DP-2000 Human Laboratory Pupillometer System

Instructions for Use





Notices & Safety Information

Copyright and Trademark Notice

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Intended Use Notice

The NeurOptics® DP-2000 Pupillometer is an optical scanner which captures and analyzes a digital video of a subject's pupil responding to light stimulation. It should only be operated by properly trained personnel.

The results obtained from the Pupillometer scans are used for information only and are not to be used for any diagnostic process.

Regulatory Notice

Federal law restricts the sale of this system except by or on order of a physician.

Classification

Type of Equipment: Medical Equipment, Class 1

Trade names: NeurOptics® DP-2000 Human Laboratory Pupillometer

Manufactured by:

NeurOptics, Inc.

9223 Research Drive Irvine, CA 92618 | USA p: 949.250.9792 Toll Free North America: 866.99.PUPIL info@NeurOptics.com

NeurOptics.com

If you have a question regarding this product, please contact NeurOptics at one of the numbers given in Appendix B.

Safety Information

Please review the following safety information prior to operating the system.

Read the Operating Instructions fully before attempting to use the DP-2000 Pupillometer. Attempting to operate the system without fully understanding its features and functions may result in unsafe operating conditions and/or inaccurate results.

If you have a question regarding the installation, set up, operation, or maintenance of the system, contact NeurOptics as shown in Appendix B.

Warnings: Warnings and Cautions appear throughout this manual where they are relevant. The Warnings and Cautions listed here apply generally any time you operate the system.

Investigational use only.

Use of the Pupillometer - The Pupillometer is intended for use by trained personnel. If a problem is recognized while operating the system, the system must be removed from use and referred to qualified personnel for servicing. Using an inoperative system may result in inaccurate readings.

Electric shock hazard - Do not open any parts of the system. There are no user serviceable parts. Refer all servicing to a NeurOptics authorized service technician. Opening up the system will void the warranty.

Use only power adapters which are shipped with the NeurOptics® DP-2000 Pupillometer system. Using unauthorized parts will void the warranty.

Trusted Media – The DP-2000 System is shipped without antivirus protection. Attach only to trusted media.

Cautions

DO NOT submerge the system or pour cleaning liquids over or into any components of the system.

DO NOT use acetone to clean any surface of the Pupillometer.

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Section 1 - Introduction

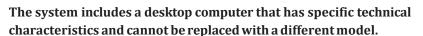
The DP-2000 Human Laboratory Pupillometer is a multi-chromatic binocular/dual camera system that tracks both pupils and stimulates both eyes. It is specifically designed for measuring the pupils of human subjects. The video stream is captured at 30Hz per second. The video frame is digitized into 640X480 pixels with 8-bit gray level resolution. The profile of the light stimulation can be customized using a graphical user interface (GUI). Pupillary variables are automatically analyzed, reported graphically in a special window and saved into a results file. Light stimulation has four different chromatic options: Red (622 nm), Green (528 nm), Blue (463 nm) and White. Light is emitted through a diffusing screen (approx. 50° x 35° of visual angle).

Section 2 - Set-up

The NeurOptics® DP-2000 system is packed into a shipping box. Included with your system are:

- The DP-2000 Pupillometer System See Figure 1
- Slim Desktop Computer with Monitor, Keyboard & Mouse
- Windows 10 Pro
- Instructions for Use

Please check the contents of your box when delivered. If anything is damaged or missing please contact NeurOptics at once at 1-949-250-9792. DO NOT ATTEMPT TO USE A DAMAGED SYSTEM!



• Place the Computer, Monitor, Keyboard, and mouse on a stable surface. Plug in all accessories to the computer. See Figure 2.



Figure 1



Figure 2

• Assemble Camera Arm by mounting base post on a stable table and lock into place by twisting the knob on the base of the arm. Loosen set screw with the included hex-head wrench and slide the arm down on post to desired height. Retighten set screw. Place the camera assembly on end of the arm in slot and lower till it locks in place. See Figure 3.









Figure 3

- Connect the two red camera video (firewire) cables to the Unibrain Hub (Unit D). The other ends should already be pre-connected to the cameras. See Figure 4.

Figure 4

• Connect the grey Serial Cable (Item #2) and 9V power supply (Labeled NEUR-PWR4) to the open ports of the Control Box (Unit A) as shown below. See Figure 5. The other end of the Serial Cable is connected to the back of the computer as shown on the right. See Figure 6.



