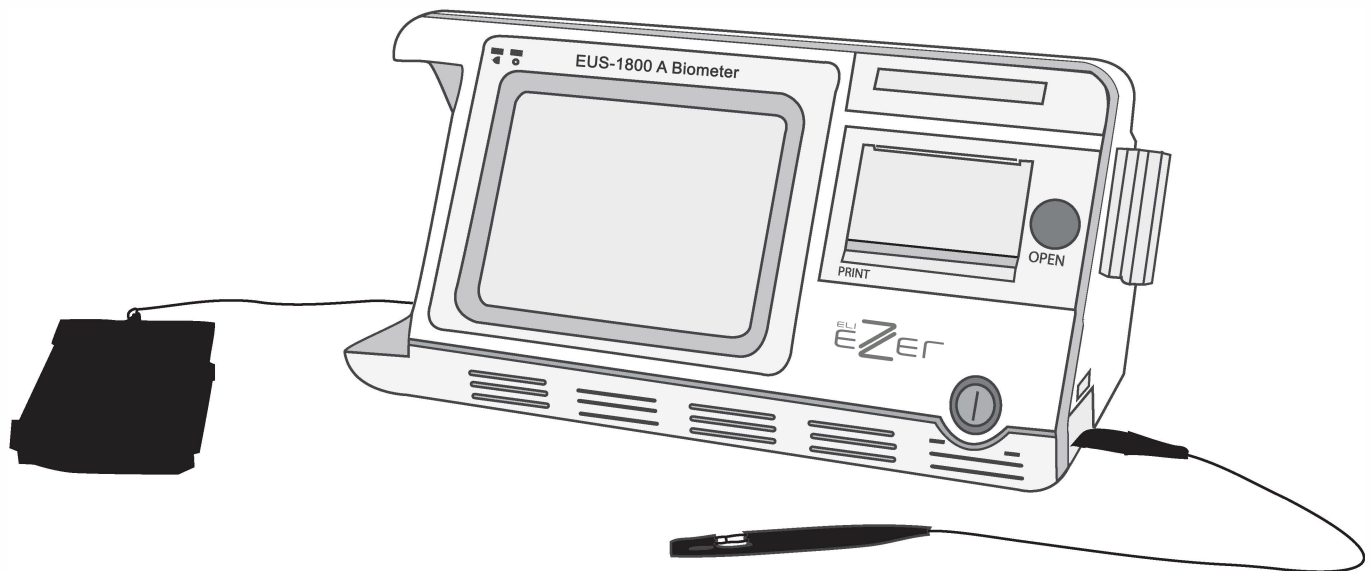


Operation Manual

EUS-1800 A BIOMETER



CONTENTS

PRECAUTIONS	IV
WARNINGS	IV
CAUTIONS	VI
EXPLANATION OF SYMBOLS	VII
CHAPTER 1 INTRODUCTION.....	1
1.1 GENERAL DESCRIPTION.....	1
1.2 INTENDED USE.....	1
1.3 INTENDED LOCATION OF USE.....	1
1.4 CONTRAINDICATIONS.....	1
1.5 EXPECTED SERVICE LIFE.....	1
1.6 SOFTWARE VERSION.....	1
CHAPTER 2 SPECIFICATIONS	2
2.1 WORKING CONDITIONS	2
2.2 MAIN SPECIFICATIONS	2
2.3 SAFETY	2
2.4 STORAGE AND TRANSPORTATION CONDITIONS	2
2.4.1 Storage Conditions	2
2.4.2 Transportation Conditions.....	3
2.5 CLASSIFICATION	3
CHAPTER 3 INSTALLATION.....	4
3.1 STRUCTURE	4
3.2 ENVIRONMENTAL REQUIREMENTS	5
3.3 CONNECTION	5
3.3.1 Connection of Probe.....	5
3.3.2 Disconnection of Probe	5
3.3.3 Connection of Power Supply.....	6
3.3.4 Disconnection of Power Supply.....	6
3.3.5 Connection of Foot Switch	7
3.3.6 Replacement of Printing Paper.....	7

CHAPTER 4 OPERATION	8
4.1 STARTUP AND SHUTDOWN	8
4.1.1 Check before Startup	8
4.1.2 Startup Procedure	8
4.1.3 Check before Use	8
4.1.4 Shutdown Procedure	9
4.2 INTERFACE INTRODUCTION	9
4.2.1 Startup Interface	9
4.2.2 ASCAN Interface	10
4.2.3 Patient Interface (Patient Information Interface)	12
4.2.4 MENU Interface	14
4.2.5 IOL Interface	14
4.2.6 SETUP Interface	17
4.2.7 DATA BASE Interface	18
4.3 OPERATION	19
4.3.1 Positioning	19
4.3.2 A Biometer Unit	19
4.3.3 DATA BASE	28
4.3.4 Display of Saved Information	29
4.3.5 Amendment and Storage of Parameters, and Clock Setting	30
CHAPTER 5 CLEANING AND DISINFECTION	34
FOR EUROPE:	34
5.1 HOW TO PREVENT CROSS-INFECTION	34
5.2 DECONTAMINATION PROCEDURE – PRE-DISINFECTION AND DISINFECTION OF PROBES	34
5.3 PREPARATION OF DECONTAMINATION AGENT	35
5.4 STANDARD METHOD	36
FOR U.S.A	39
5.5 CLEANING OF THE MAIN UNIT OF EUS-1800A	39
5.6 HOW TO PREVENT CROSS-INFECTION	39
5.7 METHOD	39
5.7.1 Preparation	39
5.7.2 Thorough Cleaning	40

5.7.3 <i>Disinfection</i>	40
5.7.4 <i>Rinsing after Disinfection</i>	41
CHAPTER 6 LABELING	42
6.1 EXTERNAL LABEL	42
6.2 INTERNAL ID	42
CHAPTER 7 MAINTENANCE, ATTENTIONS AND SIMPLE DEFECTS TREATMENT	43
7.1 MAINTENANCE AND ATTENTIONS TO INSTRUMENT	43
7.2 MAINTENANCE AND ATTENTIONS TO PROBE	43
7.3 MAINTENANCE OF LCD SCREEN	44
7.4 CHECK BEFORE USE	44
7.5 TROUBLE SHOOTING	45
7.6 DISPOSAL AFTER PRODUCT LIFE-CYCLE	45
CHAPTER 8 SERVICE AND SUPPORT INFORMATION	46
8.1 WARRANTY	46
8.2 ACCESSORIES AND MATERIALS	46
8.2.1 <i>Consumables</i>	46
8.2.2 <i>Detachable Parts</i>	46
8.2.3 <i>Materials</i>	46
8.3 SOFTWARE UPDATE AND SERVICES	46
ANNEX A PRUDENT USE STATEMENT	48
A.1 STATEMENT	48
A.2 PRINCIPLE OF ALARA (As Low As Reasonably Achievable)	48
ANNEX B PROMPT MESSAGE	49
ANNEX C.1 ACOUSTIC OUTPUT REPORT (IEC60601-2-37)	50
ANNEX C.2 ACOUSTIC OUTPUT REPORTING TABLE	51
ANNEX D GUIDANCE AND MANUFACTURER'S DECLARATION	52
ANNEX E IOL FORMULA	56
ANNEX F REFERENCES	58

PRECAUTIONS

- The device should be operated by trained technicians.
- It is prohibited to use the device on patients with eye trauma, inflammation or infection.
- Please read the manual carefully before installation and operation
- Please refer to Chapter 5. Cleaning and Disinfection to avoid cross-infection while using.
- Disconnect AC power before cleaning the housing case.
- Please refer to Chapter 7. Maintenance, Attentions and Simple Defects Treatment for maintenance attentions.

WARNINGS

- The customer is fully responsible for maintenance and management of the instrument after purchasing.
 - Do not make any modification to the software and hardware of the device without authorization.
 - The power adaptor provided with the device meets the safety standard of medical electrical devices. If damaged, contact the Manufacturer to purchase. Use of other adaptors may cause safety risk.
 - The manufacturer won't be responsible for any damage or injury caused by any failure to follow the instructions in the User Manual.
 - The manufacturer reserves the right to modify equipment characteristics without previous notice under FDA Laws and MDD (93/42/EEC) Regulation.
 - The quality guarantee of the device will be invalid if it is opened (even partially), modified or repaired in any way by anyone who is not authorized by the manufacturer.
-
- This device is not intended for therapeutic use.
 - It is prohibited to use the device on patients with eye trauma, inflammation or infection.
 - The device should be used cautiously on patients without independent behavior abilities or who are highly sensitive to contacting measurements, whose ineffective cooperation may result in inaccurate measurements.
 - According to FDA laws, the EUS-1800A is a prescription Device and is to be used by or under the supervision of a licensed physician.

- This device is not intended for fetal use.
- While plugging in the probe, make sure the red mark on the probe align with the red mark on the socket.
- While unplugging the probe, please be sure you are pulling the connector instead of the cable.
- Do not scratch the surface of the probe.
- Do not drop the probes.
- If the probe drops during using or moving, check the top and the housing of probe carefully, and then check if it works well. Stop using if there is any problem and contact the manufacturer or local distributor for service.
- Warnings of predicable potential hazards are contained in the User Manual. Please maintain vigilance at any time to those unpredictable hazards. The manufacturer won't be responsible for damages and losses caused by negligence or ignorance of the preventive measures in the **User Manual**.
- The assembly, expanding, readjustment, improvement and repair should be operated by personnel authorized by the manufacturer. Do not open the housing for repair without permission. The manufacturer won't be responsible for the consequences of safety and effectiveness caused by unauthorized repair.
- Keep the original package properly. All detachable accessories should be put into the original package before moving and transportation.
- Without written consent of the manufacturer, no individual or organization is allowed to copy, modify, or translate any part of the **User Manual**.

For any question, please contact the Manufacturer or your Local Distributor



**Address: 9990 NW 14ST
Suite 105
Doral, FL 33172**

Phone: +1-888-334-4640

Fax: +1-786-621-1842

Email: info@usophthalmic.com

CAUTIONS

HOW TO PREVENT CROSS-INFECTION

- It is prohibited to use the device on patients with eye trauma, inflammation or infection.
 - Between uses on different patients, the probe and eye cup must be thoroughly cleaned and disinfected to prevent cross-infection.
 - The Manufacturer advocates a preventive action and a cleaning procedure in Chapter 5. Cleaning and Disinfection.
-

CAUTION

- Federal (US) Law restricts this device to sale by or on the order of a physician.
 - The EUS-1800A IOL calculator will calculate negative IOL values if such is predicted by the data input.
 - These are displayed with a minus sign "-". Do not ignore this sign.
-

TISSUE EXPOSURE TO ULTRASOUND ENERGY:

- The EUS-1800A is designed for use in ophthalmology only.
- While the manufacturer is not aware of any reports of adverse effects from using ophthalmic ultrasound scanner, even at FDA pre-enactment levels, no other use is intended or implied.
- The system controls limit of the output energy within the parameters specified for its intended purpose. Please refer to Annex A of the **User Manual**.
- No control of ultrasound energy is available to the users other than the duration of exposure, considering the current concern for possible unknown hazards, and despite the extremely low output intensities used in this ultrasound system.
- The manufacturer recommends that patients' exposure time during measurement be minimized.

EXPLANATION OF SYMBOLS

Front Panel



Power On or Power Off

OPEN

Paper Housing Switch

PRINT

Thermal Printer

Rear Panel



Symbol of "Type B"



Refer to User Manual



Serial Number



CE Conformity Marking



Manufacturer



European Representative

FOOT SWITCH

Foot Switch Socket



Power Input Socket

Right Panel

A-Probe

A Biometric Probe Socket

User Manual



NOTE! Important information that operator must read carefully.

IPX1/IPX7

The degree of protection against ingress of liquids.

Chapter 1 Introduction

1.1 General Description

The EUS-1800A A Ultrasonic Biometer for Ophthalmology is an ultrasonic measuring instrument based on pulse reflection. It comprises A Mode Eye Axis Biometric Parameter Measuring Unit (A Biometer).

The A Biometer consists of a 10MHz A-Scan probe (probe model: Prb1000A/10-C) and an axis biometric parameter measuring unit. The axis is usually divided into three segments: anterior chamber, lens and vitreous body. Since the tissue within the eye varies in different areas, the acoustic velocity through these areas is also different. The summation of these three segments (ACD + LENS + VITR) provides the axial length (AL). Based on the interface reflections of these three different tissues, the ultrasonic A-biometry measures the transmitting time of ultrasound through each tissue and calculates the length of each segment to finally get the axial length.

The EUS-1800A has a built-in Thermal Printer, used to print out patient information, A-Scan measuring waveform, IOL calculating parameter and result.

The built-in memory of the EUS-1800A can store up to 180 patients' records.

1.2 Intended Use

The EUS-1800A is intended for axial biometric parameter measurement in clinical ophthalmology.

1.3 Intended Location of Use

The EUS-1800A is suitable to be applied in hospitals and ophthalmology clinics. [The device should be operated by trained technicians.](#)

As the foot switch's protective degree against ingress of liquids is IPX1, it is not suitable to be used in operating room and other locations where it is easy to splash liquid.

1.4 Contraindications

It is prohibited to use the device on patients with eye trauma, inflammation or infection.

1.5 Expected Service Life

Based on the experience of products sold (and considering technology update cycle), as well as the aging of the major part – transducer will probably reduce the basic performance, the product's expected service life is determined as five years according to the normal usage of six hours each working day. (Service life may extend if the major parts, like probe, are returned to manufacturer for update.)

1.6 Software Version

MD AP10X V1.0

Chapter 2 Specifications

2.1 Working Conditions

Environmental Temperature: 5°C—40°C

Relative Humidity: $\leq 80\%$

Atmospheric Pressure: 70kPa—106kPa

The Separate Power Supply: Input: AC100~240V, 50/60Hz,

Output: DC12V, 4A

Rate Power: 50 VA

2.2 Main Specifications

Ultrasonic frequency: 10MHz;

Display resolution: 0.01mm;

Total gain of receiver: $\geq 100\text{dB}$;

Adjustable gain scope: 0~50dB;

Measuring scope (AL): 15 mm~40mm;

Measuring accuracy: $\leq \pm 0.05\text{mm}$;

Measuring Parameter: ACD, LENS, VITR and AL;

Measuring mode: automatic mode or manual mode can be selectable. Automatic mode for Normal, Cataract and Aphakic mode;

Measuring method: contact and immersion, can be selectable;

IOL calculation: SRK/T, SRK-II, BINK-II, HOLLADAY, HOFFER-Q, HAIGIS, and IOL calculation formula after corneal refractive surgery: HISTORY DERIVED, DOUBLE-K/SRK-T, REFRACTION DERIVED, ROSA and SHAMMAS.

2.3 Safety

Satisfy the requirements of IEC 60601-1 and IEC 60601-2-37.

The acoustic output parameters: see Annex C.

2.4 Storage and Transportation Conditions

2.4.1 Storage Conditions

The device should be stored in non-corrosive gas and well-ventilated room with environmental temperature of -20°C—40°C and relative Humidity $\leq 80\%$.

2.4.2 Transportation Conditions

The accessories such as Probe should be packed into the original package before transportation. Severe impact and crash, rain and snow shall be avoided.

2.5 Classification

As per the type of protection against electric shock: Class I

As per the degree of protection against electric shock: Type B

As per the degree of protection against ingress of liquids:

- Main unit: IPX0;
- Part of Probe that can be immersed: IPX7;
- Foot switch: IPX1

As per the disinfection method recommended by the manufacturer:

- see §5 Cleaning and Disinfection.

As per the safety degree when used under flammable anesthetic gas mixed with air or under flammable anesthetic gas mixed with oxygen or Nitrous oxide: not allowed.

As per working mode: continuous working

Chapter 3 Installation

3.1 Structure

The structure of the instrument: See Figure 3.1.

