



New optical system, unique imaging impression
Hartman imaging analyzing and processing technology, accurate measurement result
TFT touch screen, can move front and back freely
Motorized chinrest
Manual focusing, auto measuring
Vertex Distance(VD): 0.0, 12.0, 13.75, 15.0
SPH: -20.00D~+20.00D (VD=12mm, 0.01, 0.06, 0.12, 0.25 Unit)
CYL: 0.00D~±6.00D (0.12, 0.25 Unit)
Axis(AX): 1°~180° (1° Unit)
Cylinder Form: -, +, ±
Puiple Distance(PD): 10~85mm
Minimum Pupil Diameter: 2.0mm
Measuring Time: < 0.5s
Pupil Diameter: 2.00–8.00mm
Measuring Light Energy: <30uw (Insure measuring safety)
Radius of Curvature: 5.0~10.0mm (0.01mm Unit)
Corneal Power: 33.00D~67.00D
(In case that the corneal equivalent refractive power is 1.3375)
Corneal Astigmatism: 0.00D~-15.00D (0.06D/0.12D/0.25D Unit)
Data Storing: Each 10 measured values of left eyes and right eyes
Axis: 1° ~180°
Chart: Auto fog
Monitor: 7" TFT LCD touch screen (Angle of view adjustable)

Built-in Printer: 57mm thermal printer

TECHNICAL INDEXES

Electrical Power: AC 100~250V, 50/60Hz

N.W.: 16kg

G.W.: 19.5kg

Dimensions(packing): (L)650mm X (W)400mm X (H)620mm

## AR (K) 7680 Auto Refrac(Kera)tometer



### Do what you want to do, save what you hope in time and money

# **AR(K)7680** Auto Refrac(Kera)tometer

## The most cost-effective Hartmann refrac(kera)tometer in the world



#### Adjustable LCD Touch Screen



#### Motorized Chin Rest





REF Refractometry - R/K Refrackeratometry

Intuitive icons provide the user an easier operating circumstances, and make the measurement become more convenient and the data to be measured more accurate and fast

#### Hartmann imaging processing technology



Hartmannaram





3 groups of data stored each measurement, maximum 10 groups of data can be stored

#### Professional Design

#### Up/down Auto Tracking





Measure peripheral keratometry precision of eyes with contact lenses fitting





## Auto Refracto(Kera)tometer User's Manual



Please be sure to read this manual carefully before using the instrument and keep it handy for ready reference.

Thanks for your choice and use of this instrument. AR(K)7680 Auto RefracKeratometer is one high precision instrument of objective measuring the patient's eyes with unique optical system inside and accurate imaging analyzing and processing in Hartman technology. It's mainly used to measure the patient's diopter, including sphere power, cylinder power, optical axis, pupil distance and corneal curvature, to provide reference datas for eyes' treating and eyeglasses choice. The measurement result can be displayed on screen or printed out on paper, and can also be transferred to auto phoroptor (fit to RS232 interface)). If the auto phoroptor can output data, this instrumentAR(K)7680 can directly print out the measured optometry data by auto photoptor.

#### **Model No./Specifications**

Item No.	Spec.(L x W x H)	Input Power	Display	Remarks
	(mm x mm x mm)	(VA)		
AR7680	750×400×630	60	TFT Color	Wavefront
			LCD	Aberration(Manual)
ARK7680	750×400×630	60	TFT Color	Wavefront Aberration
			LCD	Corneal
				Curvature(Manual)

#### **Refractometer Named**



manual/auto

universal model

product serial number

objective refractometer(K optional)

#### **Refractometer Division Description**

The serial refractometers consist of optical system, mechanical transmission system, COMS image sensing system, microcomputer control system, and printer etc., are the professional instruments of providing reference data for glasses and eye diagnosis and treatment. According to measurement function, it's divided into refractive parameters measurement and refractive parameters measurement/corneal curvature measurement. According to measurement mode, it's divided into manual measurement and automatic measurement.

#### DISCLAIMER

1. This manual has been carefully checked to insure the contents' accuracy and perfect during compiling, however, for possible errors or omissions contained herein.

- 2. The company reserves the right to make changes to this product or the specifications at any time without prior notice.
- 3. The company own the final interpretation to this manual.

#### CONTENTS

1. ELECTROMAGNETIC COMPATIBILITY GUIDE AND MANUFACT	URER
STATEMENT	3
2. SAFETY PRECAUTIONS	8
3. UNPACKING AND INSTALLATION	10
4. FUNCTIONS OF THE MAJOR COMPONENTS	11
5. MAIN TECHNICAL INDEXES	13
6. ENVIRONMENT TERMS	14
7. LCD SCREEN DISPLAY	15
8. MENU	16
9. MEASUREMENT	23
10. COMMON TROUBLE SHOOTING	26
11. PACKAGING, TRANSPORTATION, STORAGE	27
12. ENVIRONMENTAL PROTECTION	27
13. ACCESSORIES	28

## 1. ELECTROMAGNETIC COMPATIBILITY GUIDE AND MANUFACTUTER STATEMENT

This product is in compliance with the electromagnetic compatibility regulations in this manual. To ensure compliance with these regulations, the user needs to install and use the information provided in this manual. Such as the use of non - manufacturers to provide the cable may cause the increase or decrease in the immunity of the product launch.

#### Warning!

- 1. The use of non manufacturer supplied cables may cause an increase in the electromagnetic radiation of this product or decrease the immunity.
- 2. Portable or mobile radio frequency communication equipment should not be used closer to any part of the refractometer than the recommended isolation by distance, including the cable.
- 3. In addition to the transducer and cable in sale as spare parts of components from the original equipment or system manufacturer, the use of other accessories, transducers and cables may cause an increase in the device or system to launch or decrease in immunity.
- 4. The device or systems should not be close to or stacked up with other devices, and if you have to approach or stack, it should be observed to verify the normal operation of its use.
- 5. The other accessories, transducer or cable to be used together with the device and system, it may cause an increase in the device or system to launch or decrease in immunity.

#### 1.1 Electromagnetic Emission Guide and Manufacturer Statement(Form1)

Guide and manufacturer's statement—Electromagnetic emission

[**Prototype ARK7680**] expected to be used in the electromagnetic environment of the following requirements, buyers and users should ensure that it is used in this electromagnetic environment

Launching Test	Conformity	Electromagnetic Environment—Guide
Radio frequency emission CISPR11	Group 1	[ <b>Prototype ARK7680</b> ] Radio frequency energy to be used for internal function only. Therefore, its RF emission very low, and the possibility of interference in the electronic device is very small.
Radio frequency emission CISPR11	Class B	[ <b>Prototype ARK7680</b> ] Applicable for all of the facilities in use, including the
Harmonic emission	Not	home and the direct connection of
IEC61000-3-2	applicable	residential public low voltage power
Voltage fluctuation/Flicker emission IEC61000-3-3	Not applicable	supply network.

## **1.2 Electromagnetic Immunity Guide and Manufacturer Statement** (Form2)

Guide a	and manufacturer's	s statement—Ele	ctromagnetic immunity
[ <b>Prototy</b> environment it is used in t	<b>pe ARK7680</b> ] ex of the following re his electromagnetic	pected to be us equirements, buyer e environment	sed in the electromagnetic s and users should ensure that
Immunity Test	IEC60601 Test Level	Meet Level	Electromagnetic Environment—Guide
Electrostatic discharge	±6kV contact discharge	±6kV contact discharge	The ground should be wood, concrete or ceramic tile, if the ground is covered with
IEC61000-4 -2	±8kV air dischange	±8kV air dischange	synthetic material, the relative humidity should be at least 30%

Electric fast transient pulse group IEC61000-4 -4	±2kV power line ±1kV input/output line	±2kV power line	Network power supply should have a typical commercial or hospital environment in the use of quality
Surge IEC61000-4 -5	±1kV line to line ±2kV line to ground	±1kV line to line ±2kV line to ground	Network power supply should have a typical commercial or hospital environment in the use of quality
Power input line voltage dips, short interruptions and voltage variations IEC61000-4 -11	<5% $U_t$ , last 0.5 cycle (Above $U_t$ , >95% sag) 40% $U_t$ , last 5 cycle (Above $U_t$ , 60% sag) 70% $U_t$ , last 25 cycle (Above $U_t$ , 30% sag) <5% $U_t$ , last 5s (Above $U_t$ , >95% sag)	<5% $U_t$ , last 0.5 cycle (Above $U_t$ , >95% sag) 40% $U_t$ , last 5 cycle (Above $U_t$ , 60% sag) 70% $U_t$ , last 25 cycle (Above $U_t$ , 30% sag) <5% $U_t$ , last 5s (Above $U_t$ , >95% sag)	Network power supply should have a typical commercial or hospital environment in the use of quality. If the users need [Prototype ARK7680] to continuously run during power supply interruption, then it's recommended the [Prototype ARK7680] is powered by a constant power supply or battery
Power frequency magnetic field (50Hz) IEC61000-4 -8	3A/m	3A/m	The power frequency magnetic field should have the characteristics of the power frequency magnetic level in a typical commercial or hospital environment
Note: Ut refe	ers to the AC networ	k voltage before app	lying the test voltage.

## **1.3 Electromagnetic Immunity Guide and Manufacturer Statement** (Form3)

Guide and manufacturer's statement—Electromagnetic immunity

[**Prototype ARK7680**] expected to be used in the electromagnetic environment of the following requirements, buyers and users should ensure that it is used in this electromagnetic environment

Immunity Test	IEC60601 Test Level	Meet Level	Electromagnetic Environment—Guide
Radio frequency transmission IEC61000-4-6 Radio frequency radiation IEC61000-4-3	3 V (effective value) 150 kHz ~ 80 MHz 3 V/m 80 MHz ~ 2.5 GHz	3V (effective value) 3 V/m	Portable or mobile radio frequency communication equipment should not be used closer to any part of [ <b>Prototype</b> <b>ARK7680</b> ] refractometer than the recommended isolation by distance, including the cable. The distance should be calculated with the corresponding formula of the transmitter frequency. The recommended isolation distance: $d=1.2\sqrt{P}$ $d=1.2\sqrt{P}$ 80MHz~800MHz $d=2.3\sqrt{P}$ 800MHz~2.5GHz In formula: P — Maximum output rated power of the transmitter provided by the manufacturer, unit for Watt(W) d—Recommended isolation distance, unit for meter(m).

Note 1: at 80MHz and 800MHz frequency point, use the formula for higher frequency bands

**Note 2:** these guidelines may not be suitable for all cases, because the electromagnetic propagation is influenced by the absorption and reflection of buildings, objects and human bodies.

a.fixed transmitter, such as wireless(cellular/cordless) telephone and ground mobile radio base station, amateur radio, Am and FM radio and television broadcasting, etc. the electric field intensity can not be accurately predicted in theory. In order to evaluate the electromagnetic environment of a fixed RF transmitter, the survey of electromagnetic

field should be considered. If the electric field intensity measured where [Prototype

**ARK7680**] place is higher than above applicable RF Meet Level, [**Prototype ARK7680**] should be observed to verify whether it can work normally. If abnormal performances happen, the supplementary measures may be necessary, such as re-adjust the direction or position of[**Prototype ARK7680**]

b.in the entire frequency range of 150 kHz $\sim$ 80 MHz, the electric field intensity should be less than3V/m.

#### 1.4 The Recommended Isolation Distance Between Portable and Mobile Radio Frequency Communication Equipments and [Prototype ARK7680] (Form 4)

The recommended isolation distance between portable and mobile radio frequency communication equipments and [**Prototype ARK7680**]

[**Prototype ARK7680**] expected to be used in the electromagnetic environment of the radio frequency radiation disturbance controlled. According to the maximum output rated power of communication equipment, the buyer or user may prevent the electromagnetic interference by maintaining a minimum distance to be recommended as following items between the portable and mobile radio frequency communication equipment(transmitter) and [**Prototype ARK7680**]

	Isolation distance	ce of different frequent	ncy of transmitter/m
Maximum output rated power of transmitter:	150kHz~ 80MHz	$ m 80MHz{\sim} m 800MHz$	800MHz~2.5GHz
W	d=1.2√P	d=1.2√P	d=2.3√P
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

To the maximum output rated power of transmitter that not listed in the above forms,

d is recommended as isolation distance, unit for meter(m), the formula in the frequency column of the corresponding transmitter is available, here p is the maximum output rated power of transmitter provided by the manufacturer, unit for Watt(W).

Note 1: at 80MHz and 800MHz frequency point, use the formula for higher

frequency bands

**Note 2**: these guidelines may not be suitable for all cases, because the electromagnetic propagation is influenced by the absorption and reflection of buildings, objects and human bodies.

#### **2. SAFETY PRECAUTIONS**

#### 2.1 Operation

- 2.1.1 Don't optionally open and touch the inside parts of the instrument, it may cause an electric shock or the system may malfunction.
- 2.1.2 Please keep this instrument ground connection well to avoid possible injury to people or the instrument damaged.
- 2.1.3 Don't touch the screen(resistance touch-screen, contact and hold it for about 0.1 second) in too strong strength, it may damage the screen.
- 2.1.4 Don't put the instrument at the place of direct sunlight or too strong illumination, it may affect the measuring precision. It's strongly suggested to be used indoors or in darkroom.
- 2.1.5 Don't use the instrument in a hot, humid or dusty environment. Such environments cause bad influences to the instrument.
- 2.1.6 If you want to connect this instrument to other instrument, please follow our local agent's instructions.
- 2.1.7 In cold room, when temperature suddenly rise, dew maybe appear on the protection glass of measuring window or internal optical parts. In case this happen, it can be used till the dew disappear.
- 2.1.8 Keep the measuring window lens clean at all time. The dust and other substances may cause error in measuring or affect the measuring precision.
- 2.1.9 If you encounter any abnormal conditions, such as smoking or strange smells, turn off the instrument and pull out the power cord immediately. Contact the local experts/agent or original manufacturer to check and repair, you can use till the trouble is absolutely removed.
- 2.1.10 The use of materials that directly contact with the skin part: During operating the instrument, it should be used to separate the instrument from the patient's touch part with medical non-woven fabric(size for 8cm x 8cm), to avoid the direct contact with the surface of the instrument.

