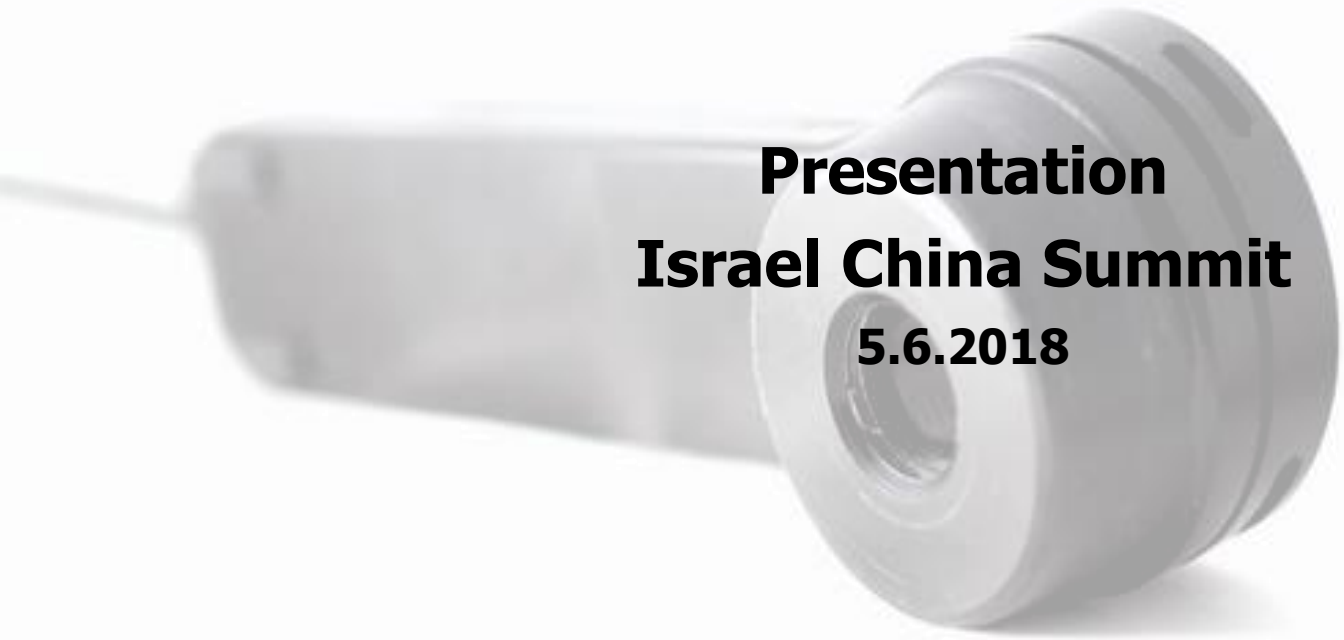

Presentation
Israel China Summit
5.6.2018

The Company

- Offices located in Ra'anana, Israel
- IP: Patents IL167559, US 11/909,103, EU 6711339.9, JPN 2007-555773
- CE Mark & ISO 13485: FIM CE Mark & ISO approval since 2012
- Received Israel Ministry of Health "AMAR" approval.

Mydriatics Today -The Problem

Time consuming:

- **Slow dilatation onset: up to 20 minutes – often a repeat application is necessary**
- **Slow eye recovery: up to 8 hours – sometimes as long as 72 hours**

Discomfort:

- **Irritated, red and stinging eyes, blurred vision**
- **Unable to read, work or drive for 4-6 hours**

Risks: Provocation of Acute Glaucoma, allergic reactions, eye infection

The use of eye drops is time consuming for both patient and physician and has inherent side effects