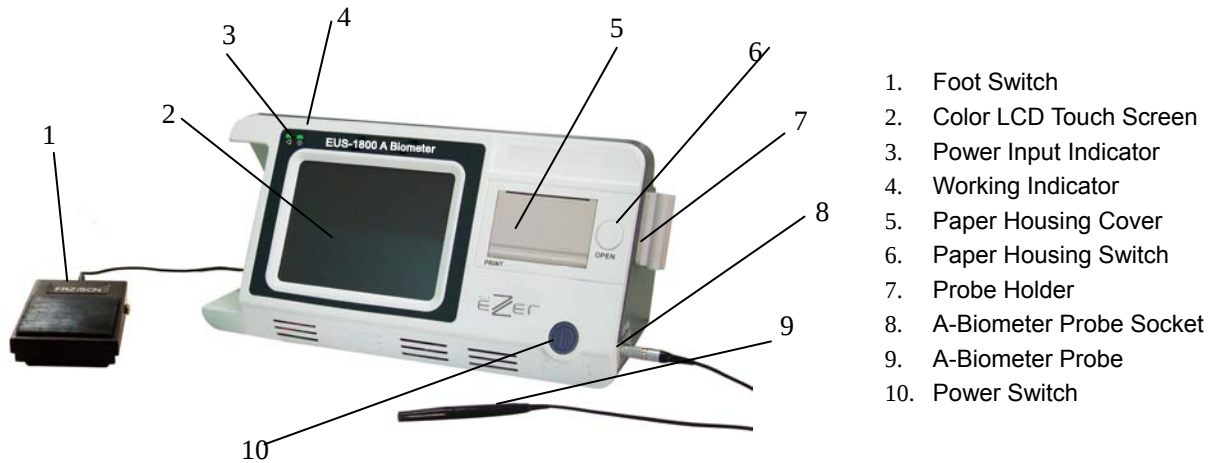


Figure 3.1 Structure of the Instrument

The Rear Panel: See Figure 3.2

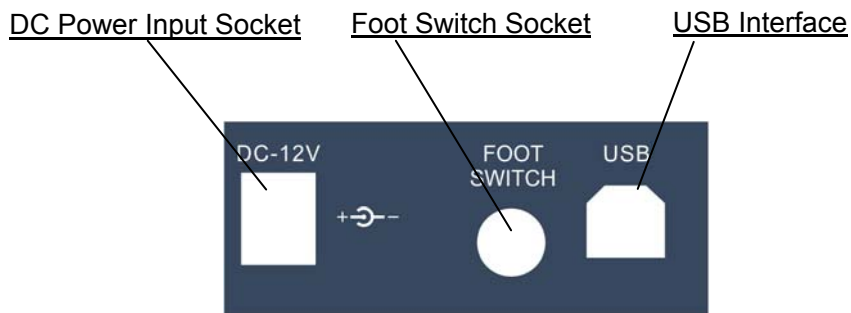


Figure 3.2 Rear Panel of the Instrument

The power adaptor of the product, with input voltage of AC100-240V and output voltage of DC12V/4A, meets the safety requirements of medical electric device, as shown in Figure 3.3:



Figure 3.3 Power Adaptor

3.2 Environmental Requirements

- 1) The instrument should be operated in a clean environment. Air-conditioned environment is recommended.
- 2) The instrument should be placed on a stable worktable or platform. Avoid direct sunlight.
- 3) Please use the supplied Power Adaptor which is in accordance with the safety standard of medical electric equipment; do not use any other adaptors.
- 4) The power outlet should be with good grounding. Improper connection of protective earth may cause not only interference, but also the risk of increasing leakage current.
- 5) Do not block the ventilation window of the instrument. When the temperature of the instrument is not normal, please contact the manufacturer for service.
- 6) Although the EMC of the instrument meets the requirements of IEC 60601-1-2, it should be placed to avoid strong electromagnetic radiation equipment (such as microwave, radio frequency therapy equipment, etc.).
- 7) Please do not put the instrument in a place where it is difficult to disconnect the power supply.
- 8) The instrument should be placed in a position where the operator can face the instrument and view the screen easily. The distance between the instrument and the patient should enable the probe to contact the patient's eye conveniently.
- 9) The instrument should always be placed safely to prevent falling of the probe.
- 10) The instrument has no special protective measures for discharge effect of cardiac defibrillators; it is not suitable for use with high-frequency surgical equipment.

3.3 Connection



Note: It is prohibited to plug in or plug out any accessories while the instrument is running.

3.3.1. Connection of Probe

Plug the A-Biometer probe into the A-Probe socket on the right panel;



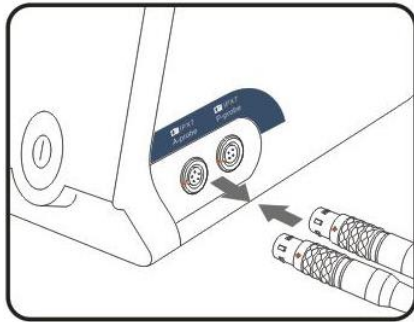
Note: While plugging in the probe, make sure the red mark on the probe align with the red mark on the socket, as shown in Figure 3.4 (a).

3.3.2. Disconnection of Probe

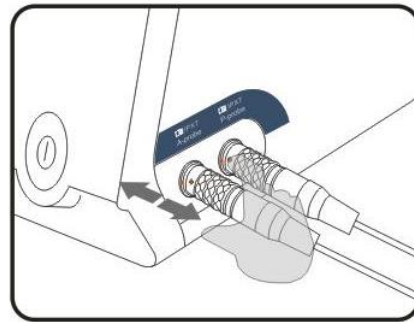
To disconnect the probe, please hold the ring of the connector and pull it out along the horizontal direction, as shown in Figure 3.4 (b).



Note: Don't pull the probe cable strongly while unplugging the probe.



(a) Connection of Probe



(b) Disconnection of Probe

Figure 3.4 Connection/Disconnection of Probe

3.3.3. Connection of Power Supply

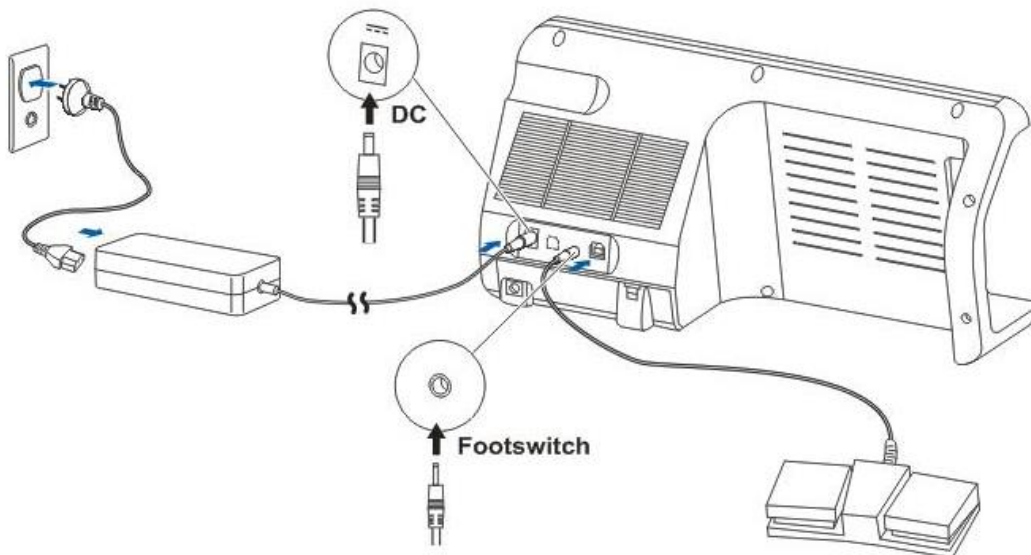


Figure 3.5 Connection of External Interface

- 1) Plug the DC Power Output Plug of the Power Adaptor in the Power Input Socket on the rear panel of the instrument;
- 2) Connect one end of the Power Cable with the Power Adaptor, and the other end to the wall outlet, as shown in Figure 3.5.

3.3.4. Disconnection of Power Supply

- 1) Turn off the Power Switch on the front panel;
- 2) Disconnect the main power plug of the Power Adaptor and the wall outlet.



Note: Take care of safety while connecting or disconnecting the power supply.

3.3.5. Connection of Foot Switch

Plug the Foot Switch connector into the Foot Switch Socket on the rear panel of the instrument, as show in Figure 3.5.

3.3.6. Replacement of Printing Paper

The built-in printer is a thermal printer that uses thermal printing paper. Please purchase and replace the printing paper according to the specific model provided by the Manufacturer, see [§8.2.1 Consumables](#).

Replacement of printing paper:

- 1) Press "OPEN" button to open the cover of the paper housing;
- 2) Take out the reel of the paper from the paper holder and add new paper, see Figure 3.6;



Figure 3.6 Replacement of Printing Paper

- 3) Pull out the printing paper from the paper slot for about 1cm, and push lightly to close the cover of the paper housing. Pay attention not to let the paper stuck in the slot. Close the paper housing finally.



Note: 1) If the paper doesn't run properly, please check installing method and reinstall.

- 2) The thermal paper has thermal side, when the paper runs normally but nothing printed out, please open the paper housing and reinstall the printing paper with the other side.

Chapter 4 Operation

4.1 Startup and Shutdown

4.1.1. Check before Startup

Before the instrument is started up, please check:

- 1) Whether the instrument is put in a proper place and whether the equipment around may cause interference;
- 2) Whether the appearance of the instrument is fine and whether there is crack on the housing and LCD screen;
- 3) Whether the probe is connected properly, and whether there is crack or damage on the surface of probe and cable; whether the probe cables are intertwined or intertwined with other cables;
- 4) Whether the foot switch is connected properly and has mechanical response when press, and whether the foot switch cable is intertwined with other cables;
- 5) Whether there is printing paper; and
- 6) Whether the connection of power supply is completed and the Power Indicator lights up; whether the Power Cable is intertwined with other cables.

4.1.2. Startup Procedure

- 1) Make sure there is no problem and the Power Indicator lights up before the instrument is started.
- 2) Start the instrument: Press the Power Switch button on the Front Panel for over 2 seconds, the instrument starts working. Now the Working Indicator lights up and the main interface appears on the screen.

4.1.3. Check before Use

After the instrument is started up, please check:

- 1) Whether the Working Indicator lights up;
- 2) Whether the touch screen is effective;
- 3) Whether waveform displayed when clicking **SCAN/FREEZE** key under the ASCAN Interface, and whether the red indicator in the front end of the probe lights up when measurement;
- 4) The parameter setting, following §4.3.6 Amendment and Storage of Parameters, and Clock Setting for operation in detail; and
- 5) Whether disinfection of the probe has been carried out as required; whether the surface of the probe is deformed or damaged and whether the probe cable is damaged. In case eye

cup is used, check whether the eye cup has been disinfected and whether there is visible damage.

4.1.4. Shutdown Procedure

While the instrument is switched on, press the Power Switch on the front panel for 2 seconds, the instrument will switch off.

Place the instrument and its accessories properly after shutdown:

- 1) Put the probe into the probe holder and make sure the probe cable can't be dragged by accident, or disconnect the probe and put it properly;
- 2) Put the foot switch properly and make sure the foot switch cable is placed properly to prevent others from stumbling by the cable and drop of instrument;
- 3) Plug out the Main Power Plug of the Power Adaptor cable and put it properly.

4.2 Interface Introduction

4.2.1. Startup Interface

Switching on the EUS-1800 A, the buzzer will beep shortly and then enter the Login Interface (see Figure 4.1).

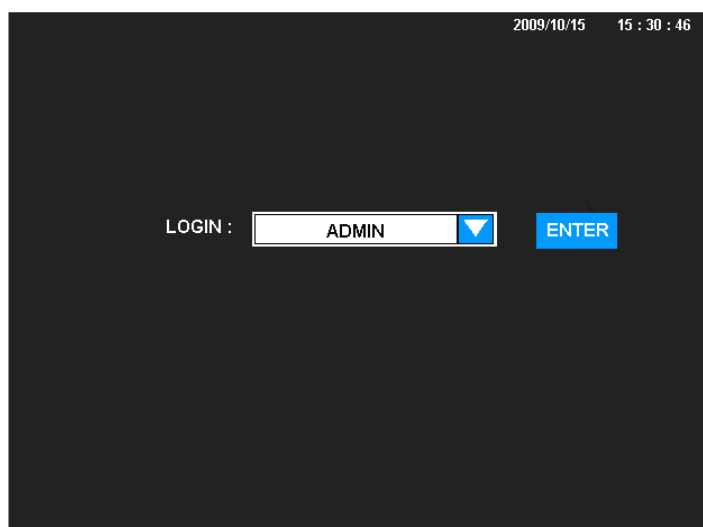


Figure 4.1 Login Interface

Click **ENTER** key directly under the Login Interface will enter the ASCAN Interface of the default ADMIN user system. Click **▼** to select other user system, see Figure 4.2; after selection, click **ENTER** key to enter the corresponding ASCAN Interface of the relevant user system. See §4.3.6 for amendment of the user name, in which ADMIN is the particular system name that can't be amended.

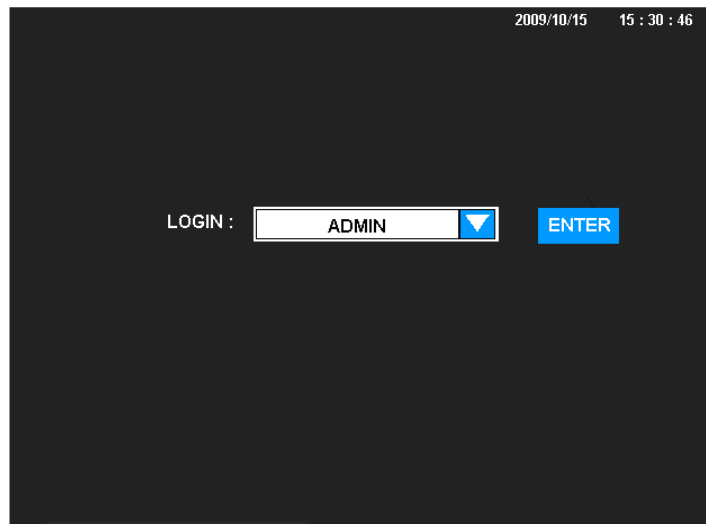


Figure 4.2 Login Interface

Click **ENTER** key directly under the Login Interface will enter the ASCAN Interface of the default ADMIN user system. Click ▼ to select other user system, see Figure 4.3; after selection, click **ENTER** key to enter the corresponding ASCAN Interface of the relevant user system. See §4.3.6 for amendment of the user name, in which ADMIN is the particular system name that can't be amended.

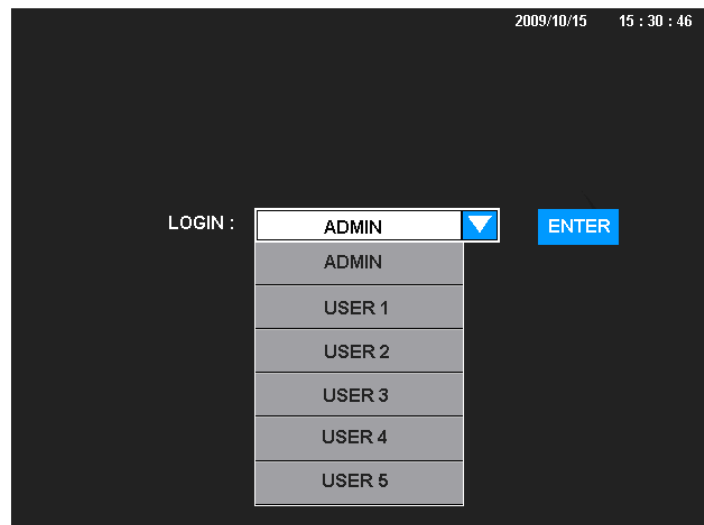


Figure 4.3 System Selection Interface

4.2.2. ASCAN Interface

Enter the ASCAN Interface, the default setting of the system is automatic measurement (AUTO), contact method (CONTACT) and normal eye (NORMAL), the interface is shown in Figure 4.4.

