REGAINING VISION, REGAINING LIFE

SG-CIB

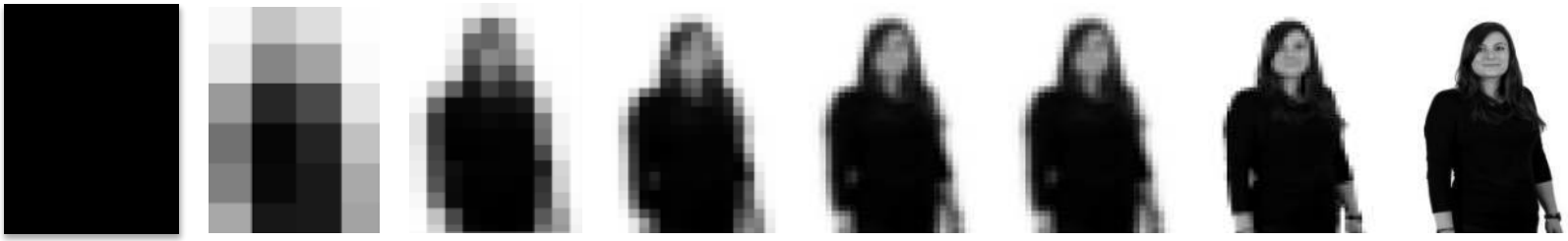
Healthcare & Biotechnology Conference

Disclaimer

This document contains information on Pixium Vision's markets and competitive position, and more specifically, on the size of its markets. This information has been drawn from various sources or from the companies own estimates. Investors should not base their investment decision on this information. This document also contains certain forward-looking statements. These statements are not guarantees of the Company's future performance. These forward-looking statements relate to the Company's future prospects, developments and marketing strategy and are based on analyses of earnings forecasts and estimates of amounts not yet determinable. Forward-looking statements are subject to a variety of risks and uncertainties as they relate to future events and are dependent on circumstances that may or may not materialize in the future. Pixium Vision draws your attention to the fact that as forward-looking statements cannot under any circumstance be construed as a guarantee of the Company's future performance and that the Company's actual financial position, results and cash flow, as well as the trends in the sector in which the Company operate may differ materially from those proposed or reflected in the forward-looking statements contained in this document. Furthermore, even if Pixium Vision's financial position, results, cash-flows and developments in the sector in which the Company operates were to conform to the forward-looking statements contained in this document, such results or developments cannot be construed as a reliable indication of the Company's future results or developments. The Company does not undertake any obligation to update or to confirm projections or estimates made by analysts or to make public any correction to any prospective information in order to reflect an event or circumstance that may occur after the date of this presentation. A description of those events that may have a material adverse effect on the business, financial position or results of Pixium Vision, or on its ability to meet its targets, appears in the sections "Risk Factors" of its "Document de Base" filed with the French Autorité des Marchés Financiers. By attending this presentation or accepting this document, you agree to be bound by the foregoing restrictions set out above.

Pixium Vision: our mission

Pixium Vision's mission is to provide the best-in-class vision restoration systems enabling the blind to regain greater autonomy



Pixium Vision

1 Proprietary systems combining French & international scientific & technological excellence

2 Attractive addressable 1 Billion Euro + market opportunity*

3 Two differentiated systems:

- IRIS® on track for launch in 2015
- PRIMA to further expand the market opportunity after 2018