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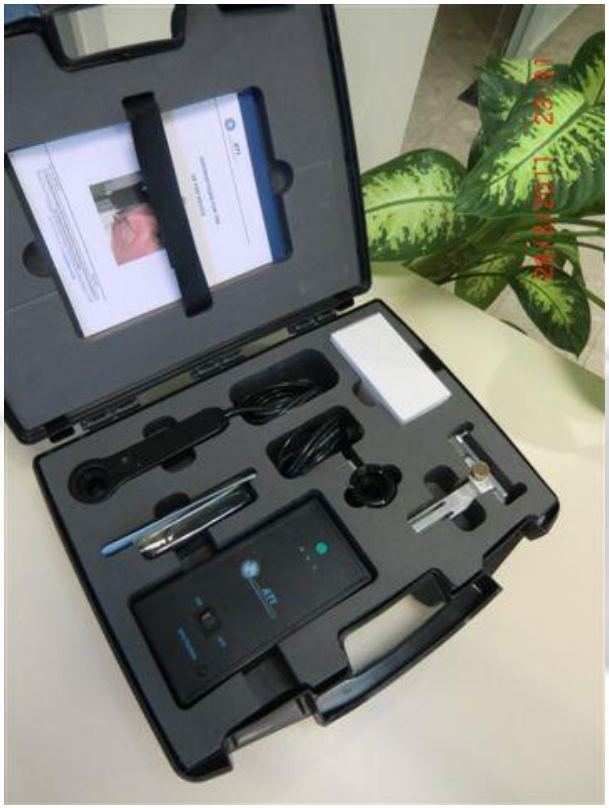
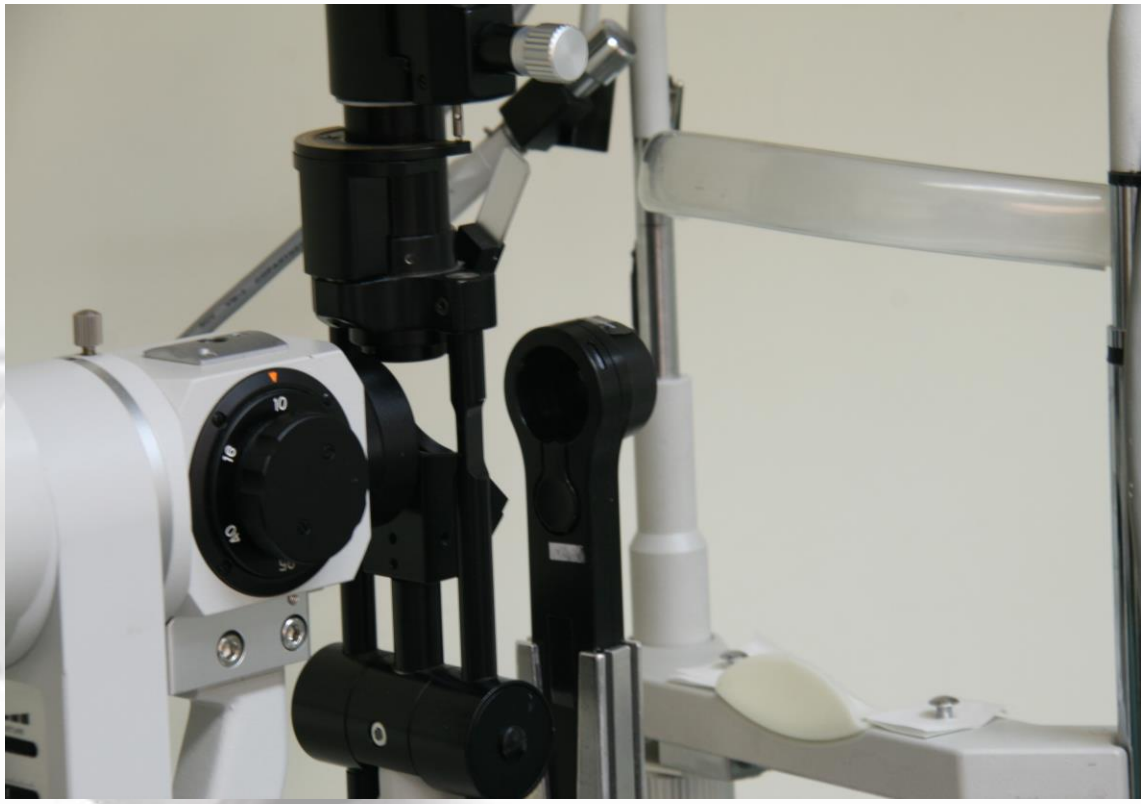
FIM 设备



FIM 设备



FIM 链接裂缝灯

Advantages

Non-contact and non-invasive, easy to operate & use

Fast dilation onset: immediate (30-120 seconds)

Fast eye recovery : immediate (30-120 seconds)

No discomfort or side effects:

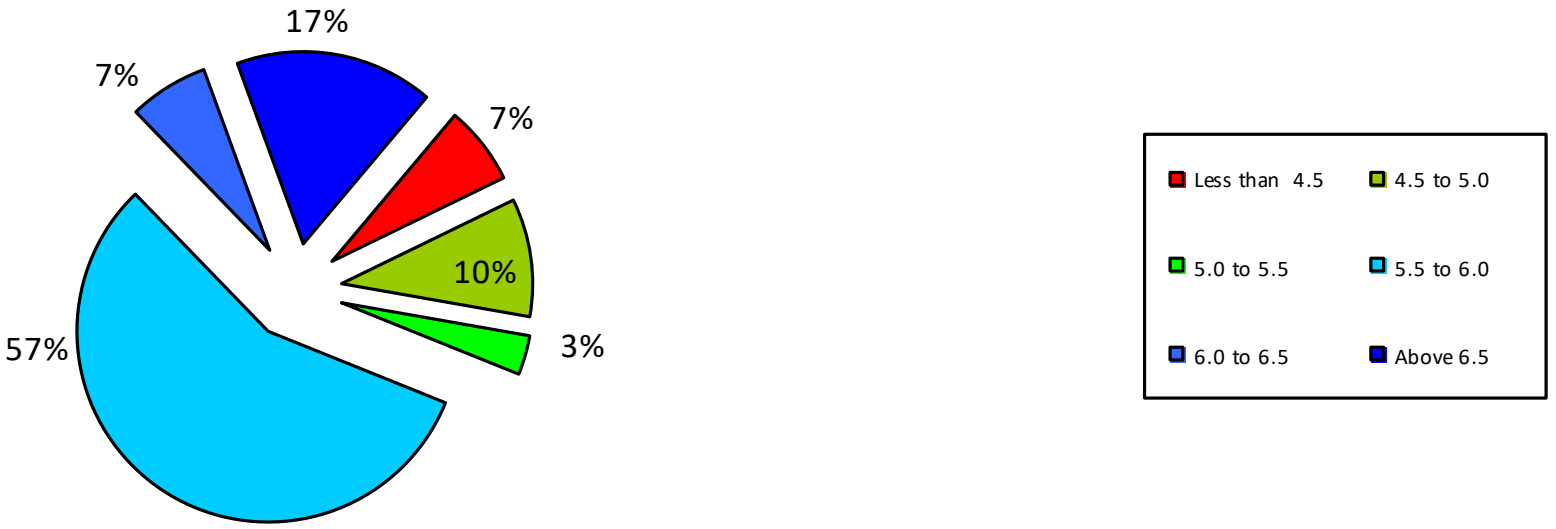
Immediately after eye examination, patient is able to read, work and drive.

The innovative & advanced FIM technology saves both physician and patient time and eliminates the side effects when using eye drops