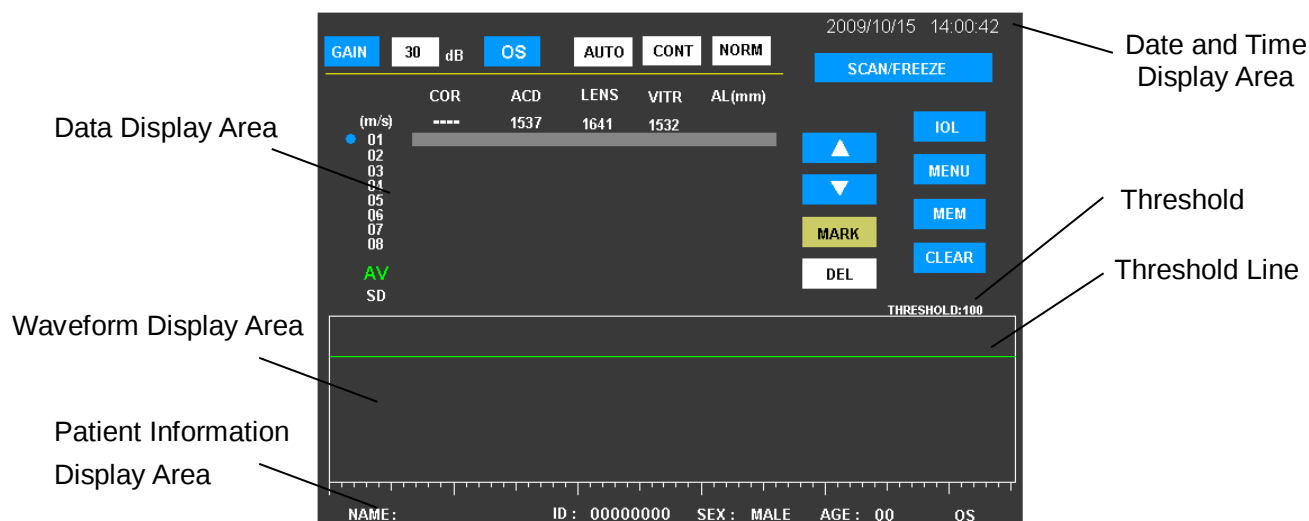


Figure 4.4 ASCAN Interface

The function of each part under the ASCAN Interface:

Data Display Area: The result of measurement display area.

Waveform Display Area: ASCAN ultrasonic echo waveform display area.

Patient Information Display Area: Display the input patient information. Click this area to enter the Patient Information Interface.

Threshold: A value associated with voltage. The echo pulse with amplitude more than the value is determined as the reflective wave of acoustic interface for different tissues. The cross point of the reflected pulse front and the threshold will be the basic point to calculate transmission time of ultrasound in different tissues.

Threshold Line: The threshold value displayed in graph.

Date and Time Display Area: Display current date and time.

GAIN: Gain control key. Range of variation: 0-50dB; the default value is 30dB. Click to select and the key turns to green. The value of gain is adjusted by ▲ and ▼ keys when it is selected;

OS: OS/OD (left/right eye) selection key. Click to switch between left eye and right eye;

AUTO: Measuring mode selection key. Click to switch between AUTO (automatic measurement) and MANL (manual measurement);

CONT: Measuring method selection key. Click to switch the measuring methods between CONT (contact method) and IMME (immersion method);

NORM: Eye model setting key. Click to open the MODE SETTING dialogue box and set eye model. The MODE SETTING dialogue box includes the settings of: Measuring Modes, Measuring Methods and Eye Models, see §4.3.2.2.c for specific operation;

SCAN/FREEZE: Scan/Freeze control key. Click to start or freeze scanning. The key is blue indicates the state of scanning stopped; while green indicates scanning;

 : Up and Down keys, used to adjust the position and value. The keys are used to adjust the position of the Threshold Line when the Threshold Line displayed, and to adjust the gain when the GAIN function is activated. In other cases, they are used to change the position of the cursor indicator (blue point) and cursor bar. The position of the cursor and cursor bar indicates the position of the current operating data group;

MARK: Mark key, used to mark unreliable data group. The marked group stores data and waveform, but doesn't participate in calculation. The cursor stays in the marked group, click again to cancel the mark;

DEL: Delete key, used to delete unreliable data and waveform. Click to call out the dialogue box. Choose YES to delete the selected data and waveform. The deleted data will not be restored. Choose NO to give up deletion and quit the dialogue box;

CLEAR: Clear key, used to clear data and waveform. Click to call out the dialogue box. Choose YES to clear all data and waveform displayed. The cleared data will not be restored. Choose NO to give up clear and quit the dialogue box, see §4.3.2.6 Save and Clear under Measuring Interface;

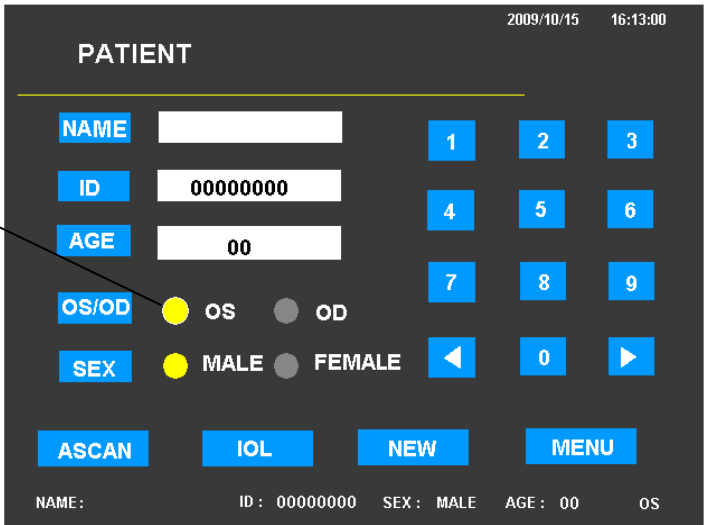
IOL: IOL (Intraocular Lens) Calculation key. Click to enter the IOL Interface;

MENU: Menu key. Click to enter the MENU Interface, from where it is able to enter other interfaces;

MEM: Memory key. Click to save data, which includes patient information, measuring data, all kinds of parameters and waveforms. If it's necessary to save complete information, please enter the IOL Interface, complete parameter input and calculation, and then click **MEM** key to save.

To enter the PATIENT Interface directly from this interface, it's able to click directly the Patient Information Display Area at the bottom of the screen.

4.2.3. Patient Interface (Patient Information Interface)



The screenshot shows the 'PATIENT' interface with a dark background and white text. At the top right, the date '2009/10/15' and time '16:13:00' are displayed. The interface includes several input fields and buttons:

- NAME**: A text input field.
- ID**: A text input field containing '00000000'.
- AGE**: A text input field containing '00'.
- OS/OD**: Radio buttons for 'OS' (selected) and 'OD'.
- SEX**: Radio buttons for 'MALE' (selected) and 'FEMALE'.
- Navigation Buttons**: A 3x3 grid of buttons labeled 1 through 9, a left arrow, 0, and a right arrow.
- Action Buttons**: Four buttons at the bottom labeled 'ASCAN', 'IOL', 'NEW', and 'MENU'.
- Status Bar**: A row at the bottom showing 'NAME:', 'ID: 00000000', 'SEX: MALE', 'AGE: 00', and 'OS'.

An arrow labeled 'Indicator' points to the 'OS/OD' radio button area.

Figure 4.5 PATIENT Interface (Patient Information Interface)

In normal measuring interface, for example the ASCAN Interface, click the Patient Information Display Area at the bottom of the screen to enter the Patient Information Interface directly. The Patient Information Interface is used to input patient information, including patient name, ID, age, OS/OD and sex accordingly.

Click the position of the corresponding key, the cursor appears, and then input the information. Patient name should be no more than 12 characters. Click the text box of **NAME** will call out the alphabetical keypad, click the keypad to enter patient name (as shown in Figure 4.6). Click the text box of **NAME** again to cancel the alphabetical keypad and complete inputting. ID is made up of 8 digits with the input range from 00000000 to 99999999, click the numerical key at the right side to input numbers when the cursor is located in the text box. Input age with the same method and the range for age is from 00 to 99.

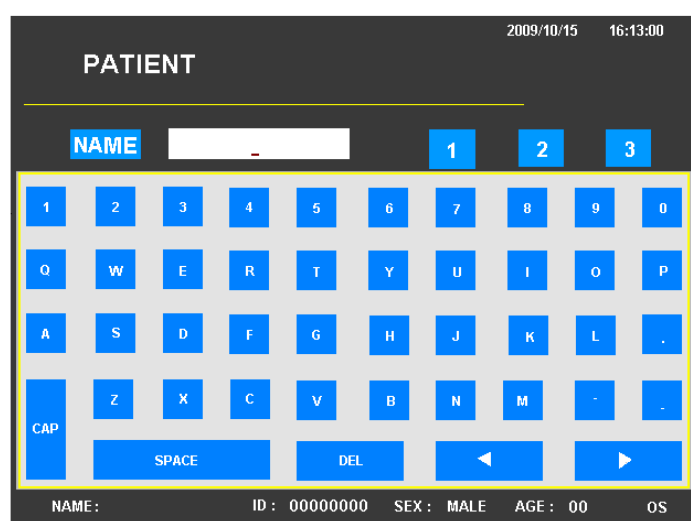


Figure 4.6 Patient Information Interface (Alphabetical Keypad)

There are two ways to select OS/OD: one is clicking **OS/OD** key, the OS and OD lights up alternatively (the indicator lights up means selected); the other is clicking the area of OS or OD directly for selection.

Patient's sex can be set with the same method.

ASCAN: Click it to choose returning to the ASCAN Interface;

IOL: Click it to choose returning to the IOL Interface;

NEW: New Patient key. Click it, the Patient ID will be restored to default value, new patient information can be input at the moment, meanwhile all data measured in other interfaces will be cleared;

MENU: MENU key. Click it to return to the MENU Interface.

Numerical keys are used to input number 0-9; the **◀** **▶** keys are used to control the left and right movement of the cursor respectively.

Under the PATIENT Interface, in case of changing into other interfaces, the patient information entered will be displayed at the bottom of the screen. The displayed information contains: name, ID, age, sex and OS/OD for reference. If the information is not input, the default patient name is empty, the ID is: ----- and the age is: --.

4.2.4. MENU Interface

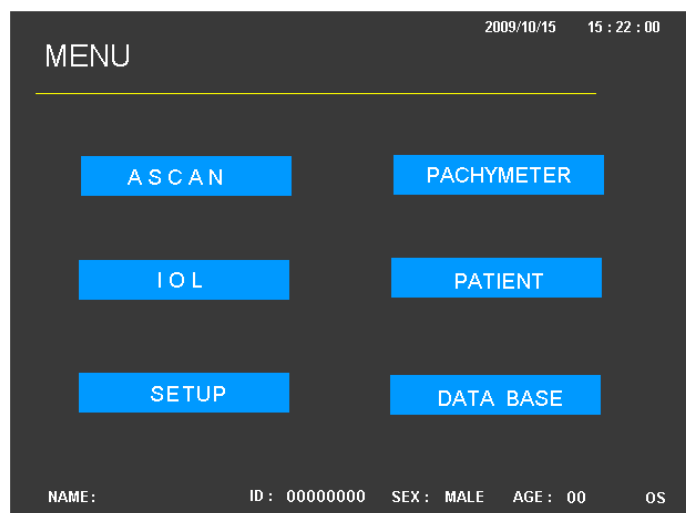


Figure 4.7 MENU Interface

Click **MENU** key to enter the MENU Interface, as shown in Figure 4.7. Click the six keys in the interface to enter the corresponding interface, which are: ASCAN Interface, PACHYMETER Interface, IOL Interface, PATIENT Interface, SETUP Interface and DATA BASE Interface.

4.2.5. IOL Interface

The IOL Calculation Interface, as shown in Figure 4.8. The keypad at the bottom of the interface is used to input numbers.

AL: Axial Length AL, which can be achieved by the measuring result of A Biometer, or input by the keypad;

AC: Anterior Chamber depth AC, which can be achieved by the measuring result of A Biometer, or input by the keypad;



Note: AC can only be used in HAGIS formula; it is inactive in other formulas.

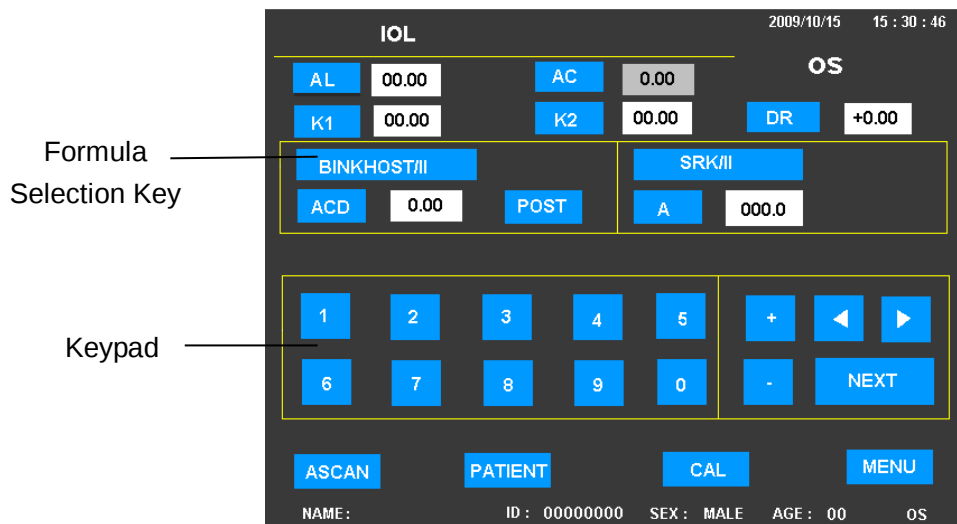


Figure 4.8 IOL Interface

There are two formula selection areas for user's choice of an applicable IOL calculation formula. Click Formula Selection key to select.

If the IOL calculation formula after corneal refractive surgery will be selected, click K1 or K2 key to call out the Parameter Input Dialogue Box (as shown in Figure 4.9). Select YES to light up the Indicator and then click the Formula Selection Key to open the pull-down menu. Select the desired formula and input parameters.

The formulas provided are: SRK-II, SRK/T, HOLLADAY, HOFFER-Q, HAIGIS, BINK-II and IOL calculation formulas after corneal refractive surgery (HISTORY DERIVED, DOUBLE-K/SRK-T, REFRACTION DERIVED, ROSA and SHAMMAS).

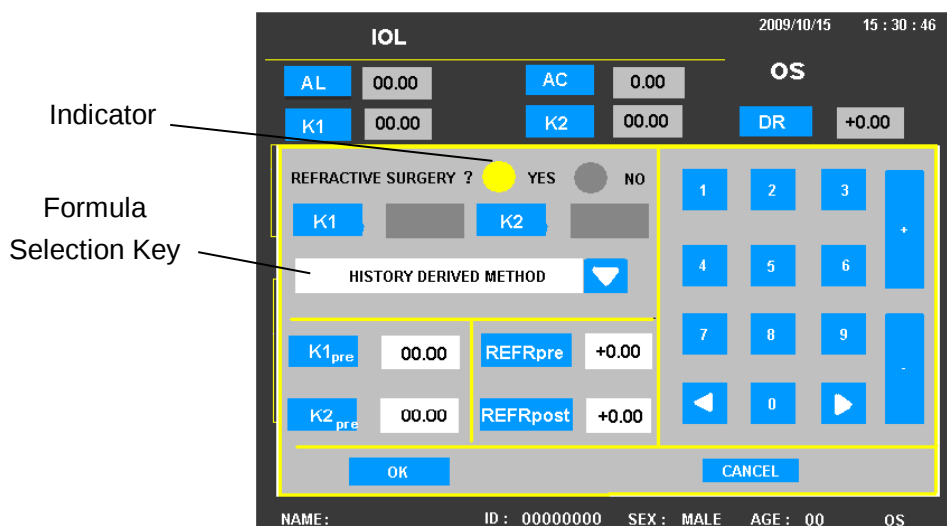


Figure 4.9 IOL Interface (Parameter Input Dialogue Box)

The parameters required for each formula can be input according to the specific circumstance. Click the parameter name key, the cursor will appear in the corresponding data box. Click the numerical and symbolic keys on the keypad to input each parameter. The constant A/ACD can be

input directly by clicking the corresponding data box or by clicking A/ACD key to call out the Constant A/ACD Input Dialogue Box to input (as shown in Figure 4.10).