Figure 5

• Connect the Firewire cable labeled #1 to the back of the computer in the Firewire port labeled #1. See Figure 7.



Figure 7

- Inner pupillary distance can be adjusted to fit the subject's eyes by turning the knob located below the cameras. Never grab the camera body to change the arm position or pivot cameras. Always use the metal support under the cameras. See Figure 8.
- Press the power button on the computer. Click on the DP-2000 icon on the screen to get started.



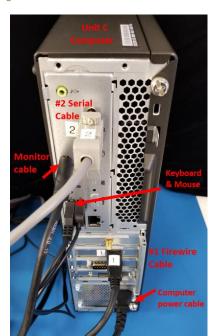


Figure 6

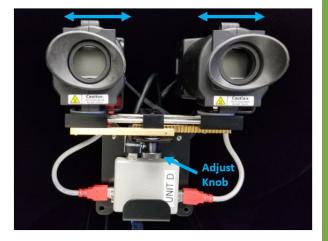


Figure 8

Section 3 – Taking a Measurement

All possible options to operate the program are listed in the left most side of the GUI (see Figure 9). The live video from the two cameras is displayed in the same GUI. Pupil tracking is represented by a circle fitting the contour of the pupil - the subject's left eye is displayed on the right and the subject's right eye is displayed on the left. This is to maintain the natural perspective of the examiner facing the subject during the examination. The left pupil (OS) is tracked with a yellow circle and the right pupil (OD) with a green circle. This association (left-yellow and right-green) is also used for display purposes (see below). Light stimulus waveform and the corresponding pupil response dynamics are shown together in the graph located at the bottom of the GUI. Note: the software version can be checked by pressing the Version button.

Subject ID

The first two windows on the top of the list (Figure 9) serve to enter information regarding the current measurement (Subject ID and Note). This information is attached to the master data result file produced at the end of the measurement and used for defining the name of the result file. Note: the following characters are not allowed to be used "./[]:+|<>=;,*?" in the Subject ID.

March March

Figure 9 – The main program graphic user interface GUI.

Edit Stimulus Profile

The Edit Stimulus Profile button serves to define and to save a new light stimulus profile or to load a predefined profile. The basic unit of a profile is a light step during which a light pulse is flashed to the eye and then turned off. Light pulses can be repeated a number of times, thereby forming a section. Pulses and/or sections can be concatenated in order to define the final light stimulus profile. The Edit Stimulus Profile GUI (Figure 10) is used to define all the elements of the light stimulus profile.

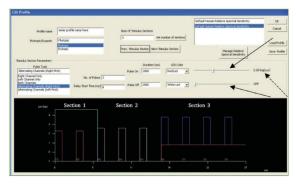


Figure 10 – The Edit Stimulus Profile GUI, Section 1.

Light intensities are defined in Lux or pure radiometric units (W/m2). Two different definitions of Lux are available. They correspond to the scotopic or photopic CIE human luminosity sensitivity function and they can be selected using the drop-down menu on the top-left side of the GUI. Lux or radiometric values can be set using the horizontal slide button (see arrows); and, they are incremented by 0.1 log unit steps. When a Lux value cannot be achieved, the number (see the dashed arrow, Figure 10) is displayed in red and a lower intensity must be selected instead.

A pulse defines single pupil stimulation and a section is a repetition of pulses having the same intensity, timing, and wavelength characteristics. One or more pulses define a section.

For example, Section 1 (Figure 10) is defined by two pulses of red light of 2.20 logLux flashed for 1000 msec and then turned OFF for another 2,000 msec (total duration 3 seconds). The eyes are stimulated alternatively with the right eye (OD) stimulated first.

Section 2 (Figure 11) is defined by three pulses of white light – 2.60 logLux, 1,000msec ON and 2,000msec OFF, both eyes stimulated simultaneously.

Section 3 (Figure 12) is defined by four pulses of blue light over a red background – intensity of the blue light is 3.80 logLux and intensity of the red background is 0.80 logLux. Only the left eye (OS) is stimulated.

All the elements of the pulse are definable using the corresponding input window in the Edit Stimulus Profile GUI (Figures 10, 11 and 12). A new Section can be added or deleted by just entering the proper number in the Num. Of Stimulus Section box. Sections can be browsed and edited using the Prev. Stimulus Section and Next Stimulus Section buttons.

