GIMA
PROFESSIONAL MEDICAL PRODUCTS

LUCE DI WORTH
WORTH LIGHT TEST
TEST DE WORTH
WORTH-TEST
TEST DE WORTH
TESTE DE WORTH
ΤΈΣΤ ΦΩΤΟΣ ΤΗΣ *WORTH*
WORTH ثروو ءضو فحص

Manuale utente - User manual - Notice d'utilisation - Gebrauchsanweisung
Manual de uso - Guia para utilização - Οδηγίες χρήσης - دليل الاستخدام

È necessario segnalare qualsiasi incidente grave verificatosi in relazione al dispositivo medico da noi fornito al fabbricante e all'autorità competente dello Stato membro in cui si ha sede.

All serious accidents concerning the medical device supplied by us must be reported to the manufacturer and competent authority of the member state where your registered office is located.

Il est nécessaire de signaler tout accident grave survenu et lié au dispositif médical que nous avons livré au fabricant et à l'autorité compétente de l'état membre où on a le siège social.

Jeder schwere Unfall im Zusammenhang mit dem von uns gelieferten medizinischen Gerät muss unbedingt dem Hersteller und der zuständigen Behörde des Mitgliedsstaats, in dem das Gerät verwendet wird, gemeldet werden.

Es necesario informar al fabricante y a la autoridad competente del Estado miembro en el que se encuentra la sede sobre cualquier incidente grave que haya ocurrido en relación con el producto sanitario que le hemos suministrado.

É necessário notificar ao fabricante e às autoridades competentes do Estado-membro onde ele está sediado qualquer acidente grave verificado em relação ao dispositivo médico fornecido por nós.

Σε περίπτωση που διαπιστώσετε οποιοδήποτε σοβαρό περιστατικό σε σχέση με την ιατρική συσκευή που σας παρέχουμε θα πρέπει να το αναφέρετε στον κατασκευαστή και στην αρμόδια αρχή του κράτους μέλους στο οποίο βρίσκεστε.

يجب الإبلاغ فورا عن أي حادث خطير وقع فيما يتعلق بالجهاز الطبي الذي زودنا به إلى الجهة الصانعة والسلطة المختصة في الدولة العضو التي يقع فيها

REF DP-5048 **(GIMA 31289)**

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MD CE

M31289-M-Rev.3.05.22

Worth light Test is used to assess a patient's flat Fusional ability.
Flat Fusional Testing is Indicated any time stereopsis falls between (50) Sec of Arc, on those patients with suspected strabismus and on preschool Children. The Worth Test should also be used when evaluating cases of reduced monocular visual acuity that does not improve with the Pinhole Test.

Procedure

1. With the best refraction correction worn by the patient, place the Anaglyph glasses over the patient's correction, with red filter over the right eye.
2. In a slightly dimmed room, turn on the flashlight and hold the Worth Test with the red dot orientated up at approximately 16" from the patient and slightly below the LOS.
3. Conduct the following Monocular Check first:
   a. Cover the right eye; ask how many dots the patient sees. They should report 3 green.
   b. Then Cover the left eye and ask how many dots they see now. They should report 2 red.
4. Next, Conduct the binocular test:
   a. With both eyes uncovered ask a third time how many dots they see. If normal flat fusion is present they should report 4.

Pediatric Note: This test can reliably be conducted on preschoolers as young as 2 if they are allowed to point to the dots.

5. Abnormal responses:
   a. If the patient reports only 2 red dots under binocular conditions, this indicates that they are suppressing the left eye.
   b. If the patient reports seeing 3 green dots under binocular conditions, then they are suppressing the right eye.
   c. If they report 5 dots, they are diplopic. The type of diplopia can then be determined by asking which side are the green dots.
   If the green dots are located on the right, the patient has a eso deviation; to the left, an exo deviation.
   d. If the green dots are reported above or below the red dots then a vertical deviation exists. A report of the green dots above the red dots would be seen with aright hyper deviation.
   e. With a report of 6 or more dots, one should question the patient's reliability.
6. The binocular view is repeated at 5 and 10 feet, then repeat all these distances under greatly reduced light as both suppression and ocular deviation can be different under varying lighting conditions.

***Disposal**: The product must not be disposed of along with other domestic waste. The users must dispose of this equipment by bringing it to a specific recycling point for electric and electronic equipment.*

GIMA WARRANTY TERMS
The Gima 12-month standard B2B warranty applies.