The screenshot shows the IOL Interface with the following elements:

- Top Bar:** IOL, 2009/10/15, 15:30:46
- Input Fields:**
 - AL: 00.00, AC: 0.00, DR: +0.00
 - K1: 00.00, K2: 00.00
 - BINKHOST/II: ACD: 5.55, POST
 - SRK/II: A: 000.0
- Numeric Keypad:**
 - Row 1: 1, 2, 3, 3.21, 3.39, 3.68
 - Row 2: 4, 5, 6, 4.15, 5.55, 5.26
 - Row 3: 7, 8, 9, 5.37, 5.61, 6.02
 - Row 4: <, 0, >, EDIT, OK
- Bottom Bar:** ASCAN, PATIENT, CAL, MENU
- Patient Info:** NAME: ID: 00000000 SEX: MALE AGE: 00 OS

Figure 4.10 IOL Interface (Constant A/ACD Input Dialogue Box)

The meaning of each parameter is as follows:

DR: Refraction Desired after IOL implantation;

K1: Current Keratometry, click it to call out parameter dialogue box;

K1pre: Keratometry before corneal refractive surgery;

K2: Current Keratometry, click it to call out parameter dialogue box;

K2pre: Keratometry before corneal refractive surgery;

A: Constant of IOL, click it to call out Constant A/ACD Input Dialogue Box;

REFRpre: Refraction before corneal refractive surgery (the amount of lens power of spectacles);

REFRpost: Refraction after corneal refractive surgery (the amount of lens power of spectacles);

ANTI/POST: Anterior Mode/Posterior Mode;

D.EM: Diopter of Emmetropia;

D.AM: Diopter of Ametropia;

EDIT: Click it to enter or exit from constant A/ACD editing status;

OK: Click it to exit from Constant A/ACD Input Dialogue Box;

ASCAN: Click it to return to the ASCAN Interface;

PATIENT: Click it to return to the PATIENT Interface;

MENU: Click it to return to the MENU Interface;

CAL: Calculation key, click it to make calculation and enter the interface of calculating result.

When clicking **CAL** key, if the input parameter is out of range, the system will prompt to make amendment and then recalculate.

When the calculation completed, the keypad will disappear and the calculating result will appear on the screen. Meanwhile, the keys will change, as shown in Figure 4.11.

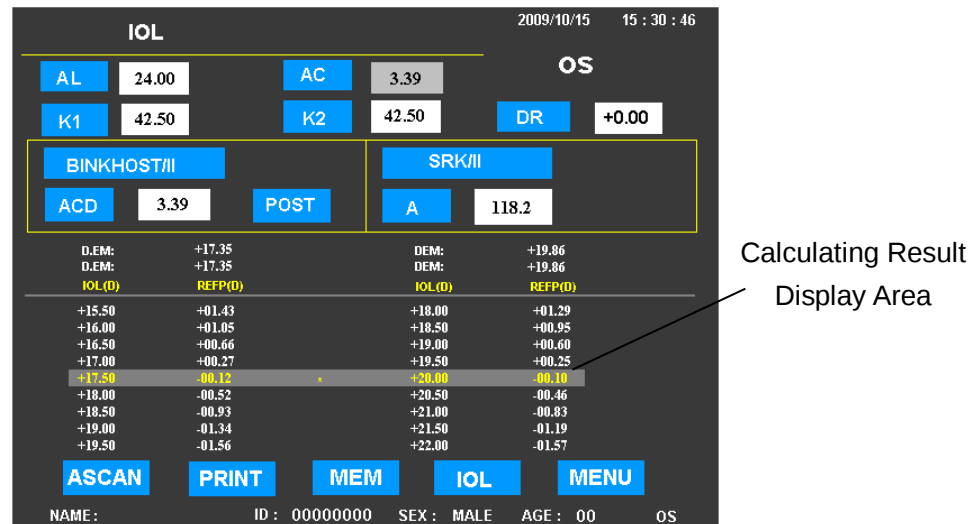


Figure 4.11 IOL Calculation Interface

MEM: Memory key, click it to save all the related information, including the patient information, measuring data, waveform and IOL calculating parameters.

PRINT: PRINT key, click it to print the saved result.

IOL: Click it to return to the IOL Interface, calling out the keypad, changing parameters and recalculating. Click the corresponding data box under this interface, the system can switch to the calculation interface.

4.2.6. SETUP Interface

The Setup Interface, as shown in Figure 4.12.

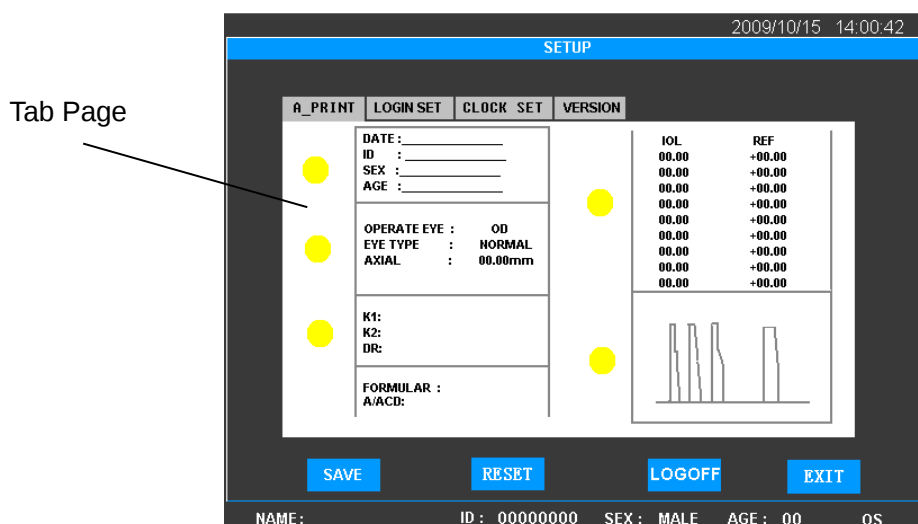


Figure 4.12 SETUP Interface

The Setup Interface consists of three parameter setup tab pages. Click the corresponding key to enter each tab page.

A-PRINT: Click it to enter the print parameter tab page for A Biometer;

LOGIN SET: Click it to enter the user setup tab page (it is not available under ADMIN system).

CLOCK SET: Click it to enter the clock setup tab page;

The functions of other keys under the SETUP Interface are as follows:

VERSION: Click it to enter version information tab page;

SAVE: Click it to save selected result;

RESET: Reset key. Click it to restore default value;

LOGOFF: Click it to cancel the system and reenter into the Login Interface after 1 second;

EXIT: Click it to exit from the interface.



Note: 1) Under other tab pages except that of **CLOCK SET**, click **SAVE** key will save all parameters in the **A-PRINT** and **LOGIN SET** tab pages; and click **PRESET** key will restore all parameters to default settings. Please confirm before operation.

2) Click **SAVE** or **PRESET** key will save the status of all parameters automatically, so please make the related change cautiously.

4.2.7. DATA BASE Interface

The DATA BASE Interface is for patient information management, as shown in Figure 4.13.

NO.	NAME	ID	AGE	SEX	OSOD	TYPE	TIME
1	MARK	00000001	50	MALE	OS	A	2009/06/03 09:30
2	JIM BROWN	00000005	50	MALE	OS	P	2009/06/06 09:00

OK CLEAR DELETE EXIT

NAME: ID: 00000000 SEX: MALE AGE: 00 OS

Figure 4.13 DATA BASE Interface

This interface lists all the saved patient information in tabular format. The contents displayed include patient name, patient ID, age, sex, OS/OD, measuring type (A Biometer) and storage time. Patient information can be checked. Click the line of corresponding patient information, the blue bar appears, then click **OK** to enter the Measuring Data Display Interface of the save information and observe the measuring result. (Please refer to §4.3.3 DATA BASE and §4.3.4 Display of Saved Information)

CLEAR key is used to clear all the patient information, clicking this key will call out a dialog box to confirm the request of clear or not. Choose YES to clear all information which can not be restored; choose NO to exit directly. Click **DELETE** key to delete the selected information only. All deletions are not recoverable, please operate cautiously. Click **EXIT** key to exit.

4.3 Operation

4.3.1 Positioning

Place the instrument beside the patient. The distance between should enable the probe to contact the patient's eye easily. The instrument should face the operator and the foot switch should be placed appropriately for operation. Please ensure the instrument is located safely and will not cause falling and hurting people by accidental operation. Switch on the instrument after completing the above steps.

4.3.2 A Biometer Unit

4.3.2.1 Input Patient Information

In the ASCAN Interface, click the Patient Information Display Area at the bottom of the screen to enter the PATIENT Interface, (or click **MENU** key to enter the MENU Interface, then click **PATIENT** key to enter the PATIENT Interface) and input patient's information accordingly. The default patient name is empty, patient ID is: ----, age: -----, and sex: MALE. Please update the patient information in time to prevent any error.

4.3.2.2 Mode Selection

a) Selection of measuring mode

There are two kinds of measuring modes: automatic measuring mode (AUTO) and manual measuring mode (MANUAL). The default mode is automatic measuring mode.

Manual measuring mode is usually used in some special cases, when the system can not identify the waveform or can't mark automatically due to the limited conditions.

Five-Point Marking Method is used to measure A Biometer data, as shown in Figure 4.14.

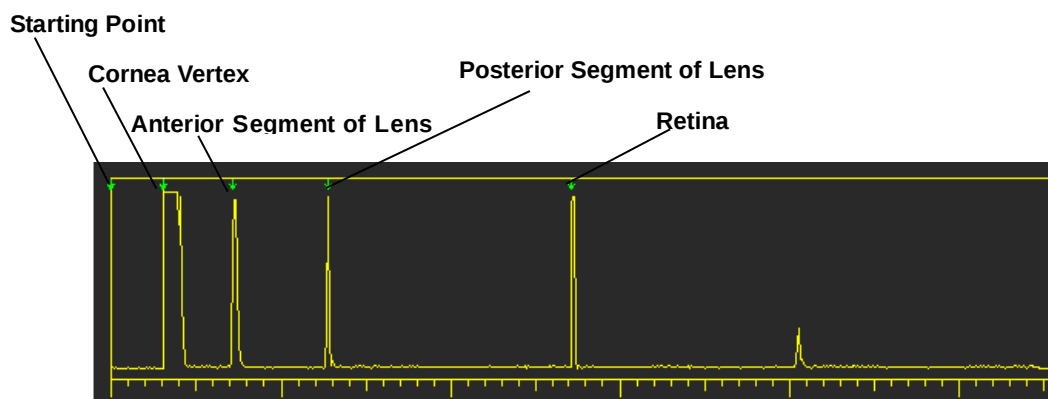


Figure 4.14 Five-Point Marking Method

The five points are respectively: cornea vertex, cornea posterior segment (sometimes in superposition with cornea vertex), anterior segment of lens, posterior segment of lens and retina.

In AUTO mode, the system will automatically determine the positions of the five points and process data. The system's default setting under AUTO mode is NORMAL eye and CONTACT method.

In MANUAL mode, the operator is required to determine the positions of the five points and mark them separately to acquire measuring data. The measuring method and eye model selection are invalid in MANUAL mode. The operator can mark the position of the arrow manually according to the selected measuring method. Pay attention that the position of the starting point of the Five-Point Marking Method for contact method is different with that of the immersion method.

With contact method, the starting point is in superposition with cornea vertex and cornea thickness is ignored, that is to say the cornea vertex is in the superposition with the cornea posterior segment.

With immersion method, the starting point is not in superposition with cornea vertex, and cornea thickness can be measured in some cases, at that time the cornea vertex is not in superposition with the cornea posterior segment.

The specific selection method for measuring mode is: click **AUTO** or **MANL** key to switch between AUTO and MANUAL, the content displayed will change accordingly (display **AUTO** or **MANL**). Select a suitable mode, or click **NORM** key to open the MODE SETTING dialogue box. Please refer to item d) of this section.

b) Selection of measuring method

According to whether A-Biometer probe will contact cornea, the measuring method is divided into CONTACT method and IMMERSION method.

CONTACT method indicates the A-Biometer probe will contact cornea vertex directly. The operator is required to handle the probe gently while measuring. It is applicable for both automatic and manual measurement.

IMMERSION method indicates the A-Biometer probe will not contact cornea directly, but through acoustic coupling medium.

The specific selection method is: click **CONT** or **IMME** key to choose measuring method, the content of key will change accordingly (display **CONT** or **IMME**). Select a suitable method, or click **NORM** key to open the MODE SETTING dialogue box. Please refer to item d) of this section.

c) Selection of eye model

According to the different conditions of patients, four EYE models can be selected, that are: NORMAL, APHAKIC, SPECIAL and CATATACT. Open the MODE SETTING dialogue box to select. Please refer to item d) of this section.

d) MODE SETTING dialogue box

Click **NORM** key to call out the dialogue box, as shown in Figure 4.15.

The dialogue box consists of three parts, namely: measuring mode setting, measuring method setting and eye model setting.

Click the keys respectively to select AUTO or MANUAL; CONTACT or IMMERSION; NORMAL, APHAKIC, SPECIAL or CATARACT.

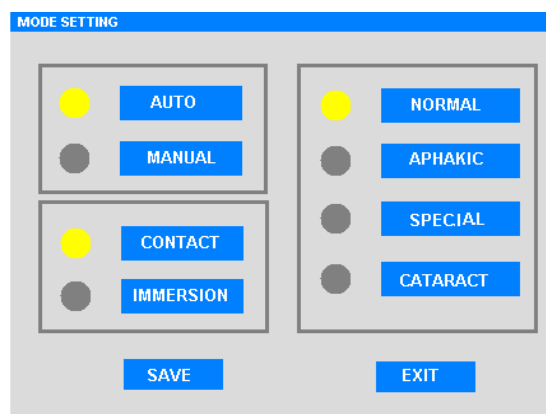


Figure 4.15 Mode Setting Interface

The acoustic velocity is the system's default setting. If CATARACT or APHAKIC is selected, the system will change the acoustic velocity automatically.

If special eye is selected, the dialogue box pops up (as show in Figure 4.16) for selection of the velocity of several common IOL. The types of IOL to be selected are: PMMA, ACRYLIC, SILICONE or USERSET, among which, USERSET is the type provided to users to input the required acoustic velocity according to patient's condition.

Click specific position to select the corresponding IOL acoustic velocity (indicator lights up), which is displayed in the dialogue box. The value can be modified but can't be saved as the instrument's parameter. It is only used under the current condition when SPECIAL eye is selected, quit the

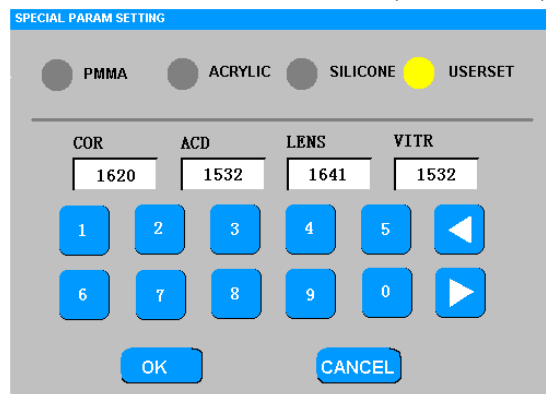


Figure 4.16 IOL Acoustic Velocity Setting

selection, then the input or selected acoustic velocity will disappear. Click **OK** key to save the set acoustic velocity; or click **CANCEL** key to exit (no parameter in this mode will be saved).