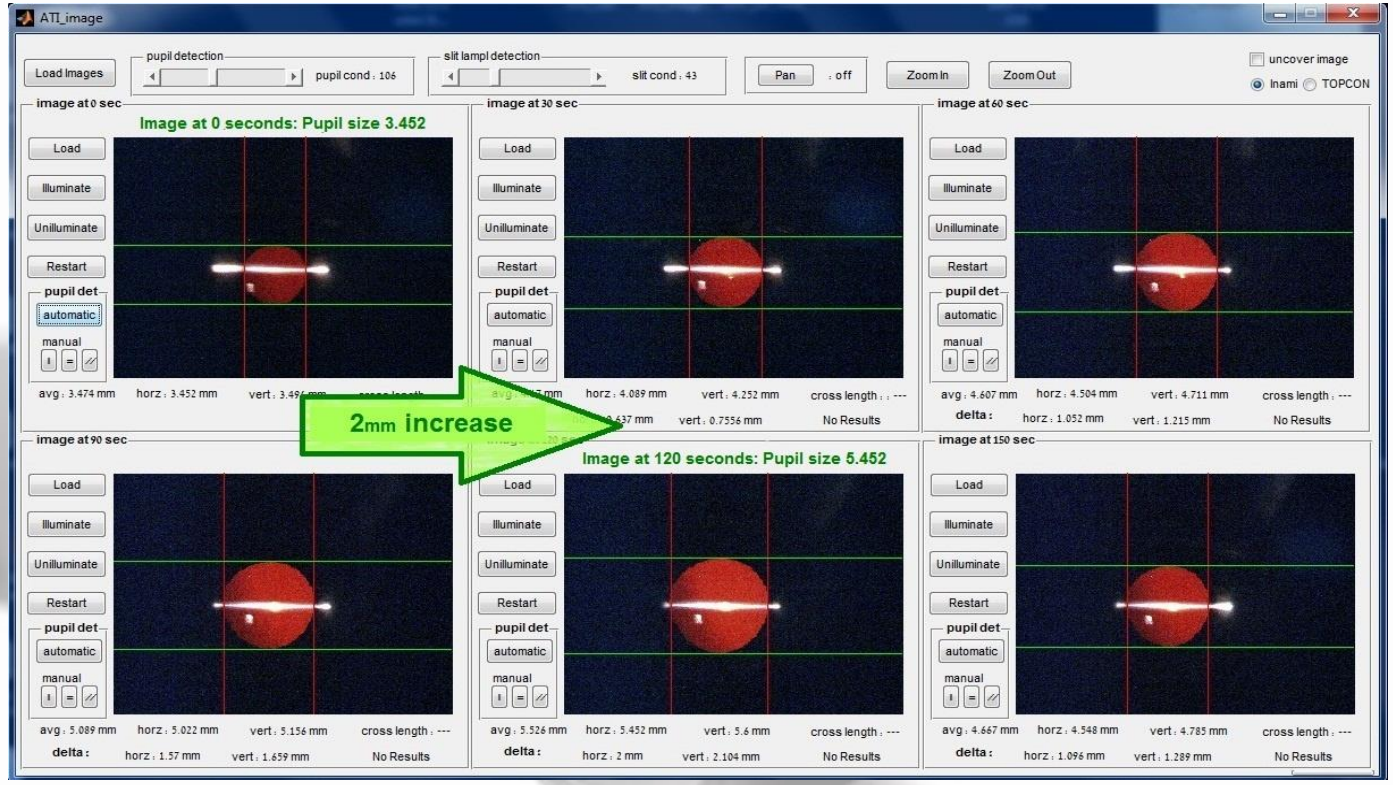
Proof of Concept

Test results clinical trial on 30 healthy volunteers at SERI Eye Clinic, Singapore, 2011

80% of patients were dilated to 5.5 mm or more



Case Study



A fundus examination requires a pupil size of ≥ 4.5 mm.
In this particular case this was achieved with the FIM in 120 seconds.

Competitive Comparison

	Eye Drops	FIM Device
Dilation onset	20-30 minutes	Seconds after start
Recovery rate	Up to 8 hours	Less 2 minutes
Post-exam effects	Blurred vision, stinging, red eyes, unable to work or drive up to 8 hours	None
Elderly	Difficult to use	Easy to use
Patient safety	Provocation of acute glaucoma, allergic reactions, eye , infections	No side effects

Without side effects: patient comfort and safety improve.
With shorter exam times: more patients can be seen.

CE & AMAR Certification + ISO 13485

CERTIFICATE

Number: 3900644CE01

CE MARKING OF CONFORMITY
MEDICAL DEVICES

Issued to:
ATI - Advanced Medical Technologies Ltd
11b Hayetzira St.
Raanana 43663
Israel

For the product category:

Field induced Mydriasis devices to provide pupil dilation

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EC-Directive which apply to them:

0344

Documents, that form the basis of this certificate:

Certification Notice 3900611CN, initially dated October 12, 2012
Addendum, initially dated October 12, 2012

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant provisions of 'Besluit Medische Hulpmiddelen', the Dutch transposition of the Directive 93/42/EEC of June 14, 1993 concerning Medical devices, including all subsequent amendments, and that for the above mentioned product category the Conformity Assessment Procedure Annex V in combination with Annex VII for class IIa products, is executed by the Manufacturer in accordance with the provisions of the Council Directive 93/42/EEC of June 14, 1993. The necessary information and the reference to the relevant documentation, of the products concerned and the assessments performed, are stated in the Certification Notice which forms an integrative part of this certificate.

This certificate is valid until: October 1, 2015
Issued for the first time: October 12, 2012

DEKRA Certification B.V.

drs. G.J. Zoetbrood
Managing Director

ing. A.A.M. Laan
Certification Manager

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All testing, inspection, auditing and certification activities of the former KEMA Quality are an integral part of the DEKRA Certification Group.