#### 2.2 In Storage

- 2.2.1 Don't store the instrument in a place where it may get wet or where poisonous gas or liquid is stored.
- 2.2.2 Be sure to store the instrument in a place away from direct sunlight and with the specified temperature and humidity.

#### **2.3 In Transference**

- 2.3.1 During carrying the instrument, please take great care to avoid colliding and falling. Sudden or strong impact may damage the instrument or performances.
- 2.3.2 Before carrying, please turn off the machine and lock tightly the sliding body. During carrying, please catch the bottom tightly by two hands.

#### 2.4 After Using

- 2.4.1 If the instrument won't be used for a long time, disconnect the power cable from the wall-outlet. It may cause a fire.
- 2.4.2 When the instrument is not used, turn the power off and put the dust cover on. Keeping the machine in electricity supplying will reduce the use life of the instrument. If the instrument is not covered for a long time, dust may affect the measuring accuracy.

#### 2.5 In Maintenance

- 2.5.1 It's one high precision optical instrument and need to be calibrated regularly.
- 2.5.2 Please fill the lube to the sliding parts regularly at the experts' guide.
- 2.5.3 Be sure to replace the fuse after disconnecting the power cord from the power inlet and use the specified fuse for replacement. Otherwise, it may cause a fire.
- 2.5.4 In case the instrument breaks down, it must be checked and repaired by the specified experts who know this instrument very well, or contacting the local authorized agent or original manufacturer. Open and repair the instrument by oneself, the agent or manufacturer don't be in charge of the consequence.
- 2.5.5 This instrument for non sterile medical device. Daily cleaning and disinfection of device by end user. Please use a soft cloth or sponge, wet cloth or detergent to clean the device. Don't use alcohol, water, benzene and other organic compounds to clean the surface of the instrument, to avoid damage to the device. The measuring window is often cleaned by a soft cloth to remove dust to maintain the accuracy of the measurement.
- 2.5.6 Determination of disinfection method:

According to the requirements of "environmental and object surface disinfection" in the hospital disinfection and sanitation standard of WS310-2, the general components of the device are cleaned in time, the chin-rest and forehead rest and operation lever are demanded to be disinfected in middle level disinfection.

In accordance with the regulations of the WS/T 367-2012 medical institutions disinfection technical specifications, the chin-rest and forehead rest and operation lever are demanded to be disinfected by using alcohol disinfectant, and use 75% (volume ratio) ethanol solution to wipe the surface of the object.

#### **3. UNPACKING AND INSTALLATION**

#### 3.1 Notices and Procedures of Taking out the Instrument

Catching the bottom and chinrest frame separately by two hands, don't catch the screen or operation lever (Figure 1)





#### 3.2 Power Line Inlet and RS232 Interface (Figure 2.3)

Connect the spare power line with the power supply socket (RS232\USB interface connection demanded, please contact the original manufacturer or local agent)



(Figure2) (Figure3) Power Supply Socket: AC power inlet (fuse F5AL 250V inside) Data Interface: RS232\USB interface to be connected with other equipments (This refractometer and the automatic phoroptor serials connected should comply with the related electrical requirements inIEC60601-1-1-2000)

#### **3.3 Chinrest Paper Installation**

Use the specified chinrest paper (Figure 4)



#### **4.** FUNCTIONS OF THE MAJOR COMPONENTS Front(Figure 5)



(Figure 5)

LCD Screen: Monitor for measurement display Height Adjustment Mark: The eyes' height position of the patients Printer Cover: Press the cover to open or close Measure Button: Performing the measurement by pressing after focusing Joystick: Adjust the focus by moving it left/right, up/down, forward/backward Stage Fixing Lever: Lock the sliding body Power Switch/Socket: AC power on/off with indicator light / AC power inlet (fuse F5AL 250V inside)

Rubber Feet: Support and adjust the instrument horizontal

Chinrest Up/Down Button: Adjust the height of the chinrest

Data Interface: USB\RS232 interface to be connected with other equipments (This refractometer and the automatic phoroptor serials connected should comply with the related electrical requirements in IEC60601-1-1-2000)

Back (Figure 6)



(Figure 6)

Chinrest: The platform for placing the patients' chin Forehead Rest: The place against the patients' forehead Measuring Window: Imaging on the retina of the patients' eyes

#### **5. MAIN TECHNICAL INDEXES**

#### 5.1 Measurement Performance Parameters

5.1.1 Corneal Vertical Distance(VD): 0.00mm、12mm、13.75mm、15mm

5.1.2 SPH: -30.00 m<sup>-1</sup>~ +25.00 m<sup>-1</sup>( VD=12mm, 0.01 m<sup>-1</sup>, 0.06 m<sup>-1</sup>, 0.12 m<sup>-1</sup>,

0.25 m<sup>-1</sup> unit ), deep myopia measurement available

5.1.3 CYL: 0.00  $m^{-1} \sim \pm 10.00 m^{-1} (0.25 m^{-1} , 0.25 m^{-1} unit)$ 

5.1.4 Cylinder Form:  $-, +, \pm$ 

5.1.5 Axis(AX): 1°~ 180° (1° unit)

5.1.6 Pupil Distance(PD): 10mm ~ 85mm(0.1mm unit)

5.1.7 Radius of Corneal Curvature: 5.0 ~ 10.0mm (0.01mm unit)

5.1.8 Corneal Power: 33.00 m-1 $\sim$  67.00 m-1 (in case that the corneal equivalent refractive power is 1.3375)

5.1.9 Corneal Astigmatism: 0.00 m-1 ~ 15.00 m-1 (0.06 m-1/0.12 m-1/0.25 m-1 unit)

#### **5.2 Other Performance Parameters**

5.2.1 8"TFT touch screen (angle adjustable)

5.2.2 Printer: 57mm thermal printer

5.2.3 Measuring Light Energy: < 30uW (prevent injury to eyes during

measuring)

5.2.4 Measuring Time: <0.5s

5.2.5 Minimum 2.0mm pupil can be measured. The application of cloud and mist chart technology allows the patients' eyes to look at the internal targets in a natural and comfortable situation and make the measurement more accurate  $5.2 \text{ (Flucturing)} \text{ Parsaux AC100} = 240 \text{ M} \cdot 50 \text{ (OUT)}$ 

- 5.2.6 Electrical Power: AC100  $\sim$  240V, 50/60Hz
- 5.2.7 Consumption: 60AV

5.2.8 N.W.: 17.25kg

5.2.9 G.W.: 22.5kg

5.2.10 Dimensions: L750mm×W400mm×H630mm

#### **5.3 Protection Level**

5.3.1 Product Grade: Medical apparatus and instruments grade II

5.3.2 Electric Shock: Level I (ground)

5.3.3 Electric Shock Protection Class: Class B

#### **5.4 Device Type**

5.4.1 Anti Electric Shock Type: Class I

5.4.2 Anti Electric Shock Degree: Applicable type B

5.4.3 Non AP device, non APG device

5.4.4 Running Mode: Continuous duty

#### **6. ENVIRONMENT TERMS**

- 6.1 Temperature:  $10^{\circ}C \sim 30^{\circ}C$
- 6.2 Relative Humidity: (30~75) %RH

6.3 Atmospheric Pressure: 86kPa ~ 106kPa

6.4 Altitude: <2000m

6.5 No strong vibration and corrosive gas around

6.6 No strong electromagnetic interference around

6.7 Brightness: <150Lx

6.8 The device should be placed at the specified instrument table that can rise and fall vertically

6.9 The device can't be used in the environment of flammable and anesthetic gas



#### 7. LCD SCREEN DISPLAY (Figure 7)

#### (Figure 7)

- 7.1 L/R Sign: Flashing sign indicates the current measured eye
- 7.2 The number of power/corneal parameters measured
- 7.3 Step selection (shortcut key)
- 7.4 Astigmatism symbol selection (shortcut key)
- 7.5 VD selection (shortcut key)
- 7.6 Lattice display
- 7.7 Power display
- 7.8 Corneal value display
- 7.9 Adult//child mode selection
- 7.10 Measurement mode selection
- 7.11 Auto/manual measurement selection (partial model)
- 7.12 Auto/manual tracking and focusing selection (partial model)
- 7.13 Data clear key

- 7.14 Data record check
- 7.15 Menu set
- 7.16 Pupil alignment target
- 7.17 Left/right eye pupil diameter

#### 8. MENU (Figure 8)

#### 8.1 Measurement Mode Selection

Touch this key to pop up three measurement mode menu (as shown in Figure 10), the user can choose to touch any measurement mode menu under need (KER for Keratometry mode, R/K for RefracKeratometry mode, REF for Refractometry mode)



(Figure 8)

#### 8.2 Auto/Manual Measurement Selection (partial model)

Touch this key to select auto measurement mode (A) or manual measurement mode (M)

#### 8.3 Auto/Manual Tracking and Focusing Selection ((partial model))

Touch this key to select auto tracking and focusing mode (AF) or manual tracking and focusing mode (MF)

#### 8.4 Data Clear Key

Touch this key to clear the measurement data

#### 8.5 Data Record Check

Touch this key to check the measurement data (directly print out the data, the measurement data won't be recorded)(Figure 9)

			DATA F	RECOR	D			
R 1 2 3 4 5 6 7 8	SPH -3.00 -3.00 -3.00	CYL 0.00 0.00 0.00	AX 180 180 180	L 1 2 3 4 5 6 7 8	SPH -3.00 -3.00 -3.00	CYL 0.00 0.00 0.00	AX 180 180 180	
9 10 AV(	G -3.00	0.00	180	10 AVG	-3.00	0.00	180	
	REF	KER		1	4			

#### (Figure 9)

Left/right eyes data can be recorded max.10 items separately. Touch REF to display the recorded refractometry data only, touch KER to display the recorded keratometry data only, touch CLEAR key to clear the recorded data, touch RETURN key to return to the measurement interface.

#### 8.6 Menu Set

Touch menu set key to enter the subsidiary menu setting (Current selection for blue)

8.6.1 Refractometry parameters setting (Figure 10)

VD: Distance between corneal and back top focus of lens, 0.0mm (contact lens), 12.0mm (Asian), 13.5mm (Middle East), 15.0mm (European)

CYL: Astigmatism symbol selection, -, +,  $\pm$ (Mix)

STEP: Measurement data precision selection

FOGG: Visual guide target atomization function switch (position of guiding target atomization)

#### (Figure 10)

8.6.2 Keratometry parameters setting (Figure 11)



(Figure 11)

MODE : Keratometry radius measurement (mm), keratometry power measurement (m-1) and average value display (AVG) optional

STEP: Keratometry power precision display

**REFRACTIVE INDEX:** Factory defaults to 1.3375

8.6.3 Mode setting (Figure 12)

#### (Figure 12)

MODE: Manual measurement mode and auto measurement mode optional (Auto

measurement icon for grey said this model without this feature)

BEEP: Sound prompt when operating. If set off, operation will keep silent

INT-M: Measurement mode selection (same as the main interface function), default startup mode for each starting

STAND BY: Instrument standby time setting (5 minutes, 10 minutes, 30 minutes and 60

minutes optical) (touch any key to wake up)

8.6.4 Printing setting and printing paper replacement (Figure 13 14)







AUTO: When ON selected, the measurement results will be printed out automatically after the both eyes measurement finished (in this case, the data is cleared automatically)When OFF selected, press the print key on panel to print out the measurement results.

REFRACTOMETRY: When OFF selected, the refractometry power won't be printed out. When STD selected, all refractometry power will be printed out. When AVG selected, only print the average value of the refractometry power.

KERATOMETRY: When OFF selected, the keratometry power won't be printed out. When STD selected, all keratometry power will be printed out. When AVG selected, only print the average value of the keratometry power.

EYE: When ON or OFF selected, the refractometry state diagram will be printed or not.

CONCENTRATION: Set the appropriate print concentration according to different thermal printing paper.

PUPIL: When ON or OFF selected, the pupil diameter will be printed or not.

SE: When ON or OFF selected, SE data (the approximate value of cylinder power converted into sphere power) will be printed or not.

BC: When ON or OFF selected, BC(base curve of corneal contact lens) will be

printed or not.

How to install the printing paper (Figure 15)

- 1.) Pull outwards the printer cover, open the cover
- 2.) Place the new printer paper in the box, keep the paper head upwards
- 3.) Draw the paper outwards and directly ride on the printer wheel
- 4.) Push the cover back and close the cover



(Figure 15)

#### 8.6.5 Data setting (Figure 16)

	SE	T MENU	J			X
REF KER	NODE F	PRINT	DATE	MES	TRANS	
DISP >	YMD	MDY	DMY			
DATE >	Y:2014	M:10	D:11			
TIME >	H:02	M:07	S:54			
COUNT >	OFF	ON				
No. >	0039					

(Figure 16)

DISP: Date, month and year display mode

DATE: Edit or modify the exact time of date and month and year

TIME: Edit or modify the exact time of second and minute and hour

COUNT: When ON or OFF selected, recording the number of patients in main interface will be refreshed or not

No.: Patient number setting, patient measuring number setting

Touch DATE, TIME and NUMBER options, enter the sub menu as shown below, select the appropriate number, press ENT key to confirm and preserve, press RETURN key to quit. Press BS key to delete one by one, press C key to clear all. (Figure 17)



(Figure17)

8.6.6 Printing message setting (Figure 18)

MSG1 for company name or product model number setting

MSG2 for company address or brand name setting. Users can edit this information freely according to the exact requires. After setting, press ENTER key to preserve and quit.