The Default Acoustic Velocity set is shown below:

Unit: m/s

IOL	PMMA	ACRYLIC	SILICONE	CATARACT
Acoustic Velocity (m/s)	2718	1946	1050	1629

4.3.2.3 Measurement under AUTO Mode

1) CONTACT Method

Switching on the instrument will enter AUTO mode automatically, or select AUTO mode according to the description of **Mode Selection** in section §4.3.2.2a, the default measuring method is CONTACT, and the operation procedure is as follows:

- a) Let the patient lie down, open his/her eye to perform surface anesthesia.
- b) Step onto the foot switch or click **SCAN/FREEZE** key to start measuring.
- c) Place the probe gently onto the patient's cornea vertex, and tell the patient to look at the probe straightly.
- d) Generally, it is not necessary to adjust the gain and threshold. For some special cases (e.g. severe cataract patients), low retina reflective amplitude or poor waveform maybe appear, adjust the gain or threshold then to get the clear reflective waveform of the tissue interface, sharp front edge is preferred.
- e) When the probe is properly aligned, you will hear a series of beep sounds, the waveform is automatically frozen and data will be displayed on the screen. The EUS-1800A supports displaying up to eight groups of data, and there will be a long beep sound after all eight groups of data are acquired. The instrument will stop measuring automatically after the data displays on the screen, and the probe can be removed then. The average value of each group and standard deviation display on the upper side of the waveform display area.
- f) If the data are not acquired, please move the probe slightly until getting the measurement. Keep the probe stably in the position found, eight groups of data will be acquired.
- g) If the standard deviation is big, it may caused by the moving of probe during measurement. Move the cursor with **▲** **▼** keys and observe each group of data and its corresponding waveform. Click **MARK** key to mark unreliable data, click again to cancel the mark. The marked data will not be calculated. Or click **DEL** key to delete the inaccurate data.



Note: 1) The deleted data will not be calculated and are unrecoverable, so please operate cautiously.

2) Since the results of each automatic measurement will subject to multiple sampling averages, the operator is required to perform the scanning gently

and smoothly. The probe can only be removed after the measurement completed, the image is automatically frozen and the data is displayed on the screen.

- 3) The probe should be placed align with the visual axis to guarantee the accuracy of measurement.
- 4) Don't move the probe directly on the patient's cornea to avoid injury of cornea.

2) IMMERSION Method

Under automatic measurement, if IMMERSION method is selected according to §4.3.2.2b, the operator should gently put the eye cup onto patient's eyelid, fix the eye cup with eyelid and inject physiological saline into eye cup as coupling medium. Place the A-Biometer probe into the coupling medium, and make sure the probe will not contact the cornea. The probe should be placed perpendicular to the cornea and moved towards the cornea. When the distance between the probe front and cornea vertex is 2-8mm, start automatic measurement. The rest operations are the same with that of the CONTACT method.

4.3.2.4 Threshold modification under the AUTO mode

The threshold can be set according to the actual demand. The threshold adjustment range is 50-120 and the default state is that the threshold line is hidden and the threshold value is 100. Double click the upper part of the waveform quickly to display the threshold line and the current threshold value, as shown in Figure 4.17. Adjust the position of the threshold line by clicking the up and down keys. Click the threshold line to hide the threshold line and the threshold value.

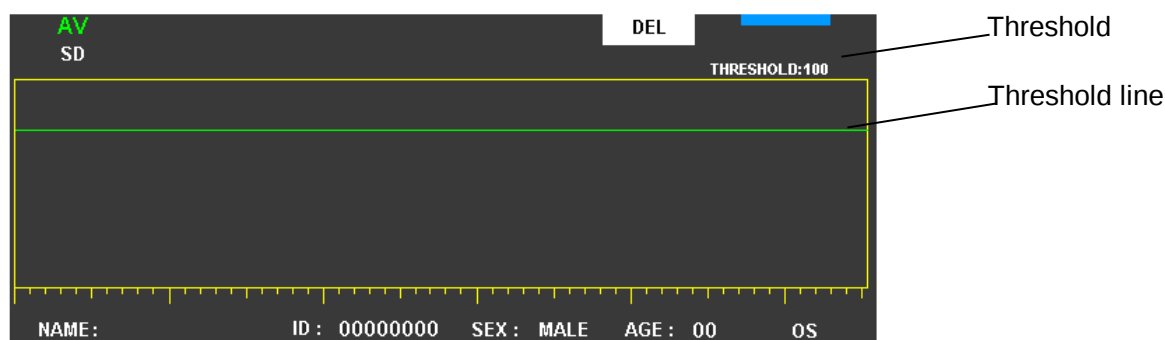


Figure 4.17 Diagram of Threshold Line



Note: 1) Normally, the threshold value is not necessary to be modified and the default setting is recommended.

2) The threshold value will return to default setting after the system is switched off.

4.3.2.5 Measurement under MANUAL mode

Select manual measuring mode according to §4.3.2.2a), the ARROW MOVE dialogue box and **ARROW MOVE** key will pop up automatically. The ARROW MOVE dialogue box is used to move

the arrow for marking the peak of wave and amend the data displayed. **ARROW MOVE** key is used to call out the ARROW MOVE dialogue box for remarking the arrow. See Figure 4.18:

The ARROW MOVE dialogue box contains **◀ ▶** keys, **ENTER** key, Zoom In and Zoom Out keys, as well as **AMEND** key and **EXIT** key. The **◀ ▶** keys are used to move the arrow; **ENTER** key is used to confirm the arrow position; Zoom In key is used to magnify the part indicated by the arrow and Zoom Out key is used to cancel magnification and return to normal state. Click **ENTER** key after all arrow positions are selected, the measuring result will be displayed in the position of the first group and then switch to the next group's measurement. Click **EXIT** key will exit from this interface directly. If amendment is required, click **AMEND** key to clear all arrow positions and re-select.

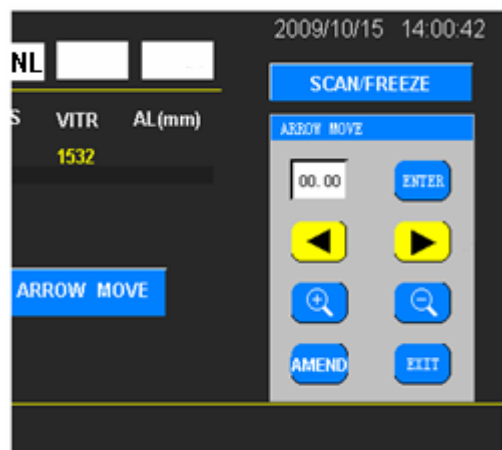


Figure 4.18 Arrow Move Dialogue Box

Pay attention that under MANUAL mode, the position of the starting point of the Five-Point Marking Method for CONTACT method is different with that of IMMERSION method. (See §4.3.2.2)

For CONTACT method, the procedure is as follows:

- a) Select manual measuring mode, click **SCAN/FREEZE** key or step on the foot switch to start scanning.
- b) Operate according to procedure a) to e) for automatic measuring mode.
- c) Move the probe slightly to observe the reflected waveform. When the reflected waveform is clear, click **SCAN/FREEZE** key again or step on the foot switch to freeze the waveform. The first group of waveform displays as —.—.—.
- d) Since contact method is used, the starting point is the cornea vertex and is in superposition with the posterior segment of cornea. Click **ENTER** key twice and confirm the position of cornea vertex and posterior segment. Then click **◀ ▶** again to move the cursor, and the moving distance of the cursor displays in the dialogue box; when the cursor is moved to the reflected pulse's rising edge for the anterior segment of lens, click **ENTER** key and the value displayed in the dialogue box will input in the column of ACD. Keep on moving the cursor to acquire VITR value with the same method. After the above measurement, click **ENTER** key again, the measuring value of AL displays on the screen, and at the moment the measuring waveform of the first group disappears, the blue dot cursor skips to the second group for the measurement.
- e) There are Zoom In and Zoom Out keys in the ARROW MOVE dialogue box. The Zoom In key is used to magnify five points within left and right range with the cursor as the center. If accurate positioning is required during measurement, click Zoom In to move the cursor accurately inside the magnified area. Click Zoom Out to cancel magnification and return to normal state.

f) If the arrow positions need to be amended during the process of determination, click **AMEND** key, the arrow disappears; then remark the arrow position.

g) Click **EXIT** key after measurement and exit from the dialogue box. And click up and down keys to load the waveforms of each group of the measuring data.

Under MANUAL mode, up to 5 groups of data can be acquired and the average value and standard deviation display at the upper side of the waveform display area. For uncertain measuring data group, the operator can click **MARK** key to mark. When necessary, click **DEL** key to delete the data or click **ARROW MOVE** key to call out the ARROW MOVE dialogue box and make amendment by clicking **AMEND** key and remarking the position of arrow. The deletion and amendment of data will influence the final calculation of the average value and standard deviation. The final results will come into the IOL calculation.

If IMMERSION method is used under manual measurement, attentions should be given for marking the measuring waveform: for the case the starting point of the waveform is not in superposition with the cornea vertex, clicking **◀ ▶** first to move the arrow to the reflected pulse's rising edge of the cornea vertex, click **ENTER** key to confirm; if the reflected waveform can't distinguish the posterior segment of the cornea, click **ENTER** key again to ignore the cornea; if the reflected waveform can distinguish the posterior segment of the cornea, then move the arrow to the reflected waveform's rising edge of the posterior segment of the cornea, click **ENTER** key then, the cornea thickness is input in the COR column of the screen. The measurement for other segments is the same with that of the CONTACT method.

4.3.2.6 Save and Clear under Measuring Interface

In the ASCAN Interface, the measuring information and data can be saved. Click **MEM** key to save the current results, the system will pop out the dialogue box to confirm the operation. Choose **YES** to save and choose **NO** to exit.

Please refer to §4.3.3 and §4.3.4 for load and display of the saved information.



Note: Save in the ASAN Interface can store patient information, waveform and measuring data; the calculated parameter and result can not be saved. If the calculated parameter and result need to be saved, please enter the IOL Interface to calculate and save.

In the ASCAN Interface, if the measured data and waveform are not satisfactory and need to be re-measured, please click **CLEAR** key to clear all the data and waveforms. Click **CLEAR** key, the system will pop out the dialog box to confirm the operation. Choose **YES** to clear, and the cleared data will not be recovered. Choose **NO** key to exit without clearing.

4.3.2.7 Manual Amendment of Measuring Result under AUTO Mode

For experienced doctors, if the standard deviation of the eight groups of data, which is acquired by automatic measurement, is big, it is able to move the cursor with up and down keys and observe the measured waveform corresponding to each group of data; and for the uncertain data that the

arrow does not align with the rising edge of the waveform, it's possible to remark manually. The detailed operation is as follows:

a) Double click the data display area quickly to call out the ARROW MOVE dialogue box as shown in Figure 4.19, and amend the data indicated by cursors.

b) Click **ENTER** key after the dialogue box is called out. The color of arrow will change to red successively. Select the position of arrow that need to be amended manually and click **AMEND** key to confirm the amendment at the arrow position.



Figure 4.19 Arrow Move Dialogue Box

c) The **◀ ▶** keys will change from un-selectable grey to selectable yellow. Click **◀ ▶** keys to move the arrow to the rising edge of the waveform, if required, select ZOOM IN and ZOOM OUT to observe the waveform. The operation procedure is the same with the ZOOM IN and ZOOM OUT function of the above Manual Measuring Mode.

d) Move the arrow to the required position and click **ENTER** key again to input the new result, the arrow's selectable status will move to the position of the next arrow. Repeat the above process to make amendment.

e) Click **EXIT** key to exit.

4.3.2.8 IOL Calculation

The measuring result will display in the IOL Interface automatically after measurement. Click **IOL** key in the ASCAN Interface or click **IOL** key in the MENU Interface can enter the IOL Calculation Interface (see §4.2.5).

In the IOL Interface, the measuring result of AL can be acquired by the calculation interface directly, or be input manually.

Select the required formula by clicking the Formula Selection Key (see Figure 4.8). Different formula will appear by each click. The formulas appear repeatedly until the required formula appears. The EUS-1800A provides the following IOL formulas: SRK-II, SRK/T, HOLLADAY, HOFFER-Q, HAIGIS, BINK-II. If the IOL calculation formula after corneal refractive surgery will be selected, click K1 or K2 to call out the Parameter Input Dialogue Box (as shown in Figure 4.9), select YES and then click the Formula Selection Key to open the pull-down menu for selection: HISTORY DERIVED, DOUBLE-K/SRK-T, REFRACTION DERIVED, ROSA and SHAMMAS.

After the YES indicator is lit up and the required formula is selected, input parameters according to the selected formula. Click **OK** key in the Parameter Input Dialogue Box to confirm the parameter and exit, at the moment, "REFRACTIVE SURGERY CASE" will be displayed on the top center of the interface (as shown in Figure 4.20). The value of K1 and K2 displayed in this interface are the values after corneal refractive surgery, which can only be amended in the Parameter Input Dialogue Box. Please refer to Annex D for the selection of the IOL calculation formulas after corneal refractive surgery and the selection of parameters. Click **CANCEL** key in the Parameter Input Dialogue Box to cancel the selection and exit.

IOL (REFRACTIVE SURGERY CASE) 2009/10/15 15:30:46

AL 24.00 AC 3.39 OS

K1 42.50 K2 42.50 DR +0.00

BINKHOST/II SRK/II

ACD 3.39 POST A 118.2

1 2 3 4 5 + ◀ ▶

6 7 8 9 0 - NEXT

ASCAN PATIENT CAL MENU

NAME: ID: 00000000 SEX: MALE AGE: 00 OS

Figure 4.20 IOL Interface (IOL calculation mode after corneal refractive surgery)

Use the keypad to input parameters required for each formula according to the specific case. The constant A/ACD can also be input by calling out the Constant A/ACD Input Dialogue Box (as shown in Figure 4.21).

IOL 2009/10/15 15:30:46

AL 00.00 AC 0.00 DR +0.00

K1 00.00 K2 00.00

BINKHOST/II SRK/II

ACD 5.55 POST A 000.0

1 2 3 3.21 3.39 3.68

4 5 6 4.15 5.55 5.26

7 8 9 5.37 5.61 6.02

◀ 0 ▶ EDIT OK

ASCAN PATIENT CAL MENU

NAME: ID: 00000000 SEX: MALE AGE: 00 OS

Figure 4.21 IOL Interface (Constant A/ACD Input Dialogue Box)

Click the constant key of **A** or **ACD** can call out the Constant A/ACD Input Dialogue Box, meanwhile the selected key turns to green indicating the status of selected (as shown in Figure 4.21). In the Constant A/ACD Input Dialogue Box, 9 groups of constants A/ACD in common use are listed. Users can make selection according to the listed items by directly clicking the corresponding data box. The selected data turns to blue, indicating the status of selected. Click **OK** to exit.

If the listed data need to be amended, click **EDIT** key, the data box will change to editing status, meanwhile the **EDIT** key turns to green (as shown in Figure 4.22).

