (F1/SWITCH), repeat (F2) or

abort (**ESC**) execution method.

METHOD:	NAME

```
ABS READ: x.xxx
ABS STD#n[xxxx]: x.xxx
```

```
F1/SWITCH-NEXT F2-REPEAT ESC-
ABORT
```

During the reading of standard all read absorbances and concentrations specified by the method are printed out. If a multi-standard method is running, after the starting on LCD is shown a graph of read absorbance and

interpolating curve that it's possible to print by pressing the button **PRINT.** In addition to the graph is printed a list of standards and the fit calculated by the instrument.



7. END-POINT METHOD

An end-point method consists of a representation of a reaction that has substance as relatively stable over time. After the operator has mixed the reagent with the analyte, the reaction begins and ends after a period of incubation time. From this moment on and for a sufficiently long period of time, the reaction products remain stable and it is possible to measure the absorbance. The value of concentration is obtained by multiplying the measured absorbance value by a factor k previously calculated on the basis of data provided by the method or obtained from a standard.



8. KINETIC METHOD

A kinetic method consists of a reaction that evolves at a rate determined by the concentrations of substances dissolved in the solution. Started the reaction, the concentrations of substances change and it's possible to determine the initial concentrations by measuring the speed of reaction over the time. The instrument measures the absorbance at fixed time, calculates the speed and after extracts the result of analysis.



9. FIXED-TIME METHOD

A fixed-time method is a kinetic method with only two significant readings: the beginning of reaction and the reaction order. The difference between these two measures related to time allows you to calculate the requested concentrations and therefore the result of analysis.

10. SETTINGS

The settings menu allows to set the operating parameters of the instrument:

```
SPECTRA 8P
SETTINGS MENU:
1-FLOW CELL SETTINGS
2-CLOCK ADJUST
3-PRINTER SETTINGS
4-ADVANCED
01/01/2009 – 12:00
```

The functions available are:

- 1. **FLOW CELL SETTINGS**: This feature allows the opearator to access the configuration menu of the cell flow to adjust the aspiration volume and the calibration of the peristaltic pump.
- 2. CLOCK ADJUST: Allows to adjust the date and time.
- 3. **PRINTER SETTINGS**: Sets the parameters of thermal printer.
- 4. **ADVANCED**: This feature allows the operator to perform maintenance operations such as calibration lamp or thermostats. It is protected by password and should be used only by qualified technical personnel.

10.1 FLOW CELL SETTINGS

The calibration of the peristaltic pump and the configuration of the aspiration air gap through the features of this menu.

FLOW CELL SETTINGS

1-ASPIRATION SETTINGS 2-WASHING SETTINGS 3-PUMP CALIBRATION 4-FLOW CELL WASH

10.1.1 ASPIRATION SETTINGS

The aspiration function settings allows the operator to configure the amount of aspirated liquid volume and optional air gap. The aspiration procedure allows to perform two consecutive aspirations. The first concerns the aspiration fluid test and the second manifold (optional) is in the air and allows the creation of a volume of separation between two consecutive aspirations that contributes both to avoid contamination between successive samples is to use small volumes of sample.

ASPIRATION SETTINGS		
1-VOLUME [u	L]:	300
2-CREATE AIR GAP: ON		
3-AIR GAP VOLUME[uL]: 210		
DEFAULT:	300/C 450/C	DN/210 DN/150

Possible options are:

- 1. VOLUME: Sample volume uL (micro-litres).
- 2. CREATE AIR GAP: ON enables the second aspiration, OFF disabled.
- 3. AIR GAP VOLUME: 2nd aspiration sample volume uL (micro-litres).

It is advisable to avoid using suction volumes lower than the recommended default.

10.1.2 WASHING SETTINGS

The washing function settings allows to set the aspiration volume of washing liquid and the waiting time, ie, the time during which the washing liquid remains in the flow cell. This pause only occurs when the washing is called from the menu, during the execution of a method is never executed

WASHING SETTINGS	
1-VOLUME [uL 2-TIME [SEC]:]: 1000 10
DEFAULT:	1000uL/ 10sec

Possible options are:

- 1. VOLUME: Washing liquid volume uL (micro-litres).).
- 2. TIME: Waiting time in seconds.

10.1.3 PUMP CALIBRATION

PUMP CALIBRATION function allows to calibrate the peristaltic pump using the relation between the rotation of the pump and the volume aspirated. It is advisable to perform regular calibration in case of intensive use of the instrument.

PUMP CALIBRATION	
1-ASPIRATED VOLUM	E[uL]: 0
2-STEPS COUNTED: 0	
3-SAVE CALIBRATION	
F1-TEST ASPIRATION	
F2-TEST WASHING	
SAVED uL/step:	0.000
GENERATED STEPS:	0
	ON TO GENERATE

The peristaltic pump is equipped with a stepper motor. At each step made by the motor corresponds n aspiration volume of the pump. The instrument calculates the number of steps in order to obtain the desidered

volume. Each time the operator press the button **SWITCH** the instrument activates the pump and on LCD

is showed the number of steps generated so that the operator can know the number of steps necessary to obtain a given volume of liquid. The calibration procedure is done as follows

- 1. Prepare a volume of distilled water used for calibration. For example, UL 1000.
- 2. Set the requested volumein the instrument by using option 1.
- 3. Place the aspiration tube into the water and hold **SWITCH** button until the instrument has aspirated all the liquid.
- 4. Read on LCD screen the number of steps generated in Item GENERATED STEPS and enter it into the instrument via option 2.
- 5. At the end of the properly calibration procedure, the operator can save the calibration by option 3 otherwise repeat steps 1 to 4

6. Testing the aspiration and washing through the options F1 and F2 to verify the accuracy of the calibration

11. PRINTER SETTINGS

The configuration of the printer allows the operator to activate it or not both generally and for printing methods and / or results

PRINTER	SETTINGS

1-PRINTER ENABLE: ON 2-AUTO PRINT METHOD: OFF 3-AUTO PRINT RESULTS: ON

The possible options are :

- 1. **PRINTER ENABLE**: activate or not the printer.
- 2. **AUTO PRINT METHOD** :automatically print the parameters of the method at the beginning of each execution.
- 3. AUTO PRINT RESULTS: automatically print the results of analysis.