(Figure 18) Touch the yellow blank space to enter the message editing menu (Figure 19)

RE	FK	ER	MOI	DE	PRIN		DATI		ES	TRA	NS
			F	PRIN	т ме	SSA	GE				X
AR	K 768	0									_
1	2	3	4	5	6	7	8	9	0	-	+
Q	w	E	R	Т	Y	U	1	0	P	SPA	CR
A	S	D	F	G	H	J	к	L	:	8	BS
z	×	С	v	в	N	м			A/a	EN	TER

(Figure 19)

ENTER key for confirming and preserving A/a for capital/small letter conversion BS for deleting single letter SPA key for space bar CRL key for clearing all letters

8.6.7 Data transfer setting (Figure20)

DE PRINT DATE MI	5 TRANS
600 19200 57600 1	200
OFF ON	

#### (Figure20)

According to the requires of the connected devices, customers choose the corresponding baud rate, and open the auto option, the measurement data will be automatically transferred to the connected devices, meanwhile the refractometer data will be automatically cleared.

#### 8.7 Shortcut Key

- 8.7.1 Step set: successively touch STEP key to quickly switch 0.01、0.06、0.12、 0.25 steps
- 8.7.2 CYL axis set: successively touch CYL key to quickly switch  $1 + 1 \pm 1$
- 8.7.3 VD set: successively touch VD key to quickly switch 0, 12, 13.5, 15
- 8.7.4 Array display: touch array key to display the patient's fundus array distribution (indirect evaluation of fundus imaging quality). (Figure.21)



(Figure.21)

#### 9. MEASUREMENT

Suitable crowd and contraindication

Target patients for adults and children, and crowd of eye power range (-30 m<sup>-1</sup>  $\sim$  +25 m<sup>-1</sup>). This product is not suitable for newborn eye measurement.

#### 9.1 Preparations before Measurement

9.1.1 Place the device on the specified instrument table, loose the stage fixing lever and keep the device in free sliding state, adjust the four rubber feet to keep the device in horizontal.

9.1.2 Fix and install the specified chinrest paper and printing paper separately

9.1.3 Connect the spare power line to the instrument socket tightly (ensure the local voltage fit to the instrument specification)

9.1.4 Turn on the left side power switch (green indicator light show right in electricity connection), the instrument goes into self-check procedures. After self-check over, it automatically switches to main interface for measurement.

#### 9.2 Notes for Operator and Patient

- 9.2.1 Adjust the chair height and screen angle in right position
- 9.2.2Ensure the patient in comfortable and relaxed posture before measurement
- 9.2.3 By adjusting the instrument tabletop, keep the instrument height same to the patient natural sitting posture
- 9.2.4 Settle patient's chin touch the chinrest front and forehead touch the rubber forehead rest in level (keep face parallel with the measurement window)
- 9.2.5 By observing the patient eyes position and height adjustment mark, press the chinrest up/down key on panel to adjust the patient's eyes same height to the measurement window
- 9.2.6 By the operation lever, move the sliding body left and right to move the patient's eyes in the measurement range (if the distance of two sides asymmetrical, adjust it by fixing the patient's head deviation)

#### 9.3 Measurement

The measurement alignment method of this device for pupil and center measurement cross target in coincidence

9.3.1 Normal Measurement Mode

Holding the operation lever, quickly shift the sliding body to left side, keeping the measurement window roughly aligning with the patient's right

8 S S C С A A **R**1 **R1 R**2 R2 AX AX RE MF 0

eye socket (Figure.22)

(Figure.22)

Observing the patient's eye location on screen, rotate the operation lever (up and down adjustment), meanwhile swing the operation lever left and right, till the yellow cross-ring target aligning at the patient's corneal vertex, then shift the operation lever front and back, till the patient's eye is clearly focused in the center measurement socket (the accuracy of focusing can be confirmed by observing whether level between the two points of split focusing and cross-ring

target) (Figure.23,24)





(Figure 24)

Prompt the patient to open eyes wide (eyelid and eyelash covering eyeball will affect the measurement accuracy), both eyes look right ahead.

Slightly adjust the operation lever, till the two points of slit focusing level with the cross-ring target, and yellow cross measurement target becomes thick and green, press the measurement button, when the measuring light flashing (the screen refreshed in black in moment), it shows the measurement over (the patient no need to see clearly the object-image during measuring, the measurement result same accuracy). The measurement result will be displayed

on screen. (Figure.25)  $\circ$ 



(Figure.25)

Shift the sliding body to right side, repeat the above steps, measure the patient's left eye.

Both eyes measurement over, pupil distance will be displayed automatically on the corresponding position. Choose whether or not to print the measurement results according to settings (auto printing or data output transmission over, the data on screen will be automatically cleared).

9.3.2 Child measurement mode

To measure children or the patients with pupil fibrillation, select child mode (Touch 7.9 key, the right small humanoid icon becomes green). (Figure 26)



(Figure.26)

#### 9.3.3 Measurement error prompt

During measurement, if the patients found having eyelid ptosis, eyelash disturbance, cataract, microcoria, keratopathy, corneal vertex and pupil center noncoincidence, the error prompt will appear on screen when the instrument can't measure normally, please select the manual measurement mode or force measurement mode (long press the measurement button). (Figure 27 28).



```
(Figure 27)
```



#### **10. COMMON TROUBLE SHOOTING**

#### **10.1 Power indicator light not work**

Check and confirm whether the local power fits to the instrument, whether the power plug loose, or whether the fuse damaged (in case this happen, please replace the same specified fuse)

#### **10.2** Chinrest not lift

Check whether the chinrest lift to limit position

#### **10.3 Printer can't work regularly**

Check whether the printing paper is finished (in case this happen, red indicator

light on panel will flash). Or whether the print setting is correct, and whether there is the measurement data (no data, not print)

#### 10.4 Sliding body not flexible

Check whether the stage fixing lever placed at right position, or whether other sundries go into the slide slot

#### 10.5 Press measurement button, but no data appear

Check whether the patient pupil smaller than 2mm, whether the eye position seriously incorrect, whether the cross measurement target aligns with the patient pupil (the target becomes thick and green), or whether the patient eyeground seriously deseased

#### **10.6 Measurement light not work**

When the measurement over, the measurement light will automatically turn off. Sway the sliding body, it will turn on automatically

## 10.7 If other problems appear, please contact the local agent or original manufacturer

- If the fault phenomenon listed in the common trouble shooting can't be resolved, please contact the original manufacturer or local agent to repair.
- Please provide us with the following information:

Instrument name and model number

Instrument serial number

Fault phenomenon (detailed as possible)

(1.) Accessory maintenance limitation

Providing maintenance accessories to maintain the instrument functions during the instrument lifetime

(2.) Processing of instrument

• To be disposed carelessly of the instrument and accessories will pollute the environment

• Please contact the professional waste disposal company or local dealer before disposing this instrument

#### **11. PACKAGING, TRANSPORTATION, STORAGE**

Storage condition between -25°C and +40°C, transportation condition between -40°Cand 70°C, relative humidity between 30% and 75%, air pressure between 86kpa and 106kpa

(1.)Packing list, certificate and manual are included in the packing box

(2.)The product packaging is not allowed to be shipped with flammable, explosive, corrosive products. Loading should be neat, stable and firm, super high and overweight is not allowed. In transit, rain and snow prevention, anti sun, anti impact, drop prevention should be noted carefully.

(3.)The product packaging should be stored in a room temperature, dry and well ventilated warehouse, and can't be stored with chemical agents, acid and alkali substances, and other harmful substances.

#### **12. ENVIRONMENTAL PROTECTION**

The instruments that have be scrapped, should be strictly deposed in accordance with the requirements of local laws and regulations

#### **13. ENCLOSURE ACCESSORIES**

Number	Specification	Quantity
1	User's Manual	1
2	Dustproof Cover	1
3	Lens Dustproof Piston	1
4	Model Eye	1
5	Power Line	1
6	Chinrest Paper	1
7	Cleaning Cloth	1
8	Printing Paper	1
9	Fuse	2
10	Chinrest Pin	2



#### **Technical Parameter**

SPECIFICATIONS: Sphere 0.00~-19.00 Diopters, step: 0.25,0.5,1,3 0.00~+16.75 Diopters, step: 0.25,0.5,1,3 ●Cylinder: 0.00~±8.75 Diopters, step: 0.25,0.5,1,3 • Cylinder Axis:  $0^{\circ}$  to  $180^{\circ}$ , step:  $1^{\circ}$ ,  $5^{\circ}$ Prism form: Base up, base down, base in, base out Near distance: Mechanical converging with an optical axis at 30 to 40 in front of eye Binocular balance: Rotary prism, polarizing filter and R & G fliter ●Cross cylinder: ±0.25D ●PD: 50 to 80mm in 1 and 0.5mm step •Retinoscope: +2.00D or +1.50D both sides Corneal alignment 12 to 20mm by 2mm step • Field of view 35 $^{\circ}$ • Power source: AC 110-120V/220-240V, 50/50Hz, 90VA •Dimension: 350 (W) x300 (H) x80 (D) mm •Net weight: 4kg • Power supply: AC 110-120V or 220-240V 50/50Hz

●Horsepower: 60w

## <sup>®</sup> *KF-ZD6000* Auto-Phoropter

P

•



# **KF-ZD6000**

## **Auto-Phoropter**

Bluetooth wireless communication provides flexible operation space for optometrists Simple operation interface provides efficient and convenient optometry experience Innovative design, tablet control, designed for fashion stores Support personalized programming, which is conducive to the unified optometry standard of chain stores









#### Cross column mirror adjustment



Auxiliary lens display area





#### **D** Spherical mirror step adjustment

FAR	RESET	PRINT	SAVE	TIP	HELP
R G H T E Y E	0.00	S(0.25)		0.00	
	0.00	C-(0.25)		0.00	L
	180	A(1)		180	F
	BI0.0	H(0.1)		B00.0	E
	BD0.0	V(0.1)		BU0.0	E
	+0.00	ADD		+0.00	



#### Functional visual display area



## COMPUTERIZED VISION TESTER OPERATOR' S MANUAL

KF-ZD6000

#### Operation environment of optometer control software

Function description of optometer control software optometer control software is a tablet control software based on Android system. It controls the optometry system to perform specified operations, interacts with customers, carries out optometry analysis and diagnosis, and finally

generates optometry report and provides corresponding Glasses Prescription. Through the WiFi module, the optometer control software takes the tablet computer as the center, and forms the isolated units in the traditional optometry system, such as the operation end, the optical head, the computer optometer, the liquid crystal visual acuity chart and the prescription printer, into a WiFi LAN, which realizes the real-time data transmission and forms an organic whole. Through the way of WiFi communication, the more rapid and stable lens combination control of the optometer is realized, the accurate and rapid combination of spherical lens power, cylindrical lens power, axial position and prism power is realized, and the fast switching of special auxiliary lenses is realized: the optometry of the computer optometer is quickly read and loaded; Realize the sensitive and reliable control of LCD visual acuity chart: realize the reliable output of printing information of prescription printer. The optometrist control software provides a more comprehensive, convenient and powerful optometry and auxiliary system for optometrists. Through the

touch operation of the tablet computer, the rapid optometry lens combination switching operation is realized: by providing a user-defined program editing system, some optometrists' optometry experience can be saved in the form of program steps, which greatly improves the optometry efficiency: by enriching and perfecting the operation of the liquid crystal visual acuity chart, the visual acuity chart can be directly switched to the specified state, The flexible control operation of LCD visual acuity chart is realized, and the user-defined visual acuity chart editing function is introduced to greatly improve the efficiency of optometry: the optometry results are completely saved and shared in real time by establishing a customer data database, so as to push the traditional optometry to the Internet data age. The optometer control software gives a new interpretation of the traditional optometry process, integrates the advanced mobile terminal into the modern optometry system, and integrates the ideas of optometrists' personality customization and the Internet data age, bringing a fast, fashionable and scientific optometry experience.

Kingfisher manufacture CO,.LTD.

#### 1. Introduction

Computerized vision tester is used to subjective exa mination of visual acuity and refractive error of subject's eyes.