The screenshot shows the IOL interface with the following layout:

- Top Bar:** IOL | 2009/10/15 | 15:30:46
- Parameter Fields:**
 - AL: 00.00 | AC: 0.00 | DR: +0.00
 - K1: 00.00 | K2: 00.00
- Input Section:**
 - BINKHOST/II | SRK/II
 - ACD: 5.55 | POST | A: 000.0
- Keypad:**
 - Grid of numbers 1-9, 0, and navigation arrows.
 - Values displayed in the grid: 3.21, 3.39, 3.68, 4.15, 5.55, 5.26, 5.37, 5.61, 6.02.
 - Buttons: EDIT (green), OK (grey).
- Bottom Bar:**
 - Buttons: ASCAN, PATIENT, CAL, MENU
 - Fields: NAME: | ID: 00000000 | SEX: MALE | AGE: 00 | OS

Figure 4.22 IOL Interface (Constant A/ACD Input Dialogue Box - Editing Status)

Select the data to be amended, click the data box to move the cursor in and input the required data with the keypad on the left. Click **EDIT** key again, the system will save the input parameter and the **EDIT** key turns to blue. Click **OK** to exit.

Click **CAL** key to make calculation after all parameters are input. The calculating result will display on the screen. The operator can select to print out or save the result according to the practical situation.

4.3.3 DATA BASE

Click **DATA BASE** key in the MENU Interface to enter the data base and check patient information.

In each measuring interfaces, if data are saved after measurement, the patient's information will be saved in the data base.

The screenshot shows the DATA BASE interface with the following layout:

- Top Bar:** DATA BASE | 2009/10/15 | 15:30:46
- Table:**

NO.	NAME	ID	AGE	SEX	OSOD	TYPE	TIME
1	MARK	00000001	50	MALE	OS	A	2009/06/03 09:30
2	JIM BROWN	00000005	50	MALE	OS	P	2009/06/06 09:00
- Bottom Bar:**
 - Buttons: OK, CLEAR, DELETE, EXIT
 - Fields: NAME: | ID: 00000000 | SEX: MALE | AGE: 00 | OS

Figure 4.23 DATA BASE Interface

The data base will display the saved patient information in tabular format, as shown in Figure 4.23. The displayed information include: patient name, ID, age, sex, OS/OD and type of measurement (A Biometer).

The operator can select the patient information (by clicking the position of the patient information, blue bar means selected). Click **OK** to call out the Measuring Data Display Interface and observe the measured result. At the moment, some of the functions are invalid in the Measuring Data Display Interface. The invalid function keys turn to grey and there is no response after clicking. Please refer to §4.3.4 for details.

CLEAR key is used to clear all patients' information. Click this key and the dialogue box will be called out to confirm the operation. Select "YES" to confirm clearance of all data. The data cleared are not recoverable. Select "NO" to exit without clearing. Click **DELETE** key to delete one patient's information. The use of **DELETE** key is similar with that of **CLEAR** key. Click **EXIT** key to exit.



Note : 1) All deletions are unrecoverable, so please operate cautiously.

- 2) The maximum storage capacity of the EUS-1800A is 180 items of patient information. Please process the data in time to prevent storage error.
- 3) When the information storage is large, the time required to read patient information will be longer, please wait patiently.

4.3.4 Display of Saved Information

The EUS-1800A has the function of saving and displaying the measuring data. The saved measuring interface and data at the time of measuring can be loaded according to the operator's selection, which is convenient for observation and analysis. See Figure 4.24.

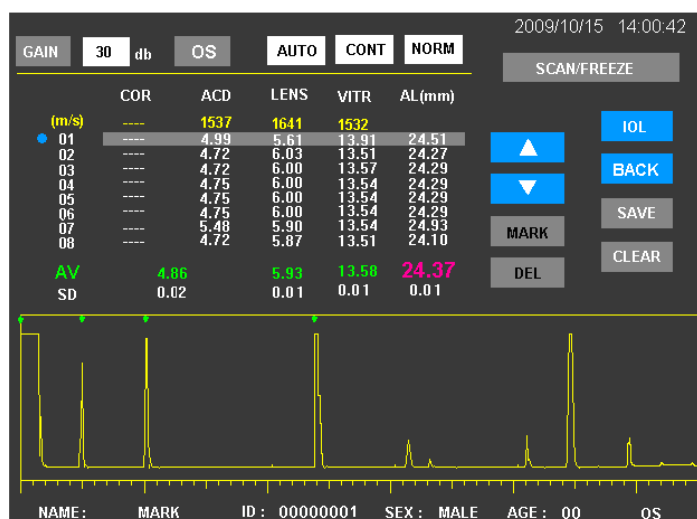


Figure 4.24 Measuring Data Display Interface

The operating method is as follows:

- 1) Enter into the DATA BASE (as shown in Figure 4.23) to check patient information list. Click the needed patient information and a blue bar displays which indicates that the current item is selected.
- 2) Click **OK** to enter the Measuring Data Display Interface. If no item is selected, the dialogue box will indicate Nothing Selected. In the Measuring Data Display Interface, some function keys under the interface are invalid. The keys are grey and there is no response when clicking.
- 3) Under the Measuring Data Display Interface of ASCAN (see Figure 4.24), the data group can be selected and the measuring waveform be displayed by controlling the up and down keys. The Measuring Data Display Interface displays the measuring result saved initially but can not make amendment for the saved data and waveforms, and without measuring functions. The threshold value can be observed but not be amended. Under the Measuring Data Display Interface, it is able to enter the IOL Interface by clicking **IOL** to check the saved data or input the parameters for recalculation. The newly input parameters and calculated result can't be saved. If measurement or storage of new data is required, please return to the normal working interface.

To exit from the IOL Interface, it's required to click **ASCAN** key to enter the ASCAN Interface and click **BACK** key to return to the DATA BASE Interface; click **EXIT** key to exit and return to the normal working interface.

Under the Measuring Data Display Interface, the data can not be processed. If clearance or deletion is required, please exit from this mode and return to the DATA BASE Interface to clear or delete the selected information. Click **EXIT** key under the DATA BASE Interface to exit and return to the normal working interface.

4.3.5 Amendment and Storage of Parameters, and Clock Setting

Click **SETUP** key in the MENU Interface to enter the SETUP Interface, in which there are four tab pages.

4.3.5.1 Printing Parameter Setup for A Biometer

Click **A_PRINT** key to enter the A_PRINT tab page, as shown in Figure 4.25, which is the tab page for ASCAN to set up printing parameters. The operator can choose the information need to be printed out. The default setting is to print out all information, including: patient information, measuring information, IOL calculating parameters, calculating results and output waveforms.

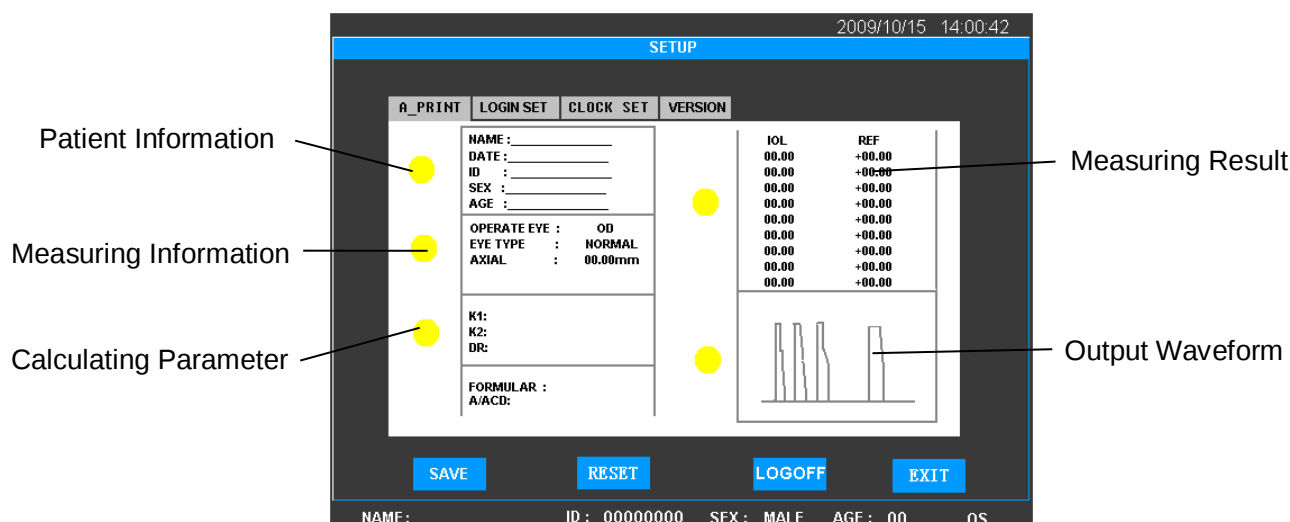


Figure 4.25 A_PRINT

Each indicator indicates the corresponding printing information on the right. Yellow indicates the selected status, while grey indicates the unselected status.

Click the position of the indicator to select, the yellow indicator lights up means that this part is selected to be printed out. Click again to turn off the indicator, which turns to grey and means that the selected part will not be printed out.

About the printing parameter setup of A Biometer, except for the patient information and measuring information, the printing status for other parts can be changed. (The calculating result synchronizes with the printing status of calculating parameter, that is to say, both items can be selected to print out or cancel printing synchronously.)

4.3.5.2 LOGIN SET Parameter Setup (Unavailable under ADMIN System)

Under startup selection, the default system is ADMIN, click **ENTER** key directly to enter. If ADMIN is selected, the system will enter the default setting, which means the gain of A Biometer, printing setup and IOL calculating parameter are all default settings. These values can be amended and saved after amendment.

If other user system (USER1, USER2, USER3, USER4 or USER5) will be selected, click system selection key to select the required user system, and then click **ENTER** key to enter. The gain of this system, printing setup and IOL calculating parameter can be set and used separately.

The name of the user system can be amended and saved in the SETUP Interface as shown in Figure 4.30. When login again, the amended name will be displayed in the LOGIN Interface.

Figure 4.26 LOGIN SET Parameter Setup

Complete all amendments mentioned in §4.3.5.1~§4.3.5.2 and select **SAVE** key to save the selected result. Click **RESET** key to recover the initial values. Click **EXIT** key to exit from the interface. Click **LOGOFF** key to exit from the system and the system can be re-selected after entering into the LOGIN Interface.



Note: The functions of **SAVE** and **RESET** will save all parameters in **A_PRINT** and **LOGIN SET** tab pages, or reset all settings to initial values. Please confirm before the operation.

4.3.5.3 Clock Set

Figure 4.27 CLOCK SET

If it is required to reset date and time, please click **CLOCK SET** key in this interface to enter the clock set tab page, as shown in Figure 4.27, where time is displayed in real-time. Click **CLOCK SET** key to change the clock setting. At that time, the clock stops, click **NEXT** key to move the cursor in the text box to the next position successively and circularly until reaching the right

position. Input the correct time with the keypad and click **SAVE** key to store. The amended value appears on the top right corner to display time. Click **RESET** key to restore to zero. Click **EXIT** key once to exit from the clock editing status; double click **EXIT** key to exit from the dialogue box.

4.3.5.4 VERSION

Click **VERSION** key to display the tab page of statement, which includes the software version and contact information of the company.

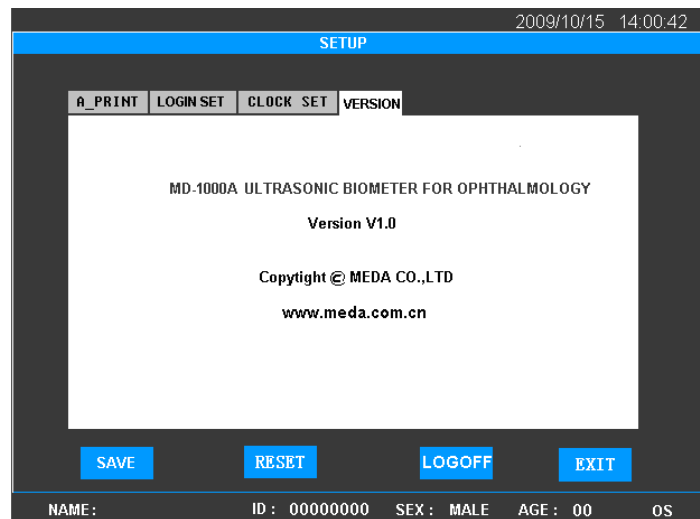


Figure 4.28 VERSION

Chapter 5 Cleaning and Disinfection

FOR EUROPE:

5.1 How to prevent Cross-Infection

Use soft cloth with water or neutral cleanser to clean the instrument surface. For further cleaning or disinfection, use soft cloth lightly dampened with the recommended cleanser or disinfectant solution to wipe the instrument surface. Remove the residual cleanser or disinfectant with clean soft cloth dampened in sterile water. Make sure the instrument is completely dry before using.

The probe and eye cup must be kept clean, before initial use and in between patient uses, the probe and eye cup should be cleaned and disinfected. The recommended procedures are as below. If other [cleansers](#) or disinfectants are used, make sure they are legally marketed and please follow their manufacturer's instructions.

5.2 Decontamination Procedure – Pre-disinfection and Disinfection of Probes

PREAMBLE

---- The operator should use standard method to ensure satisfactory decontamination - pre-disinfection and disinfection of the probe after use.

---- The operator should use method for risky-patients to ensure satisfactory decontamination - pre-disinfection and disinfection of the probe every time after use on a patient where there is a risk of infection of Creutzfeld-Jacob disease.

OPERATOR'S CLOTHING

- One-off overall.
- Disposable gloves, sterile for disinfection.
- Glasses and anti-rejection masks.

EQUIPMENT

- Soft silk brush (surgical nail brush)
- 3×500 ml stainless steel (or plastic), autoclavable-soaking trays.
- One-off hand cloths.
- Demineralized or distilled water.

PRODUCTS

- Cleaning-predisinfectant: Aniosyme ® P.L.A. (Company: ANIOS),
or predisinfectant: Alkazyme ® alcalin (Company: ALKAPHARM).

The products must be diluted at 0.5% with warm water (25 °C-30 °C) from the tap or distilled

water.

The contents of the tray must be changed every day.

- Disinfectant type Alkacide ® (Company ALKAPHARM).

The product must be diluted at 5% with distilled water.

The solution must be changed every day.

- 6 Chlorometric degree solution of sodium hypo chloride at 20 °C.

The contents of the tray must be changed after each use.

- Demineralized or distilled water.



Notes: 1) Please disconnect the probe from the instrument. Please be sure the instrument is TURNED OFF before disconnecting probes.

2) Avoid splashing liquids onto the probe connector (end of the cable, which is connected to the instrument).

5.3 Preparation of Decontamination Agent

Decontamination – Pre-disinfection

- Proteolytic enzyme based agents (2 possibilities)

0.5% Alkazyme solution in water (20g sachet)

- 1) Pour in 1L warm water (25-30°C)
- 2) Put in the unopened sachet.
- 3) Wait for 1 minute.
- 4) Pour in 4 L water and stir it.

The Alkazyme solution can be used within 8 days if kept in a sealed flask. The solution can also be made up in a 4L recipient using distilled water fill up the soaking tray from there.

OR:

0.5% Aniozyme solution in water (25g sachet):

- 1) Pour in 1L warm water (25-30°C)
- 2) Put into the unopened sachet.
- 3) Wait for 1 minute
- 4) Pour in 4L warm water and stir.

The Aniozyme solution lasts 1 day in a sealed flask.

Disinfection Agent

0.5% Alkacide solution in water:

- 1) Pour 5L distilled water in the flask.
- 2) Pour in the Alkacide
- 3) Stir it

The Alkacide solution can be used within 8 days if kept in sealed flask.

Please pour in soaking tray (500ml) when disinfection is necessary.

Replacing Contents of soaking trays

For frequent use, the contents of the trays should be replaced at the beginning of the morning and beginning of the afternoon.

Wait 10 minutes after the last decontamination before emptying out the Alkazyme or Aniozyme solutions.

5.4 Standard Method



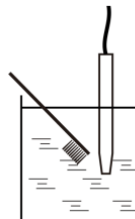
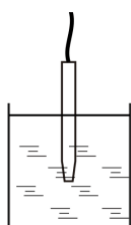
Notes: 1) Please disconnect the probe from the instrument. The instrument must be TURNED OFF first.

2) Avoid splashing any liquid onto the electrical connector.