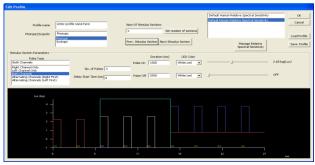


Figure 11 – The Edit Stimulus Profile GUI, Section2.

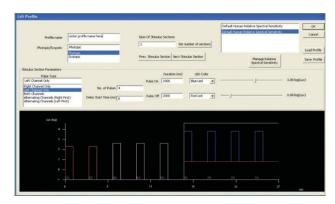


Figure 12 – The Edit Stimulus Profile GUI, Section2.

The final profile can be named, saved (it will have a .SPF extension) and retrieved later for execution using the buttons Load Profile and Save Profile. Once the profile is defined (or loaded if previously defined) press OK – that light stimulus profile is now ready to be executed. Note that the Stimulus Profile can be changed and defined also by editing the profile file using Microsoft Excel. The file contains all the instructions necessary for the editing and it has all the variables available in the GUI. Open the file with Excel, and once the new profile is defined in the spreadsheet and ready to be used in an experiment, then the file should be closed and saved as Text (Tab delimited) keeping the same extension SPF.

With the Manage Relative Spectral Sensitivity feature it is possible to upload and use a different Relative Spectral Sensitivity function than the one used by default, but this needs to be provided by the user. The new Relative Spectral Sensitivity must be provided as an ASCII file of two columns for photopic and scotopic defining the function within the range 400nm and 700nm and increment of 1nm. Each row corresponds to a 1nm increment. Please contact NeurOptics for further instructions.

Preparing the subject for the experiment and checking the tracking.

The subject should be positioned close to the camera, with the eye in front of the lens, visible and well in focus in the center of the video display in the main GUI (Figure 9 and Figure 13). It is advisable to slightly rotate the two cameras nasally, in order to form a small angle between the two cameras (as opposed to keeping the cameras perfectly straight and parallel), for a better fit to the subject's eyes. A green circle should automatically track the right eye (Displayed on the left and a yellow circle tracks the left eye (displayed on the right). It is very important to

carefully check the accuracy of the tracking before initiating the experiment. If the circles are not precisely around the pupil or there is no circle or there is only a red circle moving erratically around the video frames and tracking other portions of the image or the green (or yellow) circle turns occasionally into red – all of these are signs of bad tracking and a few corrective steps are necessary before initiating the measurement. Below is the list of all the available options.



Figure 13 – The subject pupils are well in focus in the center of the video display in the main GUI.

Digital Zoom.

The zoom button toggles between the No Zoom and the 2x Zoom video option. When the zoom is enabled, the video of both eyes is shown magnified. This feature is included in the GUI solely for assisting the user in focusing and/or adjusting the camera position or the pupil threshold. It has no effect on the tracking algorithm or the optical configuration.

Binary/Grayscale Switch.

This switch changes the video display mode between grayscale and black and white (binary) display. It does not affect the tracking algorithm and it may be used simply for visualizing the tracking algorithm's performance and helping with the setting of the threshold. When the binary view is enabled (Figure 14) the pupil must look like a black solid circle surrounded by white.

Manual Loci/Automatic Loci Switch.

When the Manual Loci feature is enabled, the user can manually locate the region of interest for searching for the pupil. This region of interest is a sub-window of the entire video frame containing the pupil. It is where the tracking algorithm will be searching for the pupil. This region is usually the entire field of view. However, in some cases, for example when strong dark patches such as heavy makeup or eyelashes are also included in the field of view or in the case of some types of iris pigmentation, the detection of the pupil can be problematic. In these cases, it is advisable to manually restrict this region to a smaller and more specific portion of the field of view. Left mouse click on the desired top-left corner of the restricted region of interest and then drag the region down to the proper size. A red rectangle overlapped on the live video indicates the position and size of the pupil searching region of interest (Figure 15). recording and the time(s) of the press (or presses) are reported in the plot and saved with the record.

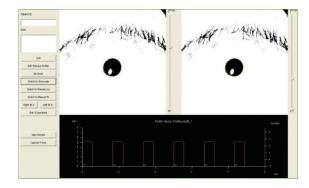


Figure 14 - Binary view enabled

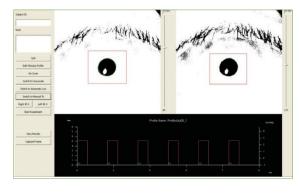


Figure 15 – The Manual Loci feature can improve pupil and tracking by restricting the region of interest for searching the pupil (see red rectangular)

The Manual/Automatic Threshold (Th) Switch.