DEKRA Certification B.V. Utrechtseweg 310, 6812 AR Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
T +31 26 356 2000 F +31 26 352 5800 www.dekra-certification.com Company registration 09065086

ADDENDUM

Belonging to certificate: 3900644CE01 1/1

CE MARKING OF CONFORMITY
MEDICAL DEVICES

Field induced Mydriasis devices to provide pupil dilation

Issued to:
ATI - Advanced Medical Technologies Ltd
11b Hayetzira St.
Raanana 43663
Israel

This certificate covers the following product(s):

FIM (Field Induced Mydriasis), model M01 (Class IIa)

Initial date: October 12, 2012

DEKRA Certification B.V.

drs. G.J. Zoetbrood
Managing Director

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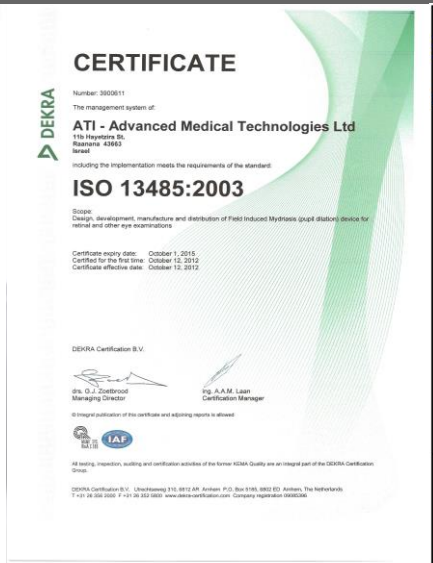

CERTIFICATE

Number: 3900611

The management system of:

ATI - Advanced Medical Technologies Ltd
11b Hayetzira St.
Raanana 43663
Israel

including the implementation meets the requirements of the standard:

ISO 13485:2003

Scope:
Design, development, manufacture and distribution of Field Induced Mydriasis (pupil dilation) device for retinal and other eye examinations

Certificate expiry date: October 1, 2015
Certified for the first time: October 12, 2012
Certificate effective date: October 12, 2012

DEKRA Certification B.V.

drs. G.J. Zoetbrood
Managing Director

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Ministry of Health
Health Technology and Infrastructure Administration
Medical Devices Department

משרד הבריאות
המנהל לטכנולוגיות רפואיות ותשתיות
היחידה לאביזרים ומכשירים רפואיים

מדינת ישראל
STATE OF ISRAEL

אישור רישום בפנקס האביזרים והמכשירים הרפואיים

ניתן בזאת אישור , כי בהתאם לבקשת רישום מס : 26240001
האביזרים / מכשירים רפואים (אמ"ר) הבאים :

שם האמ"ר	FIM – Field Induced Mydriasis
יעוד האמ"ר	FIM is a drug-free, non-invasive device, inducing a temporarily and quickly pupil dilation (mydriasis) without a need for eye drops, and with rapid recovery
התויה	1. עיניים - FIM is a drug-free, non-invasive device, inducing a temporarily and quick pupil dilation (mydriasis) without a need for eye drops, and with rapid recovery
שם בעל רישום וכתובתו	ATI ADVANCED MEDICAL TECHNOLOGIES LTD. ; 11B HAYETZIRA ST. RAANANA, 43663 ISRAEL
שם יצרן וכתובתו	ATI ADVANCED MEDICAL TECHNOLOGIES LTD. ; 11B HAYETZIRA ST. RAANANA, 43663 ISRAEL
שם אתר יצור וכתובתו	א.פ.ח.ע.ש ; קרן היסוד 15, סירת הכרמל ; ישראל
שם ספק וכתובתו	43665 ISRAEL ; אדקום ; התדהר 17, רעננה

התניות

הנחיות
DEKRA: לפי הוראות היצרן שאושרו ע"י גוף המאשר:
- תוגשה שלוש חוות דעת לצורך הארכת תוקף הרישום - אחת מכל מרכז שבו נעשה שימוש במערכת
- אישור בהתאם לאישור CE ומערכת איכות בתוקף
- מאושר לשימוש בהתאם להוראות היצרן כפי שאושרו על ידי הגוף המאשר הכל', בבית חולים ומרפאה בלבד
- השימוש מוגבל לרופאים שהודרכו והוסמכו ע"י היצרן
התניות לרישום זמני
- הגבלות נוספות: לצורך הארכת תוקפו של רישום זה יש להציג חוות דעת כמפורט לעיל