A-Probe

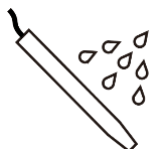
A) Decontamination-Pre-disinfection

1. Immerse the probe and the cable (except the connector) in the solution of Alkazyne or Aniozyme for 5 to 15 minutes depending on the perceived level of risk.
2. Clean the probe and the cable in the solution with the brush for 1 minute. Keep the connector dry.



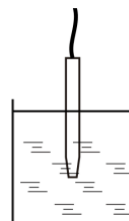
B) Rinsing

3. Rinse the probe and the cable in demineralized or distilled water. Do not wet the connector.



C) Disinfection

4. Immerse the probe and the cable in the Alkacide solution for 5 to 20 minutes depending on the estimated level of risk. Please keep the connector dry.



D) Rinsing

5. Rinse the probe and the cable with demineralized or distilled water. Keep the connector dry.

E) Drying

6. Dry it with a sterile compress or a single use dry wipe if the rinsing water was sterile.
7. Now the probe is ready for use.

METHOD FOR HIGH RISK PATIENTS

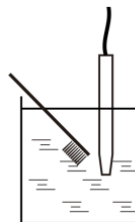
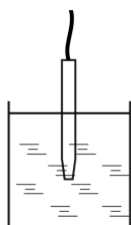


Notes: 1) Please disconnect the probe from the instrument. The instrument must be **TURNED OFF** first.

2) Avoid splashing any liquid onto the electrical connector.

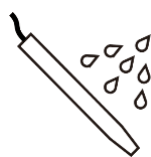
A) Decontamination-Pre-disinfection

1. Immerse the probe and the cable (except connector) in a solution of Alkazyme or Aniozyme for 5 to 15 minutes depending on the perceived level of risk.
2. Clean the probe and the cable in the Chosen solution for 1 minute using the brush. Keep the connector dry.



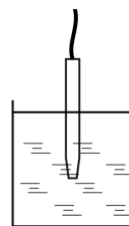
B) Rinsing

3. Rinse the probe and the cable with demineralized or distilled water. Please do not splash liquid onto the connector.



C) Inactivation

4. Immerse the probe and the cable (except connector) in a 6 chlorometric degree solution hypochloride for 60 min. at 20°C. Keep the connector dry.



D) Rinsing

5. Rinse the probe and the cable with demineralized or distilled water. Do not wet the connector.

E) Disinfection

6. Dry with a sterile compress or a single use dry wipe if the rinsing water was sterile.

F) Rinsing

7. Rinse the probe end with demineralized or distilled water keeping the connectors dry

G) Drying.

8. Dry with a sterile compress or a single use dry wipe if the rinsing water was sterile.
9. The probe is ready for use.

Reminder:

- For eye cup, please refer to the method of A-Probe.

FOR U.S.A

5.5 Cleaning of the Main Unit of EUS-1800A

Use soft cloth with water or neutral cleanser (e.g. diluted Bianex® Special Efficiency Multi-Enzymatic Cleanser) to clean the instrument surface. For further disinfection, please use soft cloth with disinfectant solution (e.g. Sporidicin Sterilizing and Disinfecting Solution) to wipe the instrument surface.

Remove the residual cleanser or disinfectant with clean soft cloth dampened in sterile water. Make sure the instrument is completely dry before using.



Note: 1) Do not use any corrosive cleanser and disinfectant when cleaning the instrument surface; keep water and any other liquids away from the instrument.

2) Do not immerse the instrument in disinfectant solution.

5.6 How to Prevent Cross-Infection

The probe and eye cup that are intended to come in contact with mucous membranes, belong to semi-critical device.

To prevent cross-infection, before initial use and in between patient use, the probe and eye cup must be thoroughly cleaned and subjected to a germicidal process with a broad spectrum of activity.



Note: It is prohibited to use the device on patients with eye trauma, inflammation or infection.

5.7 Method

Before initial use and in between patient use, the A-Probe and eye cup should be processed by high level disinfection. The recommended procedures are as below:

5.7.1 Preparation

a) OPERATOR'S CLOTHING

- Liquid-proof gowns.
- Disposable gloves.
- Eye protection and face masks.

b) EQUIPMENT

- Soft silk brush (surgical nail brush).
- Four glass containers larger than 500ml and capable of immersing the probe and eye cup.

—— One-off sterile cloth.

—— Sterile water.

c) PRODUCTS

—— Cleanser: ***Bianex® Special Efficiency Multi-Enzymatic Cleanser***

—— Disinfectant: ***Sporicidin Sterilizing and Disinfecting Solution (K003087)***

5.7.2 Thorough Cleaning

a) Prepare the cleaning solution by adding 1-2ml Bianex® Special Efficiency Multi-Enzymatic Cleanser into 500ml of sterile water. Additional amounts maybe required for hard-to-remove matter. The water temperature should be from 20°C to 64°C.

b) Immerse the A-Probe and/or eye cup that need to be disinfected into the container with 500ml cleaning solution for 5 minutes (Do not immerse the probe connector). Clean the A-Probe and/or eye cup carefully in the cleaning solution with a soft silk brush and shake it to wash; and then take it out and wipe off the cleaning solution with sterile cloth.

c) Rinse the cleaned A-Probe and/or eye cup by immersing them in 500ml sterile water for 1 minute.

d) Drain the A-Probe and/or eye cup and wipe it with sterile cloth.



Note: 1) Disposable gloves, eye protection and liquid-proof gowns should be worn when cleaning soiled probe and/or eye up.

2) All detail information for Bianex® Special Efficiency Multi-Enzymatic Cleanser, please refer to the manufacturer's instructions.

5.7.3 Disinfection

Use disinfectant solution immersion method:

a) Activate and prepare the disinfectant solution (Sporicidin Sterilizing and Disinfecting Solution) according to the manufacturer's instructions.

b) Sporicidin Indicators should be used before each use of this solution to determine whether glutaraldehyde is at or above its minimum concentration of 0.6% and phenol is at or above its minimum recommended concentration of 1.3%. Do not use beyond 14 days even if the respective concentration of active ingredients is above the minimum recommended concentration as indicated by the indicator strips.

c) Disinfect the A-Probe and/or eye cup by immersing them completely in about 500ml of disinfectant solution for a minimum of 20 minutes at 25°C. Please keep the probe connector dry during disinfection.

d) Drain it and wipe off the disinfectant solution with sterile cloth.



- Note:**
- 1) Disposable gloves, eye protection, face masks and liquid-proof gowns should be worn when disinfecting soiled probe and/or eye up.**
 - 2) Containers filled with disinfectant solution should be kept closed and stored in a well ventilated, low traffic area at room temperature.**
 - 3) All detail information for Sporidicin Sterilizing and Disinfecting Solution, please refer to the manufacturer's instructions.**

5.7.4 Rinsing after Disinfection

- a) Thoroughly rinse the disinfected probe and/or eye cup by immersing them in three separate copious volumes of 500ml sterile water. Each rinse should be a minimum of 1 minute in duration. Use fresh portions of water for each rinse.
- b) Take out the A-Probe and/or eye cup, and let it dry naturally.
- c) After disinfection processing, put the A-Probe and/or eye cup into a sterile box. Now the A-Probe and/or eye cup are ready for use.



Note: 1) Steam sterilization is prohibited.

- 2) Disconnect the probe from EUS-1800A before disinfection; the power of EUS-1800A should be turned off before disconnecting the probe, hold the probe cable connector, do not pull the probe cable; While connecting the probe, make sure the red mark on the probe align with the red mark on the socket. See Figure 3.4.**
- 3) Do not immerse probe cable connector into water or other liquid; keep the probe cable connector dry throughout the disinfection process of probe.**
- 4) After disinfection, check carefully whether the surface of the probe is deformed or damaged and whether the probe cable is damaged; check carefully whether the eye cup is deformed and whether the surface smoothness is reduced and so on.**

In case of any abnormality, stop using and contact the manufacturer or your local distributor for service.

Chapter 6 Labeling

6.1 External Label

- 1) Printing Paper Packaging Label

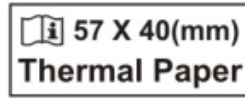


Figure 6.1 Printing Paper Packaging Label

- 2) Probe Socket Label



Figure 6.2 Probe Socket Label

6.2 Internal ID

- 1) Button cell: CR2032/3V, lithium cell



Figure 6.3 ID of Button Cell inside the Main Unit

Chapter 7 Maintenance, Attentions and Simple Defects Treatment

7.1 Maintenance and Attentions to Instrument

- 1) The instrument should be operated in a clean environment. Air-conditioned environment is recommended.
- 2) The instrument should be placed on a stable worktable or platform. Avoid direct sunlight.
- 3) Please use the supplied Power Adaptor which is in accordance with the safety standard of medical electric equipment; do not use other adaptors or adaptors of other equipment with the instrument.
- 4) Although anti-interference measures which are in accordance with IEC 60601-1-2 have been adopted, the instrument should be placed to avoid strong electromagnetic radiation equipment (such as microwave, radio frequency therapy equipment, etc.)
- 5) Routine inspection and maintenance should be carried out only if the mains power is switched off. None corrosive detergent is allowed to clean the housing. Avoid water and liquid flowing into the housing. Only a mild detergent may be used with soft tissue cloth.
- 6) In humid area and/or season, if the instrument is not used for a long time, it should be power-on for one hour per month to get the damp out.
- 7) Avoid drop or severe shock when moving the instrument. Give particular attention to protect the probe.
- 8) Do not block the ventilation window of the instrument. When the temperature of the instrument is not normal, please contact the manufacturer for service.
- 9) The instrument should always be placed in a secure location to prevent probe falling damages.
- 10) The instrument has no special protective measures for discharge effect of cardiac defibrillators; it is not suitable for use with high-frequency surgical equipment

7.2 Maintenance and Attentions to Probe

- 1) Do not autoclave the probes.
- 2) Don't wind the probe cable in coil less than 9 cm (3.5 inches); Check the probe cable regularly and stop using immediately if it is damaged or broken, contact the manufacturer or the local distributor for service.
- 3) Probe should be handled gently to avoid collision and drop, in order to prevent break and damage.
- 4) Avoid drop or scratch of the surface of probe when the probe is used or moved; make sure the probe is removed from the main unit and put into the packing bag during transportation.
- 5) If the probe drops during using or moving, check the top and the housing of probe carefully, and then check if it works well. Stop using if there is any problem and contact the

manufacturer or local distributor for service.

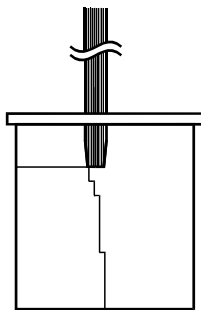
- 6) The whole probe cable including the cable plug is prohibited to be immersed into water or other liquid.
- 7) The connection and/or disconnection of probe should be done only if the system is powered off. While plugging in the probe, make sure the red mark on the probe align with that on the socket. Hold the probe plug and don't plug probe cable while plugging off the probe.
- 8) If the dialogue box on the screen prompts "PLEASE CHECK THE PROBE!", it indicates the probe connection is abnormal. Click "OK" to close the dialogue box and check if the probe is well connected.
- 9) The probes provided with the instrument should be used in the instrument only and not be used for other purposes.

7.3 Maintenance of LCD Screen

- 1) Clean the LCD screen with glasses cloth, lens tissue or other soft material.
- 2) Don't touch the LCD screen with hard object, which will cause irreparable damage to the screen.
- 3) Avoid strong shock and vibration during transportation.
- 4) Complete the touch-screen operation with finger belly.

7.4 Check before Use

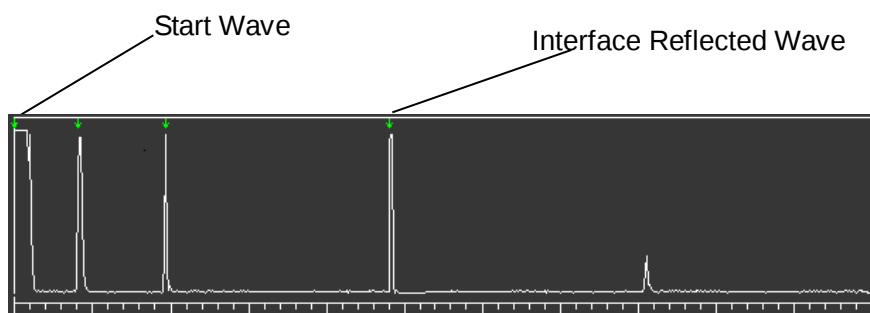
Biometric Measuring function



There is a test tank with each EUS-1800A which imitates four acoustical reflect interfaces of human eyes and used to test the biometric measuring.

Fill the tank with distilled water. Be sure that there is no air bubbles in the water. Operate according to automatic biometry. Gently put the A probe onto the highest stage perpendicularly into the tank (see left drawing).

Click **SCAN/FREEZE** key under the ASCAN Interface, the automatic measuring result to step target will be achieved, as shown below:



Keep the probe unmovable; click **SCAN/FREEZE** key to repeat measuring. If the results show good repeatability, it indicates the biometry mode of the equipment works well.

7.5 Trouble Shooting

- 1) In case of connection failure, please check first if the power supply and probe are well connected.
- 2) In case of printing failure, please check first if it is out of printing paper or paper jam.
- 3) In case of clock failure which can not be restored by restarting, possibly, the clock battery is used up. The battery can't be replaced by operator, please contact your distributor for repair.
- 4) In case of display or operation failure, please switch off the power and restart, check if it is back to normal.
- 5) In case of overheat or other abnormalities, please switch off the mains power immediately to prevent danger.
- 6) The over-current protection fuse is sealed inside the power adaptor and can't be replaced by user. If the power adaptor has no DC output, please pull out the main power plug and contact the manufacturer for repair.

If above operations are not effective, please don't open the housing without authorization. Contact your local distributor immediately. Explain the problems in detail for proper and in time support.

If required, we can provide the complete maintenance and repair manual to the authorized qualified engineers of service stations.

The Biometer is a high-tech product designed and built with high level of precision. Only qualified trained engineers are authorized to repair the instrument.

We are not responsible for problems caused by any kind of unauthorized repair.

7.6 Disposal after Product Life-Cycle

Disposal of the product after life-cycle should comply with the relevant local environment protection regulations. Same way can be taken with the electronic devices (computers, etc).

Chapter 8 Service and Support Information

8.1 Warranty

- 1) The product has a warranty of one year from the date of purchasing, on the premise of using in accordance with the User Manual.
- 2) If the device does not work properly, please contact your local distributor or the manufacturer immediately.
- 3) Following repairs will be charged within warranty period:
 - Problems caused by man-made damages;
 - Damages caused by unauthorized repair;
 - Damages caused by inappropriate operation.
- 4) We provide continuous maintenance and repair after warranty period with certain charges.

8.2 Accessories and Materials

8.2.1 Consumables

SN	Description	Manufacturer	Model	Specification
1	Printing Paper	Beijing Xun Pu Electronic Technology Company	—	57mm×40mm Thermal Printing Paper

8.2.2 Detachable Parts

SN	Description	Model	Specification
1	A-Probe	Prb1000A/10-C	10MHz
2	Power Adaptor	HES49-12040	12V/4A
3	Foot Switch	EUS-1800A.FS-2	—

8.2.3 Materials

Material of Housing	Model
Flame Retardant ABS Engineering Plastic (not less than FV-2)	MCN-6000



Note: 1) The probe, power adaptor and foot switch used in the system must be provided by the Manufacturer.

2) The consumables used in the system must be product (with specification) designated by the Manufacturer.

8.3 Software Update and Services

The Manufacturer will provide users with software update information and services.

In case of software update due to correction of defects, the Manufacturer will provide users with

free software update service.

For the expansion of software functions, the manufacturer will provide users with software update information, and according to users' demand, provide software update service with certain charges.