#### 1.1 Classify

Model number: KF-ZD6000 Security category: I class Medical electrical equipment

#### 1.2 Storage and transport environment conditions

a)temperature:-10°C~55°C b)Air pressure:700hpa~1060hpa(transport), 500hPa=1060hPa(storage) c)Humidity:10%~85%

#### 1.3 Working environment condition

a) temperature:10°C~40°C
b) Air pressure:800hpa~1060hpa
c) Humidity:10%~80%

#### 1.4 Power adapter requirement

a)input: AC100-240V<sup>~</sup>5060Hz b)output: DC24V2.5A85VA

#### 2. Main performance index

- 2. 1 Measurement range
- a) Sphere: +16.75D  $^{\sim}$  -19.00D;
- b) Cylinder: 0  $^{\sim}$  -6.00D;
- c) Axis: 0  $^{\sim}$  180 $^{\circ}$  ;
- d) Prism:  $0 \triangle \sim 20 \triangle$ ;
- e) Prism base: 0  $^{\sim}$  360 $^{\circ}$  ;
- f) PD range: 50  $^{\sim}$  80mm;

#### 2.2 Step

- a) Sphere: 0.25D/0.5D/1.0D/3.0D;
- b) Cylinder: 0.25D;
- c) Axis: 1°  $/5^\circ$  ;
- d) Prism: 0.1cm/m;
- e) Prism base: 1° ;
- f) PD range: 0.5cm;

### 2.3 Tolerance

a) The tolerance of sphere should accord with the requirement table 1.1

item	Standard	tolerance	Tolerance  s1-s2
	0≤ Ö ≤3.00	$\pm 0.06$	
	3.00< Ö ≤6.00	$\pm 0.09$	
☆/注 (D)	6.00< Ö ≤9.00	$\pm 0.12$	0.02
环境/夏0(D)	9.00< Ö ≤12.00	$\pm 0.15$	0.03
	12.00<  Ö ≤15.00	$\pm 0.18$	
	15.00< Ö	$\pm 0.25$	

Table 1.1 sphere tolerance

Table 1.2 Astigmatic power tolerance

Meridian direction		Astigma	tic power nomina	l diopter	
Maximum absolute	≤0.50	>0.50~1.00	>1.00~3.00	>3.00~6.00	>6.00
dioper			Tolerance D		
0.00~5.00			0.06	0.00	0.19
>5.00~10.00	0.06	0.06	0.00	0.09	0.12
>10.00~15.00	0.06		0.09	0.12	0.18
>15.00		0.09	0.12	0.18	0.25

item	Nominal diopter(Absolute)	Tolerance
Culindan ania	>0D~0.25D	$\pm 5^{\circ}$
Cylinder axis	>0.25D~1D	$\pm 3^{\circ}$
	>1D	$\pm 2^{\circ}$
	≪1cm/m	$\pm 5^{\circ}$
Prism base	>1cm/m~10cm/m	$\pm 3^{\circ}$
	>10cm/m	$\pm 2^{\circ}$
DD&VD (mm)	VD	$\pm 0.5$
	PD	$\pm 0.5$

Table 1.3 Cylinder axis and prism base tolerance

Nominal diopter(Absolute)D	Tolerance (cm/m)
0.00	0.12
>0. 00~6. 00	0.25
>6.00~12.00	0. 37
>12.00	0. 50

# Table 1.4 optical center tolerance of mechanical system of sphere and cylinder combinative optical system

### 2.1 Function introduction

2.1 The minimum aperture of all lenses of the optometer shall not be less than 16mm; For prisms of 6 cm / M and above, the minimum luminous aperture can be reduced to 11mm.

2.1.2 There are mirror eye distance monitoring windows on both sides of the instrument of the comprehensive optometer. Through the mirror eye distance monitoring window, it can be observed that a dividing plate is engraved with a long line and three short lines, and the spacing between two adjacent lines is 2mm. The position of the anterior apex of human cornea should be on the long line of the dividing plate, which is the reference plane of the instrument of the comprehensive optometer.

2.1.3 The comprehensive optometer shall be equipped with a covering and splitting device, and each system shall be equipped with a Jackson cross column mirror;

2.1.4 There is no stray light interference or structural occlusion in

the observation optical path of the optometer.

2.1.5 The structure of lens cavity shall not affect the examination of patients' visual function;

2.1.6 When the lens and accessories are fixed in front of the observation hole, the instrument shall be aligned and centered.

2.1.7 The operation keys of the optometer shall be clearly marked. When pressing the keys, the instrument function or state shall change to normal.

2.1.8 When the optometer has WiFi function, the communication is effective at 10m.

### 3 Security con siderations

### 3.1 Security identity

In this manual, the words used to indicate the degree or level of safety alarm are defined as follows:

Warning: indicates a potentially dangerous situation that, if not avoided, may result in death or serious injury

Note: it indicates a potential dangerous situation. If it cannot be avoided, it may lead to slight or medium injury or property loss. Under certain conditions, even if the situation is attention, it may lead to serious injury. Therefore, safety precautions must be strictly observed at all times

Safety precautions before use

### warning

1. Make sure the optometer is fixed on the support arm of the

2. If the optometer drops or falls, it may cause personal injury or equipment failure.

3. Ensure that the power box is installed and placed vertically according to the method shown in Figure 1 in 1.2 classification in this manual.

caution

1. Safety precautions and operating procedures must be thoroughly understood before using the equipment

2. Unexpected use may lead to unexpected failure or adverse results.

3. The equipment shall not be exposed to rain, water or fog.

Do not place containers containing liquid or gas on the top of

the instrument.

4. The storage environment shall not be in dusty, hot, humid places or direct sunlight

5. Do not carry the equipment to another place alone, which may damage the back or slide the equipment

6. Install the equipment on a stable and horizontal table. If the equipment slips, it may cause injury or equipment damage Do not install equipment where there is water. Contact with liquid may cause electric shock or equipment failure

7. Please install the equipment in an environment that meets the following conditions. The following conditions must also be met in the use of the equipment.

a)temperature:10℃~40℃

b)Air pressure:800hpa~1060hpa

c)Humidity:10%~80%

Installation position: dust-free and clean dark light room

Place free from vibration and impact

8. Be sure to adjust the horizontal refractometer before use.

If the refractometer is not installed horizontally, it may affect the accuracy of the data. The level adjustment knob is used to calibrate the refractometer until the foam concentration is at the level.

9. Be sure to use sockets that meet the requirements of equipment voltage parameters. If the voltage is too high or too low, the equipment cannot operate normally, and the fault may cause fire

10. To avoid the risk of electric shock, the equipment must be connected to the power supply with protective grounding.

11. Do not overload the socket, which may cause fire

12. Fully insert the power plug into the socket.

13. Improper connection may cause fire.

14. Do not use any power cord other than equipment, which may cause failure or fire.

15. Do not place heavy objects on the power cord.

15. Damaged power cord may cause fire or electric shock.

16. When installing and operating the equipment, observe the following EMC instructions (electromagnetic compatibility):

It cannot be used together with other electronic equipment to avoid electromagnetic interference between equipment operation and other electronic equipment

It cannot be used in the same room with other equipment, including life support equipment, other equipment that has a significant impact on the patient's life and treatment results, and measuring or therapeutic instruments containing small current

It cannot be used simultaneously with the equipment of portable and mobile radio frequency communication systems, because the electromagnetic interference they emit may adversely affect the operation of the device.

Do not use cables and accessories not specified by the company, which may increase the electromagnetic wave emission of the equipment or system and reduce the anti electromagnetic interference ability of the device. If there is potential electromagnetic interference between the equipment and other equipment, shielding measures shall be taken or the installation position of the equipment shall be changed to reduce the possible interference

### Maintenance and inspection

Any repair and service of this instrument must be provided by personnel trained by Hangzhou Jingfei Optical Instrument Manufacturing Co., Ltd. Be able to operate correctly and maintain by experienced personnel or dealers. Removing the safety screw may cause the instrument to separate from the support arm and cause serious injury. Do not open the outer cover of the instrument or try to repair any internal parts. Any maintenance and service of the equipment must be carried out by experienced staff of Hangzhou Jingfei Optical Instrument Manufacturing Co., Ltd. or authorized distributors trained by Hangzhou Jingfei Optical Instrument Manufacturing Co., Ltd

The adjustment of the instrument must be carried out by the technical service personnel or other authorized personnel of Hangzhou Jingfei Optical Instrument Manufacturing Co., Ltd

The use and operation of the optometer must be in strict accordance with the instructions in the user's Guide. If it is not used in the way specified by Hangzhou Jingfei Optical Instrument Manufacturing Co., Ltd., the patient's safety and the normal operation of the instrument will not be guaranteed

Do not use solvent or strong cleaning solution on any part of the instrument, otherwise the instrument may be damaged

Do not use organic solvents such as paint thinner to clean the outside of the equipment, which can damage the surface of the equipment

Do not immerse the optometer in liquid, otherwise it will cause damage to the instrument.

Avoid touching the optical components of the instrument to prevent performance degradation caused by fingerprints or oil stains on the lens group. Note: pixels may occasionally disappear or appear red, blue or green pixels on the screen. This does not mean that the LCD panel is broken: it is caused by the LCD screen in the production process.

### 4. Configuration

### 4.1 Refractor head con figuration





- (1) Examination window: The patient's eyes are observed through the window, and patient observe charts through the window.
- (2) Near vision illumination: Light up when bear vision testing, light off when far vision testing.
- (3) Corneal aligning windows: The position of patient' s cornea can be observed through the window.
- (4) Forehead rest adjust knob: Moves the forehead rest forward and backward in order to adjust the VD.
- (5) Near-point rod holder: the near\_point rod is inserted and attached here.

(6)Near\_point rod clamp screw.....Fix the near-point rod.

(7) spirit level..... Used to confirm that the refractor head.

- (8)Leveling knop..... Adjust the level of the reractor head.
- (9) Forehead rest magnet..... Fix the face shield.
- (10) forehead rest ..... The patient's forehead rests here when test.

(11)Cornea alignment scale.....Measures the VD.

(12)Face shield(2sets)..... Attached to the instrument to put the patient's face in place, use one while the other is being sterilized(13) Dust cover..... Covers and protects the instrument body from Dust and dirt during storage.



(14) Forehead rest (2 sets) ..... The patient's forehead rests here. Detachable by a touch use one while the other is being sterilized

(15) Near-point rod..... The card holder is attached at the near point examination

(16) Card holde.... The near-point card is attached

(17) Operator's manual.... Describes instructions about handli the instrument

(18) Near-point card....Contains the near-point charts, for test the near vision

(19) Dust cover....used for dust prevention of instruments during storage

### 5. Ready use

4.1Setup connection of the device.



It shall be installed by professional personnel of the company or personnel trained and authorized by the company

It shall be used together with other supports or other combination tables, and the connectors provided by the company shall be equipped during installation. As shown in the figure, it is fixed on the support or other combined arms. After the main engine is connected, adjust the horizontal adjustment knob to adjust the main engine to maintain its level. The forehead support is facing the patient.

Installation of near vision chart



Before using the instrument, the supporting rod of the visual acuity chart, the clamp seat of the near visual acuity chart and the near visual acuity chart shall be installed first

Insert the near vision chart holder into the top of the support rod. The clamp seat shall be able to slide with the extension rod.

### Tablet application interface and function introduction

Main interface: operation program the main interface is divided into three parts: optometer operation and data display area, LCD visual acuity chart operation area and system status and menu area. The layout of the main interface and the distribution of the above three parts in the main interface are shown in the figure.



Optometer operation and data display area

The optometer operation and data display area are shown in the figure below.



The data display area of the optometer mainly has two functions.

1. Display of lens parameters of optometer

2. Realize the selection of project status and parameters of refractometer



The detailed block of the optometer is shown in the figure below:



Left eye auxiliary lens selection interface



Right eye auxiliary lens selection interface



Instrument operation

Fog inspection

1. The spherical mirror degree of the right eye is manually set at a place + 3.00d higher than the approximate degree, because the

customer's current right eye degree is -1.00d, which becomes + 2.00d after adding + 3.00

2. At this time, the visual acuity value should be lower than 0.1. Manually project a visual acuity icon, and then gradually adjust the degree to reduce the spherical lens degree by 0.25D, from + 200D to + 1.76d, and then to + 500 until his visual acuity value is about 0.5.

3. Call up the radiation vision chart radiation vision chart and ask the customer "do you have any lines that look particularly clear? If the customer answers "all look the same", it means that there is no astigmatism. After the measurement, if the customer answers "one line looks particularly clear", multiply 30 ° by the small number of the line that looks the most clear. This value is a negative correction value. For example, if the third line looks clear, it is 3x30 ° = 90 °

4. Adjust the astigmatism axis, adjust the astigmatism, transfer from 0 to -0.25, and then adjust to -0.50 until every line is clear.

5. Change the ball mirror in gear 1 of 0.25D.

The customer has myopia, and the spherical lens degree is -1.75d. After general inspection, his degree is shown in the figure below.

### Accurately check the astigmatism axis position and degree

one Manually select the point group sight mark, and the cross cylindrical mirror is automatically applied to the right eye; Or cross the cylindrical lens from the auxiliary lens, with a single line of sight, or press M1 or M2 directly into the cross column mirror. Automatically jump to a, and first accurately measure the astigmatism axis position. two Precise astigmatism axis position: press M1 or m2 to exchange both sides of the cross column mirror, so that the customer can distinguish which side to see more clearly under the payment request. If M1 is clearer, it is allowed to press the plus button above the button. If M2 is clearer, it is necessary to press the minus button above the button, and the equipment will automatically change the astigmatism axis position. After pressing, repeat the above steps by pressing M1 and M2 until they think that the clarity of both sides is almost the same, and then the accurate astigmatism can be obtained.

three Accurate astigmatism degree: after accurate astigmatism axis position, press C (-) to accurately calculate the astigmatism degree (keep the right cross column mirror state). Press M1 or m2 to change both sides of the cross column mirror, so that the customer can distinguish which side he sees more clearly. If M1 is clearer, press the plus button above the button. If M2 is clearer, press the minus button above the button, The device automatically changes the astigmatism. After pressing, repeat the above steps by pressing M1 and M2 until they think that the clarity of both sides is almost the same, and then the accurate astigmatism can be obtained. When the astigmatism degree is changed, the equipment can automatically carry out ball column linkage, that is, the column mirror is changed to  $\pm$  0.50dc, the ball mirror is automatically changed to  $\pm$  0.25ds, and the result is: -0.50dc  $\times$  100°

four After the accurate measurement of astigmatism axis position and astigmatism degree is completed, it will be removed automatically according to any visual acuity chart or s (0.25) cross cylindrical mirror

### Accurately measure whether there is any deviation in the sphericity

(red and green test is used in this method. If the red and green test is not applicable to any customer, whether there is deviation can be

determined directly according to the visual acuity)

Project red and green icons and ask the customer which word in the red and green picture can be seen more clearly. The customer says that the green is slightly clearer, which means that myopia is over corrected, so the spherical mirror degree is increased, + 0.25D. Or press the Add button directly, and the equipment will automatically complete the increase, and then ask the customer. His answer in red is a little clearer, which means that myopia is not corrected, increase -0.25d, or press the decrease button directly, and the equipment will automatically complete the decrease. It indicates that the degree of the customer should be between -1.50d and -1.75d, and whether it is high or low should be selected according to the requirements and purpose of the customer's glasses. The screen displays.