Manual pupil threshold is set using the Manual/Automatic Threshold (Th) Switch button. This option must be used in conjunction with the binary view introduced above. It serves to refine the binary threshold defining the pupil. The pupil must appear as a uniform black patch. A good threshold is when the pupil is clearly isolated from the rest of the searching region of interest appearing in white (as displayed in Figure 14 or 15). The threshold can be changed using the vertical scroll bar on the right side of each video. If the threshold is too high (Figure 16, video on the left) the pupil looks fuzzy, blackness is sparse and not contiguous, and the tracking circle is red which means poor tracking. If the threshold is too low (Figure 17, video on the left) the image looks too dark and the pupil is not clearly isolated from the background.

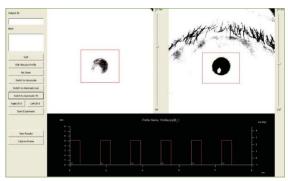


Figure 16 – Threshold too high on the left video (right pupil) - pupil looks fuzzy, blackness is sparse and not contiguous. Note the tracking circle in red which indicates poor tracking.

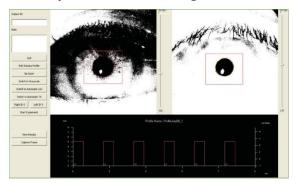


Figure 17 – Threshold too low on the left video- the image looks too dark and the pupil is not clearly isolated from the background.

The PR Suppression Switch.

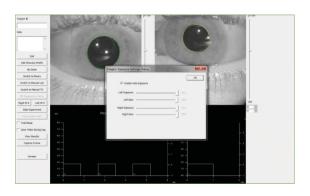
Operates on the two bright spots (Purkinje Reflections) created on the surface of the cornea - when the suppression is enabled, a special algorithm is applied to attenuate the effect of those bright spots on the pupil tracking. It is suggested only for very small pupils when the size of the reflections is relatively large compared to the size of the pupil.

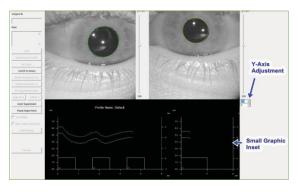
Auto-Exposure Setting

The auto-exposure or the value of the exposure and the gain of the two video cameras can now be set manually. This is to optimize the quality of the video when necessary. To change the exposure and the gain of the two cameras, right click on one of the two live videos and open the "Imager's Exposure Setting Dialog" window.

Small Graphic Inset

A small graphic inset is added in the main window, next to the original graphic. The new graphic displays in real time the current pulse and the two pupil waveforms at larger magnification and it is updated at each pulse transition. The range of the y axis range of the new graphic can be set using two small scroll boxes located on the top-right side of the graphic. The new graphic allows the user to magnify the display of each pulse in real time during the measurement.





Start Experiment.

If both pupils are tracked correctly, a press of the Start Experiment button will initiate the execution of the light stimulus protocol. The subject should be kept firmly in front of the cameras – blink or similar artifact should be minimized. A live progressing trace of the two pupils' dynamics (yellow for the left pupil and green for the right pupil) are shown on the bottom inset of the main GUI (Figure 18, note a blink at second 11). At the end of the measurement – the results file is saved in folder C:\NeurOptics\DP2000\DATA and results are displayed in the View Results GUI. Note that the Start Experiment switches to Abort Experiment as soon as the measurement is initiated, and it must be used to abort the experiment. Another switch, Pause/Resume Experiment (added in the new version of the software and not shown in Figure 10) serves to interrupt momentarily (and then resume) the ongoing measurement. The Pause/ Resume switching operations can be performed also by simply pressing the space bar on the keyboard.

Right IR – Left IR.

Each camera has two pairs of IR LEDs (0 and 1) that can be toggled using the IR buttons for the left and right channel. This can be useful for small pupils, if one of the IR corneal reflections is overlapping on the pupil, disrupting the tracking. Switching to the other pair of IR sources can often resolve the problem and improve the visibility of small pupils.

Capture Frame.

Two snapshots of the two videos being displayed in the GUI can be saved as bitmaps with a press of this button prior the start of a measurement. A new pair of bitmaps are generated and saved in memory at each press.