Annex A Prudent Use Statement

A.1 Statement

Although there is no evidence so far the diagnostic ultrasound instrument will lead to biological effects in humans, it is possible to prove the existence of biological effects in future applications. The radiated acoustic power of the instrument itself is very low; still we must use ultrasound prudently in clinical applications. Do the best to complete patients' examinations within shortest time and with lowest power, on the premise of achieving necessary clinical information.

A.2 Principle of ALARA (As Low As Reasonably Achievable)

The principle of ALARA should be implemented to perform ultrasound procedure. Try to use lowest level of energy that will not result in biological effect. The ultrasonic energy depends on acoustic output intensity and exposure time. The ultrasonic intensity required may vary depending on different patients and clinical cases.

Not all examinations can be accomplished with lowest ultrasonic energy output. Lowest ultrasonic energy will produce low-quality image or weak Doppler signal which will influence the reliability of diagnosis. However the acoustic power higher than practical use will not be helpful to improve the quality of diagnostic information, but will increase the risk of biological effect.

The operator must be responsible for the safety of patients and use ultrasound on purpose, that is to apply ultrasonic output power according to the principle of ALARA.

Annex B Prompt Message

1. **System start-up prompt:** Switch on the system, it will read parameter and self-check, displaying "THE SYSTEM IS CHECKING ...PLEASE WAIT...". After checking, it will enter the normal interface.
2. **Probe checking prompt:** If probe is not connected, the system will prompt automatically when scanning. The dialogue box will display "PLEASE CHECK THE PROBE!" reminding you to check the probe connection.
3. **Nothing Selected prompt:** If the operation is wrong and unable to select the correct item, the dialogue box will prompt "PLEASE SELECT AN ITEM!" or "NO DATA!", indicating that no correct item is selected.
4. **Confirming prompt for deleting or saving:** When deleting some item or saving the result of a patient's case, the dialogue box will pop up. "ARE YOU SURE TO SAVE?" prompts to confirm saving or not. "ARE YOU SURE TO DELETE?" prompts to confirm deletion or not. "ARE YOU SURE TO CLEAR ALL THE DATA PERMANENTLY?" prompts the cleared data are not recoverable and whether to continue the clearance. "ARE YOU SURE TO DELETE THE ITEM PERMANENTLY?" prompts the deleted data are not recoverable and whether to continue the deletion.
5. **Whistle prompt:**
 - a) Three short beeps of buzzer "beep - beep - beep" prompt measurement failed.

For A Biometer, please check if measurement is performed for eight groups of data.
 - b) A long beep of buzzer "beep-" prompts measurement completed.

Under automatic measuring status for A Biometer, it prompts the measurement for eight groups of data have been completed.
 - c) A short beep of buzzer "beep" prompts single measurement completed.

Under automatic measuring status for A Biometer, there will be a short beep each time when proper data have been acquired, then the data will display in the corresponding group of cursor.

Annex C.1 Acoustic Output Report (IEC60601-2-37)

Index		MI	TIS	TIS	TIS	TIB	TIB	TIC
Mode		-	Scanning	Non-scanning	Non-scanning	Scanning	Non-scanning	-
				$A_{\text{aprt}} \leq 1\text{cm}^2$	$A_{\text{aprt}} > 1\text{cm}^2$			
Global Maximum Index Value		0.12	-	0.08	-	-	-	-
Associated acoustic parameter	Acoustic working frequency (f_{awf})	9.27	-	9.27	-	-	-	
	Bounded output power (P_1)		-			-		
	Attenuated output power (P_{α})			1.86	-		-	-
	Attenuated spatial-peak temporal-average intensity ($I_{\text{spta}, \alpha}$)				-		-	
	Attenuated peak-rarefactional acoustic pressure ($p_{r, \alpha}$)	0.24						
	-12dB output beam area (A_{aprt})		-		-	-	-	-
	Break-point depth (z_{bp})				-			
	Equivalent aperture diameter (D_{eq})							-
	Depth for TIS (z_s)				-			
	Depth for TIB(z_b)=depth at max [$I_{\text{pi}, \alpha} \times P_{\alpha}$]						-	
	Depth at max. attenuated pulse-intensity integral (z at max. $I_{\text{pi}, \alpha}$)	3.16						
Other information	Pulse duration (t_d)	0.14						
	P_r at max I_{pi}	0.67						
	$I_{\text{pi}, \alpha}$ at max MI	2.71						
Supplementary information: Operating accessory: A probe Prb1000A/10-C Frequency: 10MHz								

Annex C.2 Acoustic Output Reporting Table

Acoustic Output Reporting Table for Track1 Non-Autoscanning Mode

System: EUS-1800A Ultrasonic Biometer for Ophthalmology

Transducer Model: Prb1000 A/10-C

Operating Mode: A-Mode

Application: Ophthalmology

Working Frequency: 10MHz

Acoustic Output			MI	I _{SPTA.3} (mW/cm ²)	I _{SPPA.3} (W/cm ²)
Maximum index value			0.079	0.003	1.68
Associated acoustic parameter	P _{r.3} (MPa)		0.241		
	W ₀ (Mw)			0.00035	0.00035
	f _c (MHz)		9.203	9.203	9.203
	Z _{SP} (cm)		2.10	2.10	2.10
	Beam dimensions	x-6 (cm)		0.169	0.169
		y-6 (cm)		0.162	0.162
	PD (usec)		0.202		0.202
	PRF (Hz)				10
	EDB	Az (cm)		Φ 0.48	
		Ele (cm)		Φ 0.48	
Operating Control Conditions	---			---	
	---			---	
	---			---	
	---			---	

Transducer Model:		Prb1000A/10-C		
Operation Mode:		A Mode		
Application:		Ophthalmology		
Working Frequency:		10MHz		
Acoustic Output		MI	I _{SPAT.3} (mW/cm ²)	I _{SPPA.3} (W/cm ²)
Sample Size		3	3	3
K (P=0.90, gamma=0.9)		4.258	4.258	4.258
Mean	(x)	0.071	0.003	1.613
StdDev	(Sx)	0.007	0.000	0.335
Limit	(X)	0.100	0.005	3.040

Annex D Guidance and Manufacturer's Declaration

Guidance and manufacturer's declaration – electromagnetic emissions- for all EQUIPMENT and SYSTEMS

Guidance and manufacturer's declaration – electromagnetic emission		
The <i>MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY</i> is intended for use in the electromagnetic environment specified below. The customer of the user of the <i>MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY</i> should assure that it is used in such and environment.		
Emission test	Compliance	Electromagnetic environment – guidance
RF emissions CISPR 11	Group 1	The <i>MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY</i> uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emission CISPR 11	Class A	The <i>MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY</i> is suitable for use in all establishments, other than domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	

**Guidance and manufacture's declaration – electromagnetic immunity –
for all EQUIPMENT and SYSTEMS**

Guidance and manufacture's declaration – electromagnetic immunity			
The MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY is intended for use in the electromagnetic environment specified below. The customer or the user of MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY should assure that it is used in such an environment.			
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
Electrostatic discharge (ESD) IEC 61000-4-2	±6 kV contact ±8 kV air	±6 kV contact ±8 kV air	Floors should be wood, concrete or ceramic tile. If floor are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast transient/burst IEC 61000-4-4	±2 kV for power supply lines	±2 kV for power supply lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	±1 kV differential mode ±2 kV common mode	±1 kV differential mode ±2 kV common mode	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5% U_T (>95% dip in U_T) for 0.5 cycle 40% U_T (60% dip in U_T) for 5 cycles 70% U_T (30% dip in U_T) for 25 cycles <5% U_T (>95% dip in U_T) for 5 sec	<5% U_T (>95% dip in U_T) for 0.5 cycle 40% U_T (60% dip in U_T) for 5 cycles 70% U_T (30% dip in U_T) for 25 cycles <5% U_T (>95% dip in U_T) for 5 sec	Mains power quality should be that of a typical commercial or hospital environment. If the user of the MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY requires continued operation during power mains dip & interruptions, it is recommended that the MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY be powered from an uninterruptible power supply or a battery.
Power frequency (50Hz) magnetic field IEC61000-4-8	3A/m	3A/m	Power frequency magnetic fields Should be at levels characteristic of a typical location in a typical commercial or hospital environment.
NOTE U_T is the a.c. mains voltage prior to application of the test level.			

**Guidance and manufacturer's declaration – electromagnetic immunity –
for EQUIPMENT and SYSTEMS that are not LIFE-SUPPORTING**

Guidance and manufacturer's declaration – electromagnetic immunity			
The MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY is intended for use in the electromagnetic environment specified below. The customer or the user of MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY should assure that it is used in such an environment.			
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
Conducted RF IEC 61000-4-6	3 V _{rms} 150 kHz to 80 MHz	3 V _{rms}	Portable and mobile RF communications equipment should be used no closer to any part of the MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance $d = \left[\frac{3.5}{V_1} \right] \sqrt{P}$ $d = \left[\frac{3.5}{E_1} \right] \sqrt{P} \qquad 80 \text{ MHz to } 800 \text{ MHz}$ $d = \left[\frac{7}{E_1} \right] \sqrt{P} \qquad 800 \text{ MHz to } 2.5 \text{ GHz}$ Where <i>P</i> is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and <i>d</i> is the recommended separation distance in metres (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, ^a should be less than the compliance level in each frequency range. ^b Interference may occur in the vicinity of equipment marked with the following symbol: 
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	
NOTE 1 At 80 MHz and 800 MHz, the higher frequency range applies. NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.			
^a Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY is used exceeds the applicable RF compliance level above, the MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY.			
^b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.			

**Recommended separation distances between portable and mobile
RF communications equipment and the EQUIPMENT or SYSTEM –
for EQUIPMENT or SYSTEM that are not LIFE-SUPPORTING**

Recommended separation distances between portable and mobile RF communications equipment and the <i>MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY</i>			
The <i>MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY</i> is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the <i>MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY</i> can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the <i>MD1000A ULTRASONIC BIOMETER FOR OPHTHALMOLOGY</i> as recommended below, according to the maximum output power of the communications equipment.			
Rated maximum output power of transmitter (W)	Separation distance according to frequency of transmitter (m)		
	150 kHz to 80 MHz $d = \left[\frac{3.5}{V_1} \right] \sqrt{P}$	80 MHz to 800 MHz $d = \left[\frac{3.5}{E_1} \right] \sqrt{P}$	800 MHz to 2.5 GHz $d = \left[\frac{7}{E_1} \right] \sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.37	0.37	0.74
1	1.17	1.17	2.33
10	3.69	3.69	7.38
100	11.67	11.67	23.33
For transmitters rated at a maximum output power not listed above, the recommended separation distance d in metres (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.			
NOTE 1	At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.		
NOTE 2	These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.		

Annex E IOL Formula

Six formulae are used in the EUS-1800A, they are:

SRK-II

SRK-T

BINKHOST-II

HOLLADAY

HOFFER-Q

HAIGIS

IOL calculation formulas after corneal refractive surgery

HISTORY DERIVED METHOD — To use this method, all historical data are required, including K1 and K2 values before corneal refractive surgery and the lens power of spectacles before and after corneal refractive surgery, that is to input all parameters of K1pre, K2pre, REFRpre and REFRpost in the dialogue box.

DOUBLE-K/SRK-T METHOD — To use this method, all historical data are required, including K1 and K2 values before corneal refractive surgery and the lens power of spectacles before and after corneal refractive surgery, that is to input all parameters of K1pre, K2pre, REFRpre and REFRpost in the dialogue box.

REFRACTION DERIVED METHOD — To use this method, some historical data are required, including the current K1 and K2 values and the lens power of spectacles before and after corneal refractive surgery, that is to input the parameters of K1, K2, REFRpre and REFRpost in the dialogue box.

ROSA METHOD — To use this method, the current K1 and K2 values of the patients are required, that is to input the parameters of K1 and K2 in the dialogue box.

SHAMMAS METHOD — To use this method, the current K1 and K2 values of the patients are required, that is to input the parameters of K1 and K2 in the dialogue box.

See below for the selection of formula and parameter:

Refractive Surgery ?	Formula	Parameter									
		AC	K1	K2	A	ACD	ANTI/POST	K1pre	K2pre	REFR pre	REFR post
NO	SRK_II	x	√	√	√	x	x	x	x	x	x
	SRK_T	x	√	√	√	x	x	x	x	x	x
	BINK_II	x	√	√	x	√	√	x	x	x	x
	HOFFER_Q	x	√	√	x	√	x	x	x	x	x
	HOLLADAY	x	√	√	√	x	x	x	x	x	x
	HAIGIS	√	√	√	√	x	x	x	x	x	x
YES	History Derived	SRK_II	x	√	√	√	x	x	√	√	√
		SRK_T	x	√	√	√	x	x	√	√	√
		BINK_II	x	√	√	x	√	√	√	√	√
		HOFFER_Q	x	√	√	x	√	x	√	√	√
		HOLLADAY	x	√	√	√	x	x	√	√	√
		HAIGIS	√	√	√	√	x	x	√	√	√
	DoubleK	DoubleK/SRKT	x	x	x	√	x	x	√	√	√
	Refraction Derived	SRK_II	x	√	√	√	x	x	x	√	√
		SRK_T	x	√	√	√	x	x	x	√	√
		BINK_II	x	√	√	x	√	√	x	√	√
		HOFFER_Q	x	√	√	x	√	x	x	√	√
		HOLLADAY	x	√	√	√	x	x	x	√	√
		HAIGIS	√	√	√	√	x	x	x	√	√
	ROSA	ROSA	x	√	√	√	x	x	x	x	x
	SHAMMAS	SHAMMAS	x	√	√	√	x	x	x	x	x

Remarks:

1. “√” means the parameter is applicable to this formula and “x” means not applicable. The shaded area in the table is applicable to cases without refractive surgery.
2. The parameters of AL and DR are required for each formula; they are not listed in this table.
3. K1 and K2 are the current keratometry; K1pre and K2pre are keratometry before corneal refractive surgery; REFRpre is the refraction before corneal refractive surgery (the amount of lens power of spectacles); and REFRpost is the refraction after corneal refractive surgery (the amount of lens power of spectacles).

Annex F REFERENCES

HAIGIS W: Biometrie, in: Augenärztliche Untersuchungsmethoden, Straub W, Kroll P, Kühle HJ (Hrsg), F.Enke Verlag Stuttgart, 255-304, 1995

RETZLAFF J: A new intraocular lens calculation formula, Am Intra-Ocular Implant Soc J 6:148-152, 1980

RETZLAFF J, SANDERS DR, KRAFF MC: Development of the SRK/T intraocular lens implant power calculation formula. J Cataract Refract Surg 16 (3):333-340, 1990

SANDERS DR, KRAFF MC: Improvement of intraocular lens power calculation using empirical data, Am Intra-Ocular Implant Soc J 6: 263-267, 1980

SANDERS DR, RETZLAFF J, KRAFF MC: Comparison of the SRK II formula and other second generation formulas. J Cataract Refract Surg 14: 136-141, 1988

HOFFER KJ: The effect of axial length on posterior chamber lens and posterior capsule position. Current Concepts Ophthalmic Surg, 1:20-22, 1984

HOLLADAY JT, PRAGER TC, CHANDLER TY, MUSGROVE KH, LEWIS JW, RUIZ RS: A three-part system for refining intraocular lens power calculations. J Cataract Refract Surg, 14:17-24, 1988

Intraocular lens power calculation after corneal refractive surgery: Double-K method. Jaime Aramberri, MD , J Cataract Refract Surg 2003; 29:2063–2068 2003 ASCRS and ESCRS INTRAOCULAR LENS POWER CALCULATION AFTER CORNEAL PHOTOREFRACTIVE SURGERY; LITERATURE. MESA-GUTIÉRREZ JC1, RUIZ-LAPUENTE C2, ARCH SOC ESP OFTALMOL 2009; 84: 283-292.

EUS-1800 A BIOMETER



9990 NW 14 Street | Suite 105 | Doral, FL 33172
Telephone: 888.334.4640