	**IT** Fabbricante **GB** Manufacturer **FR** Fabricant **DE** Hersteller **ES** Fabricante **PT** Fabricante **GR** Παραγωγός **SA** الشركة المصنعة
	**IT** Data fabbricazione **GB** Date of manufacture **FR** Date de fabrication **DE** Herstellungsdatum **ES** Fecha de fabricación **PT** Data de fabrico **GR** Ημερομηνία παραγωγής **SA** تاريخ التصنيع
	**IT** Codice prodotto **GB** Product code **FR** Code produit **DE** Erzeugniscode **ES** Código producto **PT** Código produto **GR** Κωδικός προϊόντος **SA** كود المنتج
	**IT** Numero di lotto **GB** Lot number **FR** Numéro de lot **DE** Chargennummer **ES** Número de lote **PT** Número de lote **GR** Αριθμός παρτίδας **SA** رقم الدفعة
	**IT** Conservare in luogo fresco ed asciutto **GB** Keep in a cool, dry place **FR** À conserver dans un endroit frais et sec **ES** Conservar en un lugar fresco y seco **DE** An einem kühlen und trockenen Ort lagern **PT** Armazenar em local fresco e seco **GR** Διατηρείται σε δροσερό και στεγνό περιβάλλον **SA** يحفظ في مكان بارد وجاف
	**IT** Conservare al riparo dalla luce solare **GB** Keep away from sunlight **FR** À conserver à l'abri de la lumière du soleil **DE** Vor Sonneneinstrahlung geschützt lagern **ES** Conservar al amparo de la luz solar **PT** Guardar ao abrigo da luz solar **GR** Κρατήστε το μακριά από ηλιακή ακτινοβολία **SA** يحفظ بعيدا عن أشعة الشمس
	**IT** Seguire le istruzioni per l'uso **GB** Follow instructions for use **FR** Suivez les instructions d'utilisation **DE** Folgen Sie den Anweisungen **ES** Siga las instrucciones de uso **PT** Siga as instruções de uso **GR** Ακολουθήστε τις οδηγίες χρήσης **SA** اتبع تعليمات الاستخدام
	**IT** Dispositivo medico conforme al regolamento (UE) 2017/745 **GB** Medical Device compliant with Regulation (EU) 2017/745 **FR** Dispositif médical conforme au règlement (UE) 2017/745 **DE** Medizinprodukt im Sinne der Verordnung (EU) 2017/745 **ES** Producto sanitario conforme con el reglamento (UE) 2017/745 **PT** Dispositivo médico em conformidade com a regulamento (UE) 2017/745 **GR** Ιατρική συσκευή σύμφωνα με την ΚΑΝΟΝΙΣΜΟΣ (ΕΕ) 2017/745 **SA** جهاز طبي يتوافق مع التوجيه (UE) 2017/745
	**IT** Attenzione: Leggere e seguire attentamente le istruzioni (avvertenze) per l'uso **GB** Caution: read instructions (warnings) carefully **FR** Attention: lisez attentivement les instructions (avertissements) **DE** Achtung: Anweisungen (Warnungen) sorgfältig lesen **ES** Precaución: lea las instrucciones (advertencias) cuidadosamente **PT** Cuidado: leia as instruções (avisos) cuidadosamente **GR** Προσοχή: διαβάστε προσεκτικά τις οδηγίες (ενστάσεις) **SA** الحذر: قراءة التعليمات (التحذيرات) بعناية
	**IT** Smaltimento RAEE **GB** WEEE disposal **FR** Disposition DEEE **DE** Beseitigung WEEE **ES** Disposición WEEE **PT** Disposição REEE **GR** Διάθεση WEEE **SA** التخلص WEEE
	**IT** Dispositivo medico **GB** Medical Device **FR** Dispositif médical **DE** Medizinprodukt **ES** Producto sanitario **PT** Dispositivo médico **GR** Ιατροτεχνολογικό προϊόν **SA** جهاز طبي
	**IT** Importato da **GB** Imported by **FR** Importé par **DE** Importiert von **ES** Importado por **PT** Importado por **GR** Εισάγεται από **SA** استيراد
	**IT** Rappresentante autorizzato nella Comunità europea **GB** Authorized representative in the European community **FR** Représentant autorisé dans la Communauté européenne **ES** Representante autorizado en la Comunidad Europea **PT** Representante autorizado na União Europeia **GR** Εξουσιοδοτημένος Αντιπρόσωπος στην Ευρωπαϊκή Κοινότητα **SA** الممثل المعتمد في المجموعة الأوروبية