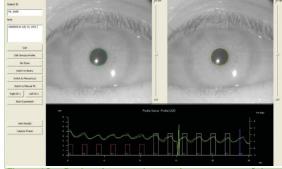


Figure 18 – During the experiment, the responses of the two pupils are displayed in real time in the graph at the bottom of the main GUI - yellow is for left pupil and green is for the right pupil.

Section 4 – Viewing the Results

Pupil variables are reported graphically and numerically in the View Results GUI (Figure 19). The small inset on the right (A) shows each single pulse and different pulses can be viewed by pressing the Previous Pulse and the Next Pulse button. The horizontal gray lines overlapped on the plot indicate the Maximum and Minimum pupil size (before the constriction and at the peak of the constriction) – the vertical line indicates the latency of the constriction. The user can select which pulse to show: both pulses, left pupil only or right pupil only. The entire pupil waveform and the light stimulus protocol are shown together on the long graph at the top of the GUI (B) – similarly to the Edit Stimulus Profile GUI; a light-blue rectangle indicates a section.

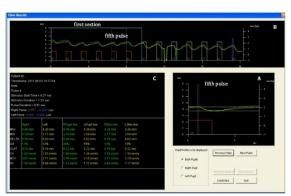


Figure 19 – The View Result GUI. The entire pupil profile (and stimulus waveform) is displayed in B. Pulse can be individually inspected in A. Pupil variables (belonging to that specific pulse in A) are reported in C. In this example, the fifth pulse of the first section has been selected in the display

In the example (Figure 19), the 5th pulse in the list is shown in the small inset A and that is part of the first section (pulses are enumerated starting from zero, so the 5th pulse is pulse no. 4). Pupil variables and pulse information are reported in C. The information includes the ID of the subject and Notes about the subject and the measurement, duration, type and intensity of the light stimulation. Below, still in the inset C, are the pupil variables for the right and left pupil light reflex – the variables measured are:

Max = pupil diameter baseline before the constriction

Min = pupil diameter at the peak of the constriction

Delta = Max - Min

CH = (Max - Min)/Max, given in percentage

CLAT = latency of the constriction (delay of the onset of the pupil constriction)

MCV = maximum constriction velocity

DV = mean dilation velocity (recovery after the constriction)

If the pulse belongs to a section composed of many pulses (repeated in time) then the analysis in C returns the average of all the above variables for all the pulses composing the section. RPupil Ave is the average of all the right pupil responses (for both right and left eye stimulation), LPupil Ave is the average of all left pupil responses (for both right and left eye stimulation), RStim Ave is the average of all the responses (left and right pupils together) corresponding to all the right eye stimulations and LStim Ave is average of all pupil responses to all the left eye stimulation. In case of a blink (or similar artifact) as, for example, for pulse no. 7 (belonging to the second section see Figure 20), pupillary variables are reported in red and some of them might not have been calculated (and reported with "--", see Figure 20). A feature in the View Results GUI allows the recovery of an artifact affected pulse (Figure 21). The blink, or gap in the pupil profile, can be linearly interpolated by selecting the option L. Pupil Fill Data Gap for the left pupil or R. Pupil Fill Data Gap for the right pupil – in this case, a line is drawn automatically over the gap (see example in Figure 21, right insert) to indicate the fitting. If the linear fitting is not properly performed (or the gap is too large to be interpolated efficiently), the MAX, MIN and LAT of the pupil reflex can be set manually using the three input cursors at the bottom of the graph (Figure 21). Changes in the MAX, MIN or LAT values are reflected by the position of the three gray lines in the plot to give a visual feedback of the values being changed (two horizontal lines for MAX and MIN, and one vertical line for LAT). The pupil variables reported in table C (Figure 19) are automatically updated after the linear fit option has been selected or a value during manual setting has been changed.

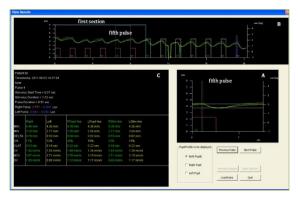


Figure 20 – In the case of a blink 9 or a similar artifact) and in the example for pulse 7, (belonging to the second section) pupillary variables are reported in red or not reported.