### COT- vision chart projector

### **Technical Specifications**

Visual mark	33 options
Visual beacon selection speed	Average 0.3 seconds
Cover plate	1 open 5 vertical line 5 horizontal line 21 single letter
Filter	Red / Green
Projection distance	2.7m ~ 7.0 m(standard is 5m)
Projection magnification	30x (at 5m)
Projection size	φ252mm, 330mm (W) x 225mm (H) (at 5m)
Tilt angle	Ball joint ( $\pm$ 10 degrees)
Light bulb	6V 30W (halogen), 2000 hours service life, 6V 10W LED
Power Supply	AC100~120V 50/60Hz AC200~240V 50/60Hz
Power	50VA
Auto Power Off	After 5 minutes
Overall dimension	Host: 265 (H) x 230 (W) x 360 (d)mm
	Remote control: 20 (H) x 64 (W) x 195 (d)mm
Netweight	Main engine: 6.0kg; Remote control: 180g
	1 polarized metal mesh (400mmx350mm)
Enclosure	2 spare fuses
	2 No. 7 batteries
Purchase accessories	Platen frame, wall frame, vertical frame, zoom lens,
	screen angle fixing Kit
Interface	RS232



# *vision chart projector*







# AUTO CHART PROJECTOR OPERATION HANDBOOK

# CONTENTS

INTRODUCTION		
PACKAGE CONTENTS	1	
CLASSIFCATIONS		2020
DESIGNATED SYMBOLS	2	
SAFETY INSTRUCTIONS	3	1000
INSTRUMENT COMPONENTS	4	100
INSTALLATION	5	1.10
FOCUSING	5	
OPERATION INSTRUCTIONS	6	1000
CHARTS DESCRIPTION	7	
MAINTENANCE	8	
BASIC TROUBLE SHOOTING	10	ŝ
TRANSPORTATION	10	
TECHNICAL SPECIFICATIONS	11	

# INTRODUCTION

Congratulations for purchasing of Auto Chart Projector.

In the area of subjective refraction, the Auto Chart Projector is best used for measuring visual acuity. It is fully recommended that you carefully read, understand and follow the steps in this manual in order to ensure safe operation, optimum performance and a longer service life from your new instrument. Please retain this manual for future reference.

# PACKAGE CONTENTS

Auto Chart Projector Auto Chart Projector and accessories were carefully checked and packed prior to the shipment. However, plesse check condition and contents upon delivery.

In addition to Auto Chart Projector, this shipment includes:

Polarized Metal Screen (400mm × 350mm)	1 PC
Power Cord	1 PC
Remote Control	1 PC
Spare fuses (2A,250V)	2 PCS
AAA type batteries	2 PCS

# **OPERATION HANDBOOK**

# CLASSIFICATIONS\_

[Classification under the provision of 93/42/EEC(MDD)] Class I Auto Chart Projector is classified into Class I system.

### [Protection method against electric shock] Class I

Auto Chart Projector is classified into Class I instrument.

Class I instrument in which protection against electric shock does not rely on basic insulation only, but which includes an additional safety precaution with the provision of the instrument to the protective earth conductor in fixed wiring of the installation in such a way that any accessible conductive parts cannot become live in the event of a failure of the basic insulation.

### [Degree of protection against ingress of liquids ] IPXO

Auto Chart Projector is the ordinary instrument (enclosed instrument without protection against ingress of liquids). Be careful not to splash water on the instrument.

### [Degree of protection against flammability]

Auto Chart Projector is classified as an equipment not suitable to be used in a Potentially flammable anesthetic mixture with air or with oxygen or nitrous oxide. Do not use near flammable materials.

### [Mode of operation]

Continuous operation

# DESIGNATED SYMBOLS

★	This symbol indicates that the degree of protection against Electric shock is for Type B Instrument.	-	This symbol on the main switch indicates that the power is ON.	0	This symbol on the main switch indicates that the power is OFF.
ф	This symbol indicates a fuse.	٢	This symbol indicates a protective earth.	Â	CAUTION This symbol indicates that important operating and maintenance instructions are included in this User's Guide.

# SAFETY INSTRUCTIONS

### • AT OPERATION

Never disassemble or touch the inside of the instrument. This may result in an electric shock or instrument malfunction. Never yank the power cord to disconnect from wall outlet but hold the plug while disconnecting. This can weaken the metal core of the cord and may result in a short circuit or an electric shock.

### AT STORAGE

Do not store the instrument in a place where it may get wet or where poisonous gas or liquid is stored. Avoid storing the instrument in an area with excessive heat, humidity, or dust. Recommended ranges:

Temperature range within	-10°C to 70°C
Relative humidity range within	30% to 85%
Atmospheric pressure range within	500 to 1060hPa

### AT INSTALLATION

Do not install the instrument near water. If water gets into the internal structure, there is the possibility of electrical Shock or instrument malfunction. Install the instrument in a stable and level place where vibration or shock does not occur. The instrument may not perform observation correctly or may malfunction.

Also, if the instrument is tripped over because of any accidental shock, it may result in possible injury.

### • 🗥 AFTER USE

If the instrument is not be used for a long time, disconnect the power cord from the wall outlet. Otherwise, it may cause a tire. Remove batteries from the battery case of the remote controller when the remote controller will not be used for a long time.

### AT DISPOSAL

Follow local governing ordinances and recycling plans regarding disposal or recycling of device components. Check the specified disposing method for the specific waste in advance, especially when disposing the batteries used in the remote controller. When disposing packing materials, sort them by the materials and follow local governing ordinances and recycling plans.

# INSTRUMENT COMPONENTS

- 1. Upper Case 2. Projection Lens 3. Window for remote detection 2 4. Table stand 3 5. Focusing Wheel 5 6. Pilot Lamp 7. Power Switch W. 8. RS-232C Connector 9. Fuse Holders 10. Power Connector 11. Infrared ray transmitter 12. Alphabet chart buttons 13. Number chart buttons 14. Illiterate E chart buttons 15. Program buttons 16. Lamp ON/OFF button 17. Red / Green button 18. Children chart buttons 19. Vectograph chart buttons 20. Direction buttons 10 21. Vertical, horizontal, single letter Mask buttons 9 22. Buzzer
  - 23. Reset





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- 1

# INSTAL

The Auto Chart Projector should be installed according to the following procedures: 1.If using a wall mount, it is best to locate a wall stud to support the weight of instrument. 2.Determine the refracting distance (from the patient's eye to the screen) and install the projection screen. 3.Position the instrument at the same distance from the screen to the patient within the range between 2m and 7m. To obtain longer refracting distances in small rooms, a mirror or systems of mirrors can be used in this case, a high quality front surface mirror is required.



4.Check the three-dimensional alignment of the system. The instrument is optimized when the projection screen is angled to direct light to the patient's head. Place a mirror on the screen. The light should project where the patient's head would be.



To obtain the correct focus:

1.Measure the distance from the patient to the screen (refracting distance).

2.Turn the Auto Chart Projector on, and using the remote control, project the 0.05 "E" onto the screen.

3.If using a wall mount, it is best to locate a wall stud to support the weight of instrument.

4.Adjust the position of the instrument forward or backward for sizing.

5.Adjust for sharpness and clarity by turning the focusing wheel left or right as needed.

FOCUSING

# OPERATING INSTRUCTIONS

### 1. Turn the power switch of the instrument on.

The projection lamp light and the chart for 0.05 vision shall be viewed in 3 seconds.

### 2. Changing the chart

Press the appropriate chart button the remote controller on. The charts have 33 selections

### 3. Isolating the visual acuity chart

- To use Vertical line masking: Press
   To move the position right or left: Press
   To move the position up or down: Press
   To move the position up or down: Press
- To use Horizontal line masking : Press 
   △
   ▲To move the position right or left : Press 
   ④
   ▲To move the position up or down : Press 
   △
   □
   □
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- To use Single letter masking : Press 
   To move the position right or left : Press 
   To move the position up or down : Press
- To use Red/Green filter : Press 💍
- To release masking and filter : Press any chart button



### 4. Programmed operation

I nere are two programs for selection, which are Program "P1	1" and "P2" .
Be sure to press 😇 to turn off the lamp before program sele	ection.

- To install Program "P1"
- Keep pressing the left button of O for 2~3 seconds and release the button upon hearing a beep sound;
- 2) Press the charts needed to store one by one, with a limit of at most 30 charts;
- Keep pressing the right button of of a second to store selected charts and release the button upon hearing a beep sound;
- 4) Press () to turn on the lamp and exit programming.
- To install Program "P2"
- Keep pressing the right button of of 2~3 seconds and release the button upon hearing a beep sound;
- 2) Press the charts needed to store one by one, with a limit of at most 30 charts;
- Keep pressing the left button of o
   for 2 ~ 3 seconds to store selected charts and release the button upon hearing a beep sound;
- Programmed Operation

### 1) press Ö

The first chart in the program will be viewed and the programmed operation started. (Press the button once again to switch over between P1 and P2) 2) Press the left of right button to change charts.

### **5.Reset**

Press RESET button To re-position charts to 0.05E".

### 6.Mute

Press the button it to switch over between mute and sound states.

# CHARTS DESCRIPTION





Children

•< >=

XXXXA

Antik

\*\*\*\*\*

**Cross lines** 

83

+

٠

380 08

Worth For Dot

+

**Red/Green** 

0.1

0.2

0.4

0,6

0.8

1.0

Schober



**Binocular Balance** 



**Duochrome Balance** 

50 05 60 03

**Cross Cylinder Dots** 



**Astigmatic Clock Dial** 







**Phoria With Fixation** 

1	
1	
ाः 	

### **Coincidence Vertical**

1 × 1	

**Minute Stereo** 



Fixation



# MAINTENANCE

A Risk of electric shock. Always disconnect the power cord from the wall and the instrument Prior to performing any of the following procedures.

### Maintenance

There is no periodic or routine user maintenance required.

### Cleaning

There are no cleaning requirements other than regular office housekeeping, such as dusting.

### ▲Cleaning the main body

< Projection lens >

1) Blow dusts with a blower brush.

2) If it is not clean enough, wipe with lens cleaning paper.

<Cover and Screen>

1) Wipe with a dry and soft cloth.

 If for some reason the instrument becomes soiled, wipe it with the damp lint-free cloth and mild detergent. Then, wipe off with a dry cloth to finish.

### Disassembling the upper case

1) Turn off the power and pull out the power cord.

2) Unfasten the screw in counterclockwise at the back side of case.

3) Disassamble the Upper Case by lifting it.

4) In case of reassembling, place the upper case on the main body and adjust the front position. Then, fasten the screw in clockwise.

### ▲ ● Replacement of lamp

The lamp may be HOT! Do not touch the Lamp directly and allow the sufficient amount of time to cool down the heat.

- 1) Unfasten the screw at the Lamp Cover.
- 2) Disassemble the Lamp Cover by lifting it and pull out the Lamp.



- 3) Place the new Lamp at the hole of Lamp Holder.
- 4) Turn on the power switch.
- Place the white paper in front of the projection lens in order to project the filament image of lamp.



# MAINTENANCE

6) Adjust the filament image at the center of paper with the screw at the lamp plate. (Note: Do not look into the lamp directly)



7) Place the Lamp Cover and fasten the screw.

8) Turn the power switch off.

### Replacement of Fuses

1) Turn off the power switch and disconnect the power cord.

2) Push and turn the fuse holder a quarter counterclockwise with your finger and remove it.

3) Pull out the fuse from the fuse holder and check the condition.

4) Replace the fuse with the provided spare or equivalent as specified below.

VOLTAGE	FUSE		
100V~120V	2A 120V; T2AL		
2000~2400	2A 250V; 12AL		

5) Simply return it with the fuse holder to the socket. Push and turn it a quarter clockwise to lock.

# BASIC TROUBLE SHOOTING

Note: Risk of electric shock. Always disconnect the power cord from the wall and the instrument prior to performing any of the following procedures.

### • If the instrument does not function at all:

- 1) Check the facility power source.(Circuit Breakers)
- 2) Check the electrical connections. (Power Cord)
- 3) Check the main fuse located on the rear of the unit.

### If the projector lamp dose not light:

- 1) Check the main ON/OFF Switch.
- 2) Replace the lamp with the spare lamp.

1 The lamp may be HOT! Do not touch the Lamp directly and allow the sufficient amount of time to cool down the heat.

 If the projector turns on but does not function: Check the batteries in the remote control.

# TRANSPORTATION

- 1. Make sure the instrument is disconnected from the power source.
- 2. If the instrument was installed on a custom mount, please remove the unit from the mount.
- 3. Pack the instrument in a sturdy carton with suitable packing materials.
- 4. After packing, do not hurl and impact cause of breakable goods.

Do not use this machine near by other machines or electric bulb (ex.Halogen lamp), it will improperly operate on the effect of electronic waves.