נרשמו בפנקס האביזרים והמכשירים הרפואיים במשרד הבריאות
תוקף האישור לשיווק האמ"ר הינו ליעדים ולהתוויות המתוארים לעיל בלבד
האישור בתוקף עד :31/01/2015

מדינת ישראל משרד הבריאות אגף אביזרים רפואיים MEDICAL DEVICES MINISTRY OF HEALTH STATE OF ISRAEL

29/11/2013

ד"ר נדב שפר
מנהל היחידה לאמ"ר

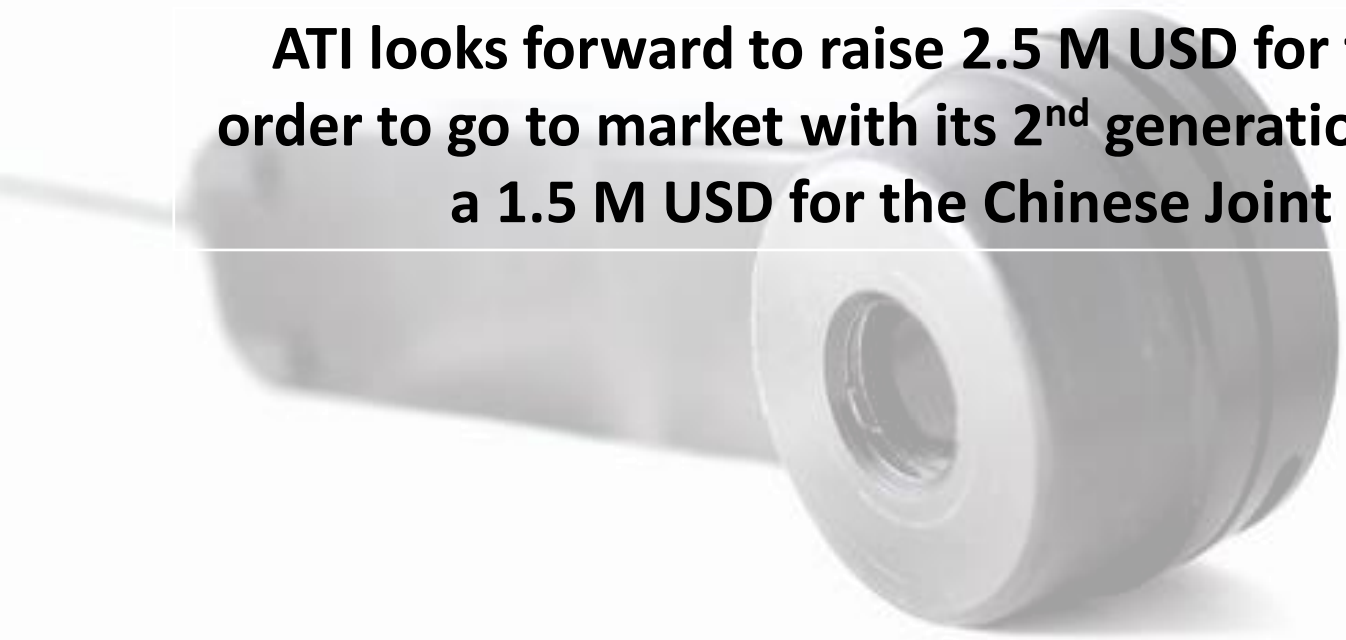
חתימה | תאריך חתימת האישור | שם ותפקיד המאשר

The ATI Managing Team

- **Professor Liora Katzenstein** – Chair Person
- **Mr. Tomas Mendelsohn** – Co-founder, CEO & Board Director
- **Mr. Ilan Ron** – Co-founder & VP R&D
- **Mr. Yoram Wilamovsky –** Board Director
- **Mr. Yaacov Cohen -** Board Director

Funds Needed

ATI looks forward to raise 2.5 M USD for the company in order to go to market with its 2nd generation FIM device and a 1.5 M USD for the Chinese Joint Venture.



Thank You

For More Information
Please contact:

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mailto: tomasm@ati-amt.com
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