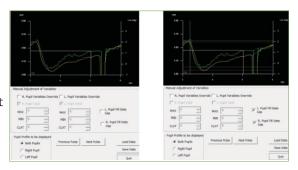


Figure 21 – The View Results GUI allows the recovery of pulses affected by pupil tracking artifacts (such as eye blinks). If the Fill Data

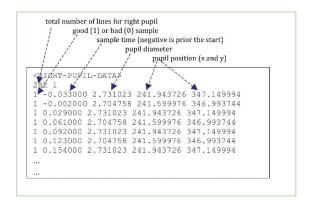
Gap option (see right panel) is selected, a linear fit is applied to the artifact and pupil variables are re-analyzed accordingly (and reported again in C). If the linear fit cannot be properly conducted, the user can also set manually the MAX, MIN and LAT of the pupil light reflex.

Finally, the options R. Pupil Valid or L. Pupil Valid (Figure 21) serve to flag as abnormal (and to report in red) a reflex that is otherwise considered normal; this is to manually correct those pulses where a tracking error was missed by the algorithm (and thus the reflex is forced not to be included in the analysis).

Data are automatically saved after the measurement in the C:\NeurOptics\DP2000\DATA and they can be loaded and viewed later using the Load Data button in the same GUI. Remember to use a different name for the data file if the fitting or a manual override has been performed on the data – this is to preserve the original file. Data files can also be viewed with Excel.

Different segments corresponding to different information are defined in a data file:

The segment <STIMULUS-DEFINED> reports the stimulus settings as defined in the SPF stimulus profile file.



The segment <STIMULUS-GENERATED> reports the exact sequences and values of the lights emitted during the experiment. The first column indicates the section that particular light intensity belongs to; the second column Indicates which eye was stimulated (0=right, 1=left, 2=both, 3 or 4 if background light); then Stimulus start time; (seconds elapsed from start of measurement); the Color of the stimulation for the Right eye (0=white, 1=red, 2=green, 3=blue); Intensity of the stimulation for the Right eye (given in lux); the Color of the stimulation for the Left eye (0=white, 1=red, 2=green, 3=blue); Intensity of the stimulation for the Left eye (given in radiometric units); Intensity of the stimulation for the Left eye (given in lux). The right and left stimulus intensities could slightly differ due to the differences in the LEDs light emittance.

Raw pupil tracking data are included in the data file; they are reported under the label <RIGHT-PUPIL-DATA> or <LEFT-PUPIL-DATA>; for each time sample (line) the second column is time, the third column is pupil diameter, the third and fourth column is the x and y position of the pupil. The first column indicates whether that sample was successfully tracked (1) or it was affected by a blink or similar artifact (0).

The results of the analysis are reported in different parts of the file and different formats. Tables <RIGHT-PUPIL-ANALYSIS> and <LEFT-PUPIL-ANALYSIS> are obsolete and can be ignored: In this table there is one line for each pulse in the stimulus waveform and a total of 25 values. From left to right those values correspond to: Pulse index; Section index; which eye was stimulated (0=right, 1=left, 2=both); Stimulus start time; (seconds elapsed from start of measurement); Stimulus duration (sec); Pulse duration (sec); Intensity of the stimulation for the Right eye (lux); Intensity of background illumination for the Right eye (lux); Intensity of the stimulation for the Left eye (lux); Intensity of background illumination for the Left eye (lux); the 10th value is obsolete and should be ignored; the Color of the stimulation of the Right eye (0=white, 1=red, 2= green, 3=blue); the Color of the background illumination for the Right eye; the Color of the stimulation of the Left eye; the Color of the background illumination for the Left eye; the 16th value is binary indicating the presence of a blink or artifact (1=tracking ok, 0=tracking not ok, pulse should be discarded); Max pupil diameter (mm); Min pupil diameter (mm); Max pupil area (mm2); Min pupil area (mm2); Latency of the constriction (sec); Average constriction velocity (mm/sec); Maximum constriction velocity (mm/sec); Average dilation velocity (mm/sec). Each line represents a single pulse in the list of the stimulus waveform; the two sections <RIGHT-PUPIL-ANALYSIS> and <LEFT-PUPIL-ANALYSIS>, have the initial fields, those describing the stimulation, identical.

What follows in the file is a list of tables in the same format as they were displayed in the View Results GUI of the program, one for each of the pulses and for all the pupil and measurement variables. The right and left pupil profile are reported again below this list of tables.