## TECHNICAL SPECIFICATIONS

Chart	33 Selections						
Chart selection speed	Average 0.3 sec.						
Mask	1 open	5 horizonta	l lines	5 vertical lines	21 single letters		
Filter	Red / Green						
Projection distance	2.7m ~ 7.0m(5m is standard)						
Projection magnification	30X (at 5m)						
Projection size	<b>φ</b> 252mm and 330mm(W) × 225mm(H) (at 5m )						
Tilt angle	Ball joint (±10 degree)						
Lamp	8V 5W LED/6V 20W (Halogen), 2,000 hrs lifetime						
Power source	AC 90-240V 50/60HZ						
Power consumption	50VA						
Auto-OFF function	after 5 minutes						
Dimensions	Main body: 265(H) × 230(W) × 360(D)mm						
	Remote controller: $20(H) \times 64(W) \times 195(D)mm$						
Net weight	Main body: 6.0 Kg Remote controller: 180g						
Accessories	Polarized Meta	al Screen (400m	m × 350mm )	1 PC			
	Spare fuses		2 PCS				
	AA A size batte	ery		2 PCS			
Optional accessories	Table stand	wall bracket	floor stand	variable focus lens	screen angle fixing set		
Interface	RS232						

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# **FDA Registration Confirmation**

Manufacturer: SHANGHAI TOP VIEW INDUSTRIAL CO.,LTDAddress: No.3388 GongHeXin Road,JingAn Shanghai , CN 200436

US Agent:Regrek LLCAddress:19 Holly Cove Ln., Dover, Delaware, 19901, UNITED STATES

#### The facility registration and device listing information:

Registration Number:3012707847Owner/Operator Number: 10059986Device Listing#: See annex

This attestation does not denote endorsement or approval of the attestation-holder's device or establishment by the U.S. Food and Drug Administration.

Pursuant to 21 CFR 807.39, "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products. Any representation that creates an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding." The U.S. Food and Drug Administration does not issue a attestation of registration, nor does the U.S. Food and Drug Administration recognize a attestation of registration.

Authorized Signature(s)

Signature: Place: China/ Shanghai Expiration Date: Dec. 31. 2021



# **FDA Registration Confirmation**

#### ANNEX:

Listing No	Code	Device Name					
D345115	нко	Refractometer, ophthalmic (TPV800 handheld refractor; TPV series auto refractometer)					
D345116	HKN	Refractor, manual, non-powered, including phoropter (phoropter)					
D345117	HKZ	Sterilizer, tonometer (TPV500 tonometer)					
D345118	PJZ	Camera, ophthalmic, general-use (Hand-held Fundus Camera; TPV series Digital Fundus Camera)					
D345119	PUE	Biomicroscope, slit-lamp, AC-powered, exempt (TPV series slit lamp)					
D345120	HRJ	Table, instrument, powered, ophthalmic(Ophthalmic unit)					
D345121	HLM	Instrument, measuring, lens, ac-powered (TPV series lens meter)					
D345122	HPA	Frame, trial, ophthalmic (trial frame)					
D345123	HPC	Set, lens, trial, ophthalmic (trial lens set)					
D345124	НОХ	Chart, visual acuity (visual charter)					
D345125	HPT	Perimeter, automatic, ac-powered (perimeter)					
D345341	HOS	Projector, ophthalmic (projector charter)					
D345342	HLH	Pupillometer, manual (pupillometer)					
D345343	HRM	MICROSCOPE, OPERATING & ACCESSORIES, AC-POWERED, OPHTHALMIC (operating microscope)					

D245244	W7A	DEVICE, VEIN LOCATION, LIQUID CRYSTAL
D345344	KZA	(vein finder)
D266902	шт	Ophthalmoscope, battery-powered
D300893	HLJ	(ophthalmoscope retinoscope)
D266804		INSTRUMENT, MANUAL, SURGICAL, GENERAL USE
D300894	MDM	(ophthalmic surgical instrument)
D266905	MMO	Topographer, corneal, ac-powered
D300895	MMQ	(cornealtopographer)

END OF THE ANNEX

CED Authorized Signature(s)

Signature:

Place: China/ Shanghai

Expiration Date: Dec. 31. 2021

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# Assessment Report

Presented to

# Shanghai Top View Industrial Co., Ltd.

#### 上海拓扑威实业有限公司

Gold Supplier & Assessed Company	Self-owned Wholly Owned Shareholder/Partner					
Relationship:	Cooperation Partner					
Compony Address	Room 906, Yongding Plaza, No. 3388, Gonghe New Road,					
	Jingan District, Shanghai, China					
City / Country:	Shanghai / China					
Consigner of Assessment:	Alibaba					
Gold Supplier Member ID:	shtopview					
Gold Supplier Company Name:	Shanghai Top View Industrial Co., Ltd.					
Contact Person:	Mr. Tao Shangguan					
Phone Number:	0086-21-61178155					
Fax Number:	N/A					
Email:	info@shtopview.com					
Website Address (URL):	https://shtopview.en.alibaba.com					

Service Provided by Intertek Report No.: 22069995\_T







Assessment Report

Report Number:	22069995_T	Assessment Type	Trading Assessment
Date of Assessment:	31/Aug./2021	Report Date:	31/Aug./2021
Assessor's Name:	Zad Zhang	Validity Period:	01/Sep./2021 - 31/Aug./2022
Reviewed By:	Mack Long	Online Verification: https://www.intertek.com.cn/ASVService	
			ome/IndexCN

#### Important Notes:

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Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
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**Assessment Report** 

#### Contents

Section 1: Company Over	view					5
1.1 Legal Validity						5
Section 2: Human Resour	ces					6
2.1 Company Chart						6
2.2 Employee Headcount						6
2.3 Management						6
Section 3: Current Export	Situation					7
Section 4: Export Busines	ss Capacity	y				9
4.1 Market Distribution (P	revious 12 I	Months)				9
4.2 Main Clients						10
Section 5: Quality Assura	nce					11
5.1.1 Quality Management	System Ce	rtification				11
5.1.2 Product Certification	1					11
5.2 Supplier Management						11
5.3 After Sales Service						12
5.4 Overseas After Sales	Service					12
Section 6: R & D Capacity						13
6.1 Current Situation						13
6.2 R&D Real Case Descri	ption					14
6.3 Design Process						15
6.4 Design Devices						15
Section 7: Company Deve	elopment /	Expansion Pla	ns			16
Section 8: Certification &	Photos					17
Section 9: Company and	Product Sa	amples				19
Section 10: Competitive A	dvantages	S				21
10.1 Product Group Capa	- city					21
Report No: 22069995 T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIAL	All Rights Res	erved			Page No:	3 of 25



1	0.2 Real Case for Lower MOQ & Lead Time	.21
1	0.3 Real Case for Large Contract	.22
1	0.4 Overseas after-sales service capacity	.22
1	0.5 After-sales service capacity	.22
See	ction 11: Service Capabilities	24
1	1.1 Experience with Large-scale Procurement Contracts	.24
1	1.2 Overseas Showroom	.24
1	1.3 Offline Trade Show	.24
Se	ction 12: Supply Chain Capability	25
1	2.1 Centralized Procurement	.25
1	2.2 Overseas Warehouse	.25

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIAL All Rights Reserve			served			Page No:	4 of 25



**Assessment Report** 

#### Section 1: Company Overview

Company Overview										
1.1 Legal Validity										
Does the company have a		Business License	9131011506602810							
valid business license?		Number:	2F							
Vear Established:	18/Apr /2013	Validity Period:	18/Apr./2013-							
	10/Api./2013		17/Apr./2033							
Export Experience	8 Vears	Industry	8 Years							
		Experience:								
Registered Address	Room 150, Building 3, No. 2558,	Zhouzhu Road, Pudo	ng New Area,							
Registered Address.	Shanghai, China									
Company Address	Room 906, Yongding Plaza, No.	3388, Gonghe New Ro	oad, Jingan District,							
	Shanghai, China									
Is It Listed Company?	Ves 🕅 No	Company Stock	N/A							
		Code:								
Annual review conducted by		Reviewed By:	Market Supervision							
the Industrial & Commercial	X Yes I No		Administration							
Bureau?			Bureau of Pudong							
			New Area, Shanghai							
Registered Capital:	RMB 2,000,000									
Corporate Representative:	Mr. Tao Shangguan									
Industry:	Optical Instrument									
Business Type:	☐ Manufacturer ⊠Trading Company ☐Manufacturer & Trading Company									
Tupo of Ownorship:	Private Owner Public Company Joint Venture									
Type of Ownership.	Sole Proprietorship Other									
Draduata (Camiaa)	Auto Lens Meter, Slit Lamp, 3D Auto Lens Edger, Rebound Tonometer, Auto									
Products /Service:	Refractometer									
1.2 Company Building Information	on line line line line line line line lin									
Certification Type:										
Land Certification	al Estate Certification 🛛 🛛 Leas	e Agreement 🛛 🗌 F	actory Officer Claimed							
Total Building Size: 326	m2									
Office Size: 326	m2									

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	IDENTIAL All Rights Reserved					Page No:	5 of 25



**Assessment Report** 

#### **Section 2: Human Resources**



Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIAL All Rights Reserved						Page No:	6 of 25



**Assessment Report** 

# **Section 3: Current Export Situation**

Cu	rrent Export Situation								
The	ere are <u>8</u> fore	eign trading emp	loyee(s)	in th	e company.				
	Working Exporionco	Hoadcount			Accepted	Listenin	g &	Readin	g &
		Headcount			Language	Speaki	ng	Writir	ng
	Over 30 Years	0		Enç	glish	Yes		Yes	
	21-30Years	0		N/A	١	N/A		N/A	
	11-20 Years	0		N/A	N	N/A		N/A	
	6-10 Years	0		N/A	N	N/A		N/A	
	2-5 Years	8		N/A	N	N/A		N/A	
	Less than 2 years	0		N/A	۱	N/A		N/A	
Do	es the company have a va	alid export licens	e?		🛛 Yes	🗌 No			
Ex	oort License Registration I	No.:			01314518				
Tot	al Revenue (Previous Yea	ır, USD):			Confidential				
Tot	al Export Revenue (Previo	ous Year, USD):			с				
Est	imated Export Revenue (	Current Year, US	SD):		Confidential				
Tra	de Agents Employed Ove	rseas:			🗌 Yes	🛛 No			
Ne	arest Port:				Shanghai Por	rt			
Ace	cepted Payment Terms				S FOB		ØΕΧ	W	
					🖾 L/C	🛛 Т/Т		🛛 Credit	Card
Ace	cepted Payment Type:				🖾 Cash 🛛 West Union 🖾 Money Gram				
					🛛 Paypal 🛛 Moneybookers				
Ave	erage lead time from prod	uct order confirn	nation to	proc	duction delivery (products exiting the factory):				
	Product C	ategory			Num			Unit	
	Auto Lens	s Meter			3 Day			Days	
	Slit La	imp			3			Days	
	3D Auto Le	ns Edger			3			Days	
	Rebound To	onometer			3			Days	
Auto Refractometer				3 Days					
Ave	erage Sampling Time				1				
	Product Category					Lead	Time		
	Auto Lens	s Meter				3 Da	ays		
	Slit La	imp				3 Da	ays		
	3D Auto Le	ns Edger				3 Da	ays		

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
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Report No.: 22069995\_T

Rebound Tonometer	3 Days
Auto Refractometer	3 Days
The Shortest Sampling Time	
Product Category	Shortest Lead Time
Auto Lens Meter	3 Days
Slit Lamp	3 Days
3D Auto Lens Edger	3 Days
Rebound Tonometer	3 Days
Auto Refractometer	3 Days

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	AL.	All Rights Res	served			Page No:	8 of 25



Report No.: 22069995\_T

**Assessment Report** 

# Section 4: Export Business Capacity

Export Business Capacity							
4.1 Market Distribution	(Previous 12 Months)						
Market	Main Product(s)	Revenue (USD)	Total Revenue (%)				
	Auto Lens Meter, Slit Lamp, 3D Auto						
North America	Lens Edger, Rebound Tonometer, Auto	Confidential	5				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
South America	Lens Edger, Rebound Tonometer, Auto	Confidential	50				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
Eastern Europe	Lens Edger, Rebound Tonometer, Auto	Confidential	2				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
Southeast Asia	Lens Edger, Rebound Tonometer, Auto	Confidential	5				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
Africa	Lens Edger, Rebound Tonometer, Auto	Confidential	5				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
Oceania	Lens Edger, Rebound Tonometer, Auto	Confidential	2				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
Mid East	Lens Edger, Rebound Tonometer, Auto	Confidential	5				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
Eastern Asia	Lens Edger, Rebound Tonometer, Auto	Confidential	5				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
Western Europe	Lens Edger, Rebound Tonometer, Auto	Confidential	10				
	Refractometer						
	Auto Lens Meter, Slit Lamp, 3D Auto						
Central America	Lens Edger, Rebound Tonometer, Auto	Confidential	2				
	Refractometer						

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	AL.	All Rights Res	served			Page No:	9 of 25



	Auto L	ens Meter, Slit Lamp, 3D Auto				
Northern Europe	Lens E	dger, Rebound Tonometer, Auto	С	onfidential	2	
		Refractometer				
	Auto L	ens Meter, Slit Lamp, 3D Auto				
Southern Europe	Lens E	dger, Rebound Tonometer, Auto	Confidential		2	
	Refractometer					
	Auto Lens Meter, Slit Lamp, 3D Auto					
South Asia	Lens E	dger, Rebound Tonometer, Auto	Confidential		5	
	Refractometer					
4.2 Main Clients						
Client Name		Main Product(s)		Total Revenue (%)		
Confidential	Confidential		Confidential			

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	L	All Rights Re	served			Page No:	10 of 25