Finally, a complete table is provided at the very bottom which includes all stimulus and variables values. This table can be selected and pasted onto a clean new Excel spread sheet for further analysis and statistics. The names of the columns should be self-explanatory. Note that the third and fourth columns named Stimulus Sections Start Index and Stimulus Sections End Index indicate the index of the pupil profile in sections <RIGHT-PUPIL-DATA> or <LEFT-PUPIL-DATA> corresponding to that specific row (pulse) in the table. This is to allow further analysis of the corresponding raw pupil profile.

For the check box "Trial Mode"

The trial mode enables a measurement without saving the results. The use of this mode is recommended only when the experimental logistics are being tested, to avoid superfluous "test" result files in the DATA folder.

For the check box "Save Video During Exp."

If the box is checked, video frames are saved to the hard drive during the experiment and they can be accessed and viewed later. Only one parameter needs to be specified, which is the length of the video, in terms of number pulses to be recorded and saved. The two videos from the left and right eye are saved in a binary format, they can read using, for example, a Matlab script provided by Neuroptics. Contact Neuroptics for information.

No Zoom Switch to Binary Switch to Manual Loci Switch to Manual Th PR Suppression None Right IR 0 Left IR 0 Start Experiment Pause Experiment Trial Mode Save Video During Exp. View Results Capture Frame

Edit Stimulus Profile

Section 5 - Cleaning & Maintenance

The DP-2000 Human Laboratory Pupillometer is a non-sterile product.

To clean the surfaces of the DP-2000 pupillometer, use a soft, lint-free wipe with a quaternary disinfectant.

DO NOT attempt to sterilize any part of the DP-2000 system, as sterilization may damage the system.

DO NOT immerse or drip liquids on any component of the system.

The laptop LCD and keyboard should be cleaned using a soft, lint-free wipe with IPA.

Clean the lens with lens cleaning solution and a lint free cloth.

Appendix A — Troubleshooting

Issue	Possible Reason	Solution
	Control Box not powered or connected	• Quit the DP-2000 program
		• Check power connection to the control box and serial cable connections
		• Restart the DP-2000 program
Error Window: "Stimulus Hardware Not Detected"	Firewire cable to Computer not connected	 Quit the DP-2000 program Check all power and video connections Restart the DP-2000 program
	Connection error or malfunctioning hardware	 Quit the DP-2000 program Check power connection to the control box and serial cable connections
		Restart the DP-2000 program
		• Reboot computer, contact Manufacturer if error persists
Error Window:	ubCore stack is not active	• Quit the DP-2000 program
"FiInitialize failed with error: ubCore stack initialization error."		• Reboot computer, contact Manufacturer if error persists
Error Window:	System unable to read settings	Contact Manufacturer
"Initializing Please Wait"		
Error Window:	Missing or corrupted system data	Contact Manufacturer
"Settings file DP2000Settings.dat can't be read."	file	
Program becomes unresponsive.	Serial Cable unplugged	 Quit the DP-2000 program Check cable connections Restart the DP-2000 program.
Program fails to start or crashes	Program corrupted	Contact Manufacturer if error persists
Window Fogging	Seal around window failed	Contact Manufacturer

$Appendix\,A-Trouble shooting\,Continued$

Issue	Possible Reason	Solution
Blue Screen Error	Windows 10 Error	Check all cable connections
		 Perform a "Hard" computer reboot by pressing and holding the computer power button until itreboots
		Contact Manufacturer if errorpersists
Error Window:	System failed to process video	• Quit the DP-2000 program and restart
FiCreateIsochReceiveEngine failed with error: Error in video streaming initialization.		• If error persists, reboot computer
Error Window:	System failed to process video	• Quit the DP-2000 program and restart
FiLocateCamerasEx failed with error: error in video streaming initialization		• If error persists, reboot computer
Error Window:	Control system does not detect the	• Quit the DP-2000 program.
"FiOpenCameraHandleEx	camera(s).	Check all power and video connections.
failed with error: FIREi STATUS CAMERA READ FAIL"		• Restart the DP-2000 program
Error Window:	Camera(s) not powered or	• Quit the DP-2000 program.
"FiOpenCameraHandleEx failed with error: FIREI STATUS INVALID PARAMETER"	connected.	 Check all power and video connections.
		• Restart the DP-2000 program
Error Window:	System failed to process video	• Quit the DP-2000 program and restart
FiQueryCameraRegister failed with error: Error in video streaming initialization		• If error persists, reboot computer
Error Window: FiStartCamera failed with error: Error in video streaming initialization	System failed to process video	Quit the DP-2000 program and restart