**Assessment Report** 

# Section 5: Quality Assurance

5.1.1 Quality Management System Certification         Certification         Certificate By         Certificate No.         Business Scope         Validity Date           N/A         N/A         N/A         N/A         N/A         N/A           5.1.2 Product Certification         Certificate No.         Product Name & Model No.         Validity Date           N/A         N/A         N/A         N/A         N/A           Cartification         Certificate No.         Product Name & Model No.         Validity Date           N/A         N/A         N/A         N/A         N/A           S.2 Supplier Management         Content         Observations /Comments         Secondary           5.2.1         Does the company have a supplier assessment procedure?         No         No         Secondary           5.2.2         updated list of approved suppliers?         No         Secondard         Secondard           freeded with a procedure for purchasing contract review and approval?         No         Secondard         Secondard           5.2.4         Does the company keep its supplier assessment reports?         Yes, assessment reports are available for more than 3 years         Yes, assessment reports are available for the last 1-3 years           5.2.5         Are the company's purchasing documents sufficient to ensur- product safety control and th	Quality Assu	irance	<del>)</del>							
Certification         Certified By         Certificate No.         Business Scope         Validity Date           N/A         N/A         N/A         N/A         N/A         N/A           5.12 Product Certification         Certified By         Certificate No.         Product Name & Model No.         Validity Date           N/A         N/A         N/A         N/A         N/A         N/A           5.2 Supplier Management         Does the company have a supplier assessment procedure?         No         Yes         No           5.2.1         Does the company have a supplier?         Yes         No         No         No           5.2.2         updated list of approved supplier?         No         No         No         Secondard procedures           5.2.3         Has the company established and procedure for purchasing contract review and approval?         No         No         No           5.2.4         Does the company keep its supplier assessment reports?         Yes, with written procedure but lack of consistent standard procedure for purchasing contract review and approval?         No         No           5.2.4         Does the company keep its supplier assessment reports?         Yes, assessment reports are available for the previous 12 months         Yes, assessment reports are available for the previous 12 months           5.2.5.         Are	5.1.1 Quality	Mana	igement S	ystem Ce	rtification					
N/A         N/A         N/A         N/A         N/A           State of the company have a supplier assessment procedure?         N/A         N/A         N/A         N/A         N/A           5.2.1         Does the company have a supplier assessment procedure?         No         State of the company have a supplier assessment procedure?         No         State of the company have a supplier assessment procedure?         No           5.2.2         Does the company have a supplier assessment procedure?         No         No         State of the company have a supplier assessment procedure?         No           5.2.2         Does the company have a supplier assessment procedure?         No         No         State of the company have a supplier assessment procedure?         No           5.2.2         Does the company have an updated list of approved suppliers?         No         No         State of the company established and procedure for purchasing contract review and approval?         Yes, with written procedure but lack of consistent standard review and approval?         No           5.2.4         Does the company keep its supplier assessment reports?         Yes, assessment reports are available for the last 1-3 years         Yes, assessment reports are available for the previous 12 months           5.2.5         Are the company's purchasing document sufficient to ensure product safety control and their customers' requirements?         Yes, the purchasing document includes all	Certificatio	n	Certifie	d By	Certificat	e No.	Busir	ess Scope	,	Validity Date
5.1.2 Product Certification       Certificate No.       Product Name & Model No.       Validity Date         N/A       N/A       N/A       N/A       N/A       N/A         Supplier Management         Item       Content       Observations /Comments         5.2.1       Does the company have a supplier assessment procedure?       No       No       Second and a standard procedures       No         5.2.2       Does the company established and implemented a standard procedure for purchasing contract review and approval?       Yes, with written standard procedure but lack of consistent standard         5.2.3       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for more than 3 years         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.5       Are the company's purchasing document includes all the information required       Yes, however the purchasing document includes all the information         5.2.6       Is there a procedure to conduct random product inspection and their customers' requirements?       Yes, with clear standard and written inspection records         5.2.6       Is there a procedure to conduct random product inspection safet       Yes, with inspection records but no procedures         Yes, with inspection records but no procedures       Yes, with	N/A		N/	A	N/	I/A N/A N/A				
Certification       Certificate No.       Product Name & Model No.       Validity Date         N/A       N/A       N/A       N/A       N/A         S.2 Supplier Maragement       Content       Observations /Comments         5.2.1       Does the company have a supplier assessment procedure?       No       No         5.2.1       Does the company have a supplier?       Yes       Secondary       Secondary         5.2.2       Does the company have a supplier?       Yes       Secondary	5.1.2 Produc	t Cert	ification							
N/A     N/A     N/A     N/A       5.2 Supplier Management       Item     Content     Observations /Comments       5.2.1     Does the company have a supplier assessment procedure?     No       5.2.1     Does the company have a updated list of approved updat	Certificatio	on	Certifie	d By	Certificat	e No.	Product N	ame & Mode	l No.	Validity Date
5.2 Supplier Management           Item         Content         Observations /Comments           5.2.1         Does the company have a supplier assessment procedure?         No           5.2.2         Does the company have an updated list of approved suppliers?         No           5.2.3         Has the company established and implemented a standard procedure for purchasing contract review and approval?         Yes, with written standard procedures           5.2.4         Does the company keep its supplier assessment reports?         No           5.2.4         Does the company keep its supplier assessment reports?         No           5.2.4         Does the company keep its supplier assessment reports?         Yes, assessment reports are available for more than 3 years           5.2.4         Does the company keep its supplier assessment reports?         Yes, assessment reports are available for the last 1-3 years           5.2.4         Does the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?         Yes, the purchasing document includes all the information required           Yes, however the purchasing document includes incomplete information         No           5.2.6         Is there a procedure to conduct radom product inspection safter final packaging?         Yes, with inspection records but no procedures Yes, with inspection records but no procedures No, inspections are not necessary	N/A N/A M					A		N/A		N/A
Item         Content         Observations /Comments           5.2.1         Does the company have a supplier assessment procedure?         No           5.2.2         Does the company have an updated list of approved suppliers?         No           5.2.3         Has the company established and implemented a standard procedure for purchasing contract review and approval?         Yes, with written standard procedures           5.2.4         Does the company keep its supplier assessment reports?         Yes, assessment reports are available for more than 3 years           5.2.4         Does the company keep its supplier assessment reports?         No           5.2.5         Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?         Yes, the purchasing document includes all the information customers' requirements?           5.2.6         Is there a procedure to conduct random product inspections after final packaging?         Yes, with clear standard and written inspection records incomplete information           Yes, with procedures but no inspection records         Yes, with procedures but no inspection records in onspection records but no procedures	5.2 Supplier	Mana	gement							
5.2.1       Does the company have a supplier assessment procedure?       No         5.2.2       Does the company have an updated list of approved suppliers?       No         5.2.3       Has the company established and procedure for purchasing contract review and approval?       Yes, with written standard procedures         5.2.3       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for more than 3 years         5.2.4       Does the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Yes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Yes, with clear standard and written inspection records but no procedures         9       Yes, with procedures but no inspection records but no procedures       No	ltem		Co	ontent			Obse	ervations /Co	omments	
5.2.1       supplier assessment procedure?       No         5.2.2       Does the company have an updated list of approved suppliers?       No         5.2.3       Has the company established and implemented a standard procedure for purchasing contract review and approval?       Yes, with written standard procedure but lack of consistent standard         5.2.4       Does the company keep its supplier assessment reports?       No         5.2.4       Does the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Yes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       No         8       Start in a packaging?       No	5.0.1	Does	s the compa	any have a	a	🛛 Yes				
5.2.2       Does the company have an updated list of approved suppliers?       No         5.2.3       Has the company established and implemented a standard procedure for purchasing contract review and approval?       Yes, with written standard procedure but lack of consistent standard         5.2.3       Does the company keep its supplier assessment reports?       No         5.2.4       Does the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Yes, subplier assessment reports are available for the previous 12 months         5.2.6       Are the company's purchasing document includes all the information required       Yes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Yes, with clear standard and written inspection records but no procedures         Yes, with procedures but no inspection records       Yes, with procedures but no inspection records	J.Z. I	supp	lier assess	ment proc	edure?	🗌 No				
5.2.2       updated list of approved suppliers?       No         3.2.3       Has the company established and implemented a standard procedure for purchasing contract review and approval?       Yes, with written standard procedures         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for more than 3 years         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.4       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Yes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Yes, with clear standard and written inspection records No, inspections are not necessary		Does	s the compa	any have a	an	🖂 Yes	i			
suppliers?         4as the company established and implemented a standard procedure for purchasing contract review and approval?       Yes, with written standard procedure but lack of consistent standard         5.2.3       Does the company keep its supplier assessment reports?       No         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.4       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Yes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Yes, with recent standard and written inspection records         Yes, with written standard procedures       Yes, with written procedures but no inspection records         No       Yes, with vertice product safety control and their customers' requirements?         No       Yes, with clear standard and written inspection records         No       Yes, with procedures but no inspection records         Yes, with procedures but no inspection records       No, inspections are not necessary	5.2.2	upda	ated list of a	approved		🗌 No				
5.2.3       Has the company established and implemented a standard procedure for purchasing contract review and approval?       Yes, with written procedure but lack of consistent standard         5.2.3       Does the company keep its supplier assessment reports?       No         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the previous 12 months         5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Yes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Yes, with no procedures but no inspection records but no procedures         Yes, with procedures but no inspection records       No       Yes, with procedures but no inspection records		supp	liers?							
5.2.3       implemented a standard procedure for purchasing contract review and approval? <pre>Yes, with written procedure but lack of consistent standard         <pre>Standard</pre>       No         5.2.4       Does the company keep its supplier assessment reports?              Yes, assessment reports are available for the last 1-3 years             Yes, assessment reports are available for the previous 12 months             No         5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?              Yes, the purchasing document includes all the information required             Yes, however the purchasing document includes incomplete information             No         5.2.6       Is there a procedure to conduct random product inspections after final packaging?              Yes, with clear standard and written inspection records             No, inspections are not necessary             Yes, with procedures but no inspection records             No, inspections are not necessary</pre>		Has	the compai	ny establis	shed and	🖂 Yes	, with written s	tandard proc	edures	
5.2.3       procedure for purchasing contract review and approval?       standard         No       No         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for more than 3 years         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Xes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Xes, with clear standard and written inspection records Yes, with procedures but no inspection records         Report No:       22069995 T       Report date: 31/Aug/2021       Assessed By       Zad Zhang	523	imple	implemented a standard			🗌 Yes	, with written p	rocedure but	lack of co	nsistent
review and approval?       No         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Yes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Yes, with clear standard and written inspection records         Yes, with procedures but no inspection records       Yes, with procedures but no inspection records	5.2.5	proc	procedure for purchasing contract			standar	ď			
5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.4       Does the company keep its supplier assessment reports?       Yes, assessment reports are available for the last 1-3 years         5.2.4       Yes, assessment reports are available for the last 1-3 years         Yes, assessment reports are available for the previous 12 months         No         5.2.5         Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?         Yes, however the purchasing document includes all the information required         Yes, however the purchasing document includes incomplete information         No         S.2.6       Is there a procedure to conduct random product inspections after final packaging?         Yes, with clear standard and written inspection records         Yes, with procedures but no inspection records         Yes, with procedures but no inspection records         Yes, with procedures but no inspection records         No, inspections are not necessary		revie	w and app	roval?		🗌 No				
5.2.4       Does the company keep its supplier assessment reports?       years         Solution       Yes, assessment reports are available for the last 1-3 years         Yes, assessment reports are available for the previous 12 months         No         Solution						🖂 Yes	, assessment i	reports are av	vailable for	more than 3
5.2.4       Does the company keep its supplier assessment reports?          Yes, assessment reports are available for the last 1-3 years         Yes, assessment reports are available for the previous         12 months         No         Yes, the purchasing document includes all the         information required         Yes, however the purchasing document includes         incomplete information         Yes, with clear standard and written inspection records         Yes, with inspection records but no procedures         Yes, with procedures but no inspection records         Yes, with procedures but no inspection records         Yes, with procedures but no inspection records         No, inspections are not necessary         Z2069995 T         Report No:         Z2069995 T         Report No:         Z2069995 T         Report date:         31/Aug/2021         Assessed By         Zad Zhang						years				
5.2.4       Does the company keep its supplier assessment reports?       years         Supplier assessment reports?       Yes, assessment reports are available for the previous 12 months         No       No         5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Yes, the purchasing document includes all the information required         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Yes, with clear standard and written inspection records but no procedures         Yes, with procedures but no inspection records       Yes, with procedures but no inspection records		Does	s the comp	any kaon i	ite	Yes, assessment reports are available for the last 1-3				
5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?	5.2.4	SUDD	lier seesee	ment rend	nte?	years				
5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       Image: Stress of the stress		Supp	nei 833633	mentrept	113:	☐ Yes, assessment reports are available for the previous				
5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?       No         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       No         Seport No:       22069995 T       Report date:       31/Aug/2021						12 months				
5.2.5       Are the company's purchasing documents sufficient to ensure product safety control and their customers' requirements?              \[                  Yes, however the purchasing document includes incomplete information						🗌 No				
5.2.5       Are the company's parchasing documents sufficient to ensure product safety control and their customers' requirements?       information required Yes, however the purchasing document includes incomplete information No         5.2.6       Is there a procedure to conduct random product inspections after final packaging?       No         8eport No:       22069995 T       Report date:       31/Aug/2021       Assessed By       Zad Zhang		Δre t	he compan	w's nurch	asina	🛛 Yes	, the purchasir	ng document	includes a	ll the
5.2.5       accountent of current of		docu	iments suffi	icient to e	nsure	information required				
5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Is there a procedure to conduct inspections after final packaging?       Is there a procedure to conduct inspections after final packaging?       Is there a procedure to conduct inspection records but no procedures incomplete information         8       Yes, with clear standard and written inspection records but no procedures incomplete information         9       Yes, with inspection records but no procedures         10       Yes, with procedures but no inspection records         10       No, inspections are not necessary         11       Report No:       22069995 T         12       Report date:       31/Aug/2021	5.2.5	nrod	uct safety c	control and	their	Yes, however the purchasing document includes				
5.2.6       Is there a procedure to conduct random product inspections after final packaging?       No         Report No:       22069995 T       Report date:       31/Aug/2021       Assessed By       Zad Zhang		custo	omers' real	uirements?	>	incomplete information				
5.2.6       Is there a procedure to conduct random product inspections after final packaging?       Yes, with clear standard and written inspection records         Yes, with inspection records but no procedures       Yes, with inspection records but no procedures         No, inspections are not necessary		0400				🗌 No				
5.2.6       random product inspections after final packaging?		Is the	ere a proce	dure to co	onduct	🛛 Yes	, with clear sta	ndard and w	ritten inspe	ection records
Image: Section products and products an	5.2.6	rand	om product	t inspectio	ns after	Yes, with inspection records but no procedures				
Report No:     22069995 T     Report date:     31/Aug./2021     Assessed By     Zad Zhang		final	packaging	?		🗌 Yes	, with procedu	res but no ins	spection re	cords
Report No: 22069995 T Report date: 31/Aug/2021 Assessed By Zad Zhang	,					🗌 No,	inspections ar	e not necess	ary	
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Report No.: 22069995\_T

5.3 After Sal	es Service	
Item	Content	Observations /Comments
5.3.1	Is customer feedback, including complaints, clearly recorded and maintained?	<ul> <li>Yes, with a standard feedback form and records</li> <li>Yes, with a standard feedback form but no records</li> <li>Yes, with records but no standard feedback form</li> <li>No</li> </ul>
5.3.2	Are there any clear procedures for handling customer complaints?	<ul> <li>Yes, with clear procedures and written records</li> <li>Yes, with clear procedures but no written records</li> <li>Yes, with written records but no clear procedures</li> <li>No</li> </ul>
5.3.3	Is there a closed-loop corrective action system in place?	⊠ Yes □ No
5.3.4	Can finished/packaged products be traced by lot identification to the appropriate raw material test reports?	<ul> <li>Yes, with procedures to trace raw materials</li> <li>Yes, main raw material can be traced</li> <li>No, only the production date can be traced</li> <li>No</li> </ul>
5.3.5	Is there a product alert and recall procedure?	⊠ Yes □ No
5.3.6	Do you have a complete after sales service capability?	⊠ Yes □ No
5.4 Overseas	s After Sales Service	
5.4.1	Is it possible to provide expatriate engineer services?	⊠ Yes □ No
5.4.2	If possible provide expatriate engineer,what kind of onsite after-sales services are included?	🛛 Debugging 🖾 Maintain 🖾 Repair 🔲 Other

Report No: 22069995_T	Report date: 31/Aug./2021 Assessed By	Zad Zhang
CONFIDENTIAL	All Rights Reserved	Page No: 12 of 25



Assessment Report

#### Section 6: R & D Capacity

R&D Capacity								
6.1 Current Situat	ion							
There is	There is   0   R&D engineer(s) in the company.							
Education Level	Headcount			Work	Experience		Headcount	
Doctorate	0			Over 3	0 Years		0	
Post-Graduate	0			21-30	/ears		0	
Graduate	0			11-20 \	⁄ears		0	
Junior College	0			6-10 Ye	ears		0	
Technical School	0			2-5 Yea	ars		0	
High School	0			Less th	an 2 years		0	
Patent Situation								
Patent No.	The Name of th	e Patent		Th	e Patent Type		Available Date	
N/A	N/A				N/A		N/A	
Brand Situation								
Registration/ap plication No.	Brand Name	For Appr Go	For Approval to Us Goods		Jse Validity Date		Ref.	
27928760	Refer to Photo	Cate	norv 9	gory: 9 14/Nov./2018- 13/Nov./2028		Pł	Photo in Section 8	
			.goi j. c			(Tr	ademark Photos)	
The Average Time	For New Products La	aunched	1					
Pr	oduct Category		Num				Unit	
	N/A		N/A				N/A	
The Shortest Time	e For A New Item Lau	nched	1					
Pr	oduct Category			Nu	m		Unit	
	N/A			N/	A		N/A	
Does the company	provide ODM service f	for others?	☐ Yes ⊠ No					
Are there relevant	design input/output, rev	/iew, and		20				
verification documents available for the assessment				0				
company?				0				
Based on inspection with adequate spect	n, are R & D employee cialized equipment?	s equipped	🗌 Ye   🖾 N	es o				
If yes, please list a	II key equipment used:		N/A					

Repo	ort No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CON	IFIDENTIA	L	All Rights Re	served			Page No:	13 of 25



		1				
Do R& D employees use any s designing new products?	pecific software for	☐ Yes ⊠ No				
If yes, please list the main soft	ware used:	N/A				
Please list all certifications and R & D department:	/or qualifications of the	N/A				
Has the company established s procedures for new products?	standard design	<ul> <li>☐ Yes, with clear</li> <li>☐ Yes, without w</li> <li>☑ No</li> </ul>	r written in ritten instr	structions uctions		
Have the designed products be validated?	een internal verified or	<ul> <li>Yes, with clear</li> <li>Yes, only part</li> <li>Yes, without w</li> <li>No</li> </ul>	<sup>-</sup> written re written rec rritten reco	cords cords ords		
Have the designed products be third-party inspection body?	een tested by a	<ul><li>☐ Yes, all design</li><li>☐ Yes, only part</li><li>☑ No</li></ul>	ed produc	ts have been d products ha	tested we been tested	
Are the designed products con customers?	firmed by the	<ul> <li>Yes, all design</li> <li>Yes, part of de according to client</li> <li>No</li> </ul>	ed produc signed pro t's requirer	ts have been oducts have b ments	confirmed een confirmed	
Does the company has qualific designers?	ation requirements for	<ul> <li>Yes, with writte</li> <li>Yes, without w</li> <li>No, but at leas</li> <li>needed</li> <li>No</li> </ul>	en job des ritten job c st two year	cription description s design expe	erience is	
Are the designers' qualification company?	s recognized by the	<ul> <li>☐ Yes, with writte</li> <li>☐ Yes, without w</li> <li>☑ No</li> </ul>	en records rritten reco	rds		
What level of design services a	are provided?	<ul> <li>Only add logo/change color/material</li> <li>Sample processing</li> <li>Graphic processing</li> <li>Create an entirely new product</li> </ul>				
New Products Launched qua	intity for each year	L				
Product Cate	gory	Num		ι	Jnit	
N/A		N/A		1	N/A	
6.2 R&D Real Case Description	on					
Customer's Name		N/A				
Report No:     22069995_T       CONFIDENTIAL	Report date: 31/Aug./2021 All Rights Reserved	Assessed By	Zad Zhang	Page No:	14 of 25	



Customer's Location N/A				
Customer's Industry:	N/A			
Order's Requirement Description:		N/A		
6.3 Design Process				
Process 1 Pro		ocess 2	Process 3	
N/A		N/A	N/A	
Description: N/A	Desci	ription: N/A	Description: N/A	
6.4 Design Devices				
Device 1	D	evice 2	Device 3	
N/A		N/A	N/A	

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	NL.	All Rights Res	served			Page No:	15 of 25



**Assessment Report** 

# Section 7: Company Development / Expansion Plans

Company Development / Expansion Plans							
ltem	n Company Development Action Timeframe						
1	The organization is going to develop more markets.	1 Year					

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	L	All Rights Res	served			Page No:	16 of 25



Report No.: 22069995 T

**Assessment Report** 

#### Section 8: Certification & Photos



Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	AL.	All Rights Re	served			Page No:	17 of 25



www 中 年 人 民 共 和 国 海 美 进出口货物被发货人根关注册登记证书 MARSHARE MINAKONS MARSHARE MINAKONS MARSHARE MINAKONS MI	上市市市市市市市市市市市市市市市市市市市市市市市市市市市市市市市市市市市市	N/A
Patent Photos		
Certification & Photos	–N/A	Certification & Photos –N/A
Droduct Contification D		N/A
Cortification & Photos	10105	Cartification & Photos
	 N/A	N/A
Quality Management S	vstem Certification Photos	
Certification & Photos		Certification & Photos
	N/A	N/A
Trademark Photos		
Certification & Photos	Tardemark	Certification & Photos N/A
日本日期 2018年11月1日 有次日 日本日期 2018年11月1日 有次日 日本日期 2018年11月1日 有次日 日本日期 2018年11月1日 有次日 日本日期 2018年11月1日 有次日 日本日期 2018年11月1日 有次日 日本日期 2018年11月1日 有次日	<ul> <li>○ 注 川丁 正</li> <li>A 27928760 号</li> <li>● 注 川丁 正</li> <li>HTOPVIEW</li> <li>A 70286, MERCES, 电测定器, 用量设备, 并明计.</li> <li>I 25588号 端150定</li> <li>A 2028年11月13日 支 証 机关</li> <li>● ジェンスの第二1月13日 支 証 机关</li> </ul>	N/A

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	NL.	All Rights Re	served			Page No:	18 of 25



**Assessment Report** 

#### **Section 9: Company and Product Samples**



Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	L	All Rights Res	served			Page No:	19 of 25



**Assessment Report** 

Report No.: 22069995\_T



Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	AL.	All Rights Re	served			Page No:	20 of 25



Assessment Report

# Section 10: Competitive Advantages

10.1 Product Group Capacity				
10.1.1 Products Sold (Within12 I	Months)			
Products Name	Quantity		Revenue (USD)	
Auto Lens Meter	400 Sets		Confidential	
Slit Lamp	150 Sets		Confidential	
3D Auto Lens Edger	70 Sets		Confidential	
Rebound Tonometer	300 Sets		Confidential	
Auto Refractometer	500 Sets		Confidential	
10.1.2 Suppliers Cooperated Wit	h (Within12 Months)	•		
No. of cooperation suppliers (total)		Confidential		
No. of suppliers (which cooperated	l over 2 times)	Confidential		
No. of provinces which cooperation	n suppliers belong to	Confidential		
Would the company like to provide	design solution	🛛 Yes		
service for integration project?		🗌 No		
If yes, these projects include		According to client requirement		
Would the company like to provide	a total solution for	🛛 Yes		
purchasing?		🗌 Part, 🔄		
		🗌 No		
If yes, please describe it		According to	client requirement	
10.1.3 Real Case Description:		T		
Customer Name			Confidential	
Customer Country			Confidential	
Customer Region			Confidential	
Products Category			Confidential	
Order Value (USD)		Confidential		
Order Processing Process			Confidential	
Customer's Feedback			Confidential	
10.2 Real Case for Lower MOQ	& Lead Time			
Products Name	MOQ (In the last	12 Months)	Shortest Lead Time	
Auto Lens Meter	Confider	ntial	Confidential	
Slit Lamp	Confider	ntial	Confidential	
3D Auto Lens Edger	Confider	ntial	Confidential	
Rebound Tonometer	Confider	ntial	Confidential	

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	L	All Rights Res	served			Page No:	21 of 25



Auto Refractometer	Confidentia	al	Confidential			
MOQ (In the last 12months)	Less than 10					
	□ 10-20	 10-20				
	☐ 20-50	20-50				
	50-100					
	☐ 100-300					
	More than 300					
10.3 Real Case for Large Cor	ntract					
Products Name	Order (In the past 1	2 Months)	Shortest Lead Time			
Auto Lens Meter	Confidentia	al	Confidential			
Slit Lamp	Confidenti	al	Confidential			
3D Auto Lens Edger	Confidenti	al	Confidential			
Rebound Tonometer	Confidenti	al	Confidential			
Auto Refractometer	Confidenti	al	Confidential			
10.4 Overseas after-sales ser	rvice capacity					
If yes, what onsite after-sales se	ervices are included	N/A				
Equipment -installation, mainten	ance and other services	🗌 Yes [	🛛 No			
Technical advice		🗌 Yes 🛛 No				
Personnel training		🗌 Yes [	🗌 Yes 🛛 No			
Other:		N/A				
Average response time:						
Num	1	Unit				
N/A			N/A			
10.5 After-sales service capa	city					
Average Guarantee Time						
Product Category	Num		Unit			
Auto Lens Meter	1		Year			
Slit Lamp	1		Year			
3D Auto Lens Edger	1		Year			
Rebound Tonometer	1		Year			
Auto Refractometer	1		Year			
The Longest Guarantee Time						
Product Category	Num		Unit			
Auto Lens Meter	1		Year			
Slit Lamp	1		Year			
3D Auto Lens Edger	1		Year			
Report No: 22069995_T R	eport date: 31/Aug./2021 A	ssessed By	Zad Zhang			
CONFIDENTIAL	Il Rights Reserved		Page No: 22 of 25			



Rebound Tonometer	1		Year
Auto Refractometer	1		Year
Deep the company eccent small and		🛛 Yes	
Does the company accept small orde	er <i>:</i>	🗌 No	

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIA	NL.	All Rights Res	served			Page No:	23 of 25



**Assessment Report** 

# Section 11: Service Capabilities

11.1 Experience with Large-scale Procurement Contracts						
Have you had a procurem	nent contract with a Fortune 500	🗌 Yes				
company?		🛛 No				
Do you have an overseas	onsite service center?	🗌 Yes				
		🖂 No				
The country/region of you	r overseas service center	N/A				
11.2 Overseas Showro	om					
Do you have an overseas	showroom?	🗌 Yes	Yes			
		🖂 No				
Country/Region		N/A	N/A			
11.3 Offline Trade Show	N					
Have you participated in o	offline trade shows?	🛛 Yes				
Tradeshow name	CHINA (SHANGHAI) INTERNA	TIONAL	Official images from the trade show			
	OPTICS FAIR					
Date attended	11/Feb./2020		Confidential			
Host Country/Region	Other China					

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIAL All Rights Reserved			Page No:	24 of 25			



**Assessment Report** 

# Section 12: Supply Chain Capability

12.1 Centralized Procurement	
In the past 12 months, have you sold products from at	Yes
least three sub-categories?	🖂 No
12.2 Overseas Warehouse	
Do you have an overseas warehouse?	Yes
	🖂 No
Country/Region	N/A

-- End of Report --

Report No:	22069995_T	Report date:	31/Aug./2021	Assessed By	Zad Zhang		
CONFIDENTIAL All Rights Reserved		Page No:	25 of 25				