Appendix A — Troubleshooting Continued

Issue	Possible Reason	Solution
Error Window: FiStartCamera failed with error: FIREI STATUS CAMERA ALREADY RUNNING	System failed to process camera image	Quit the DP-2000 program and restart
Number of cameras detected by application is less than two	Camera(s) not powered or connected	 Quit the DP-2000 program. Check all power and video connections Restart the DP-2000 program
"Initialization Video" error on one or both cameras	System failed to process camera image	Quit the DP-2000 program and restart
Error Window: Can't save data	The DATA subfolder is not write- enabled or missing	 Verify the DATA subfolder is located under the C;/NeurOptics/DP-2000 directory and is not write protected Create a DATA folder if necessary
Error Window: Loading Profile Can't open selected file - Can't open profile file	Camera(s) not powered or connected.	 Quit the DP-2000 program. Check all power and video connections. Restart the DP-2000 program
Error Window: FiQueryCameraRegister failed with error: Error in video streaming initialization	Possible missing file or the file has no read permission	 Verify file properties are not write protected Verify parent folder properties are not write protected
Error Window: Can't read from selected file - can't load the profile file data	Possible corrupted file	Delete corrupted file and reedit
Error Window: Lux level Error!!! Please readjust parameters	Saving profile with invalid lux parameters (shown in red)	Adjust Lux level

Appendix B — Contact & Ordering Information

The following **options and accessories** can be ordered from NeurOptics:

DP-2000	NeurOptics Human Laboratory Pupillometer – US
NEUR-PWR4	Control Box Power Supply – US
NEUR-COMP-04	Computer
NEUR-3089	DP-2000 Eyecup

For more information, please contact NeurOptics or visit www.NeurOptics.com

Distributed by:



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Warranty

- A. Standard Limited Warranty. NeurOptics (the Manufacturer) warrants to the original end user purchaser (the Purchaser) that the enclosed product (Product) purchased by the Purchaser, at the time of delivery to the Purchaser, shall be substantially free from defects in material and workmanship. The Manufacturer makes no warranty (express, implied, or statutory) for Products that are modified (except as expressly contemplated herein) or subjected to unusual physical stress, misuse, improper operation, neglect, improper testing, use in combination with other products or components other than those for which the Products were designed, or use in any manner or medical procedure for which the Products are not indicated. If the Product is opened up by anyone other than an authorized service technician of NeurOptics, the warranty shall bevoid.
- **B.** Remedy. Purchaser's exclusive remedy and the Manufacturer's sole liability for breach of the foregoing warranty shall be, at the Manufacturer's sole option and election, to replace the Product or credit Purchaser for the net amount actually paid for any such Product; provided that (i) the Manufacturer is notified in writing within one (1) year after Purchaser's receipt of the Product that such Product failed to conform, including a detailed explanation in English of any alleged nonconformity; (ii) such Product is returned to the Manufacturer within one (1) year after Purchaser's receipt of the Product as designated by the Manufacturer using a Returned Good Authorization number issued by the Manufacturer and (iii) the Manufacturer is reasonably satisfied that the claimed nonconformities actually exist. Except as expressly provided in this paragraph, Purchaser shall not have the right to return Products to the Manufacturer without the Manufacturer's prior written consent.
- C. Exclusion of Other Warranties. EXCEPT FOR THE LIMITED WARRANTY PROVIDED IN (A) ABOVE, NEUROPTICS GRANTS NO OTHER WARRANTIES OR CONDITIONS, EXPRESS OR IMPLIED, AND MANUFACTURER SPECIFICALLY DISCLAIMS THE IMPLIED WARRANTIES AND CONDITIONS OF MERCHANTABILITY AND FITNESS FOR A PARTICULAR PURPOSE. NEUROPTICS NEITHER ASSUMES NOR AUTHORIZES ANY OTHER PERSON TO ASSUME ANY OTHER LIABILITIES ARISING OUT OF OR IN CONNECTION WITH THE SALE OR USE OF ANY PRODUCT.

Returned Goods Policy

The DP-2000 Pupillometer is a custom-made product and no returns will be allowed unless the Product is defective. Products must be returned using a Returned Goods Authorization number issued by the Manufacturer and sent well wrapped and protected via Fedex or UPS with insurance for replacement value, to be accepted for replacement or repair. Determination of a product defect will be made by NeurOptics, which determination will be final.



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