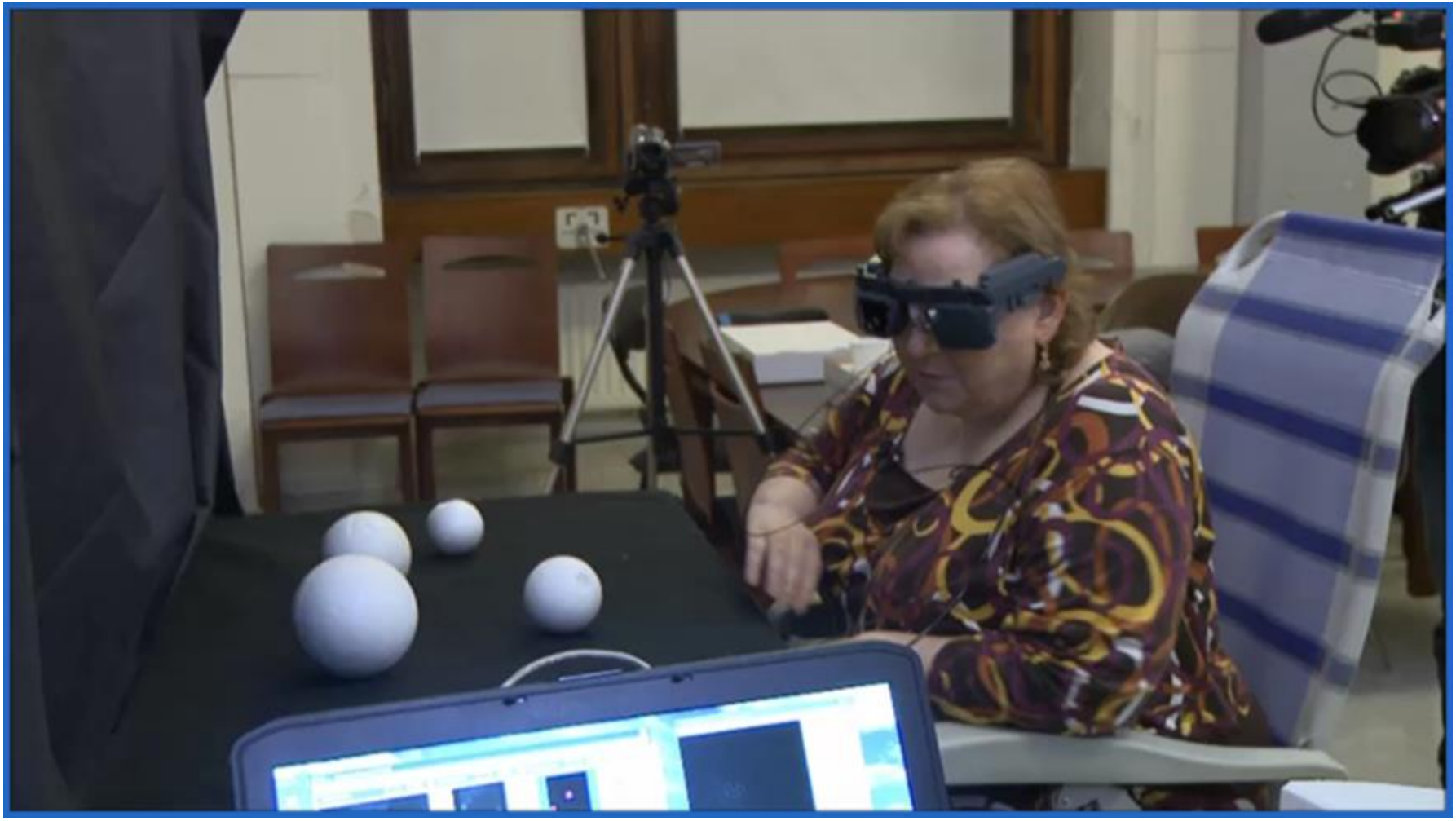
4 Strong and dedicated management

Establish Pixium Vision's position as a leader in Vision Restoration Systems

* Company Estimate

Imagine how much blind people miss out on…

Testimonies of patients participating in the clinical trial



Blindness

Costs and target pathologies

Solving blindness represents a major market opportunity

285 million people in the world are visually impaired

40–45 million people in the world are totally blind

In the US and Europe, blindness costs exceed tens of billions of USD per annum

No treatment exists for blind patients

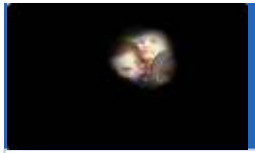
Retinitis Pigmentosa (RP) and

Age-Related Macular Degeneration (AMD)

are major causes of blindness

Sources: World Health Statistics. World Health Organization -http://www.amd.org - NORC Cost of Vision Problems: The Economic Burden of Vision Loss and Eye Disorders in the United States -Study commissioned by Prevent Blindness in America and conducted by University of Chicago -European Forum Against Blindness (EFAB)

Two major pathologies lead to photoreceptor degeneration and ultimately, blindness



Retinitis Pigmentosa (RP)

- Genetic disease
- Blindness occurrence: ~ 35 - 40 years old
- Worldwide prevalence: 1.5 to 2 million
- Prevalence in the US + EU: 350,000 - 400,000
- Incidence (US + EU): 15k-20k patients annually

Age-related Macular Degeneration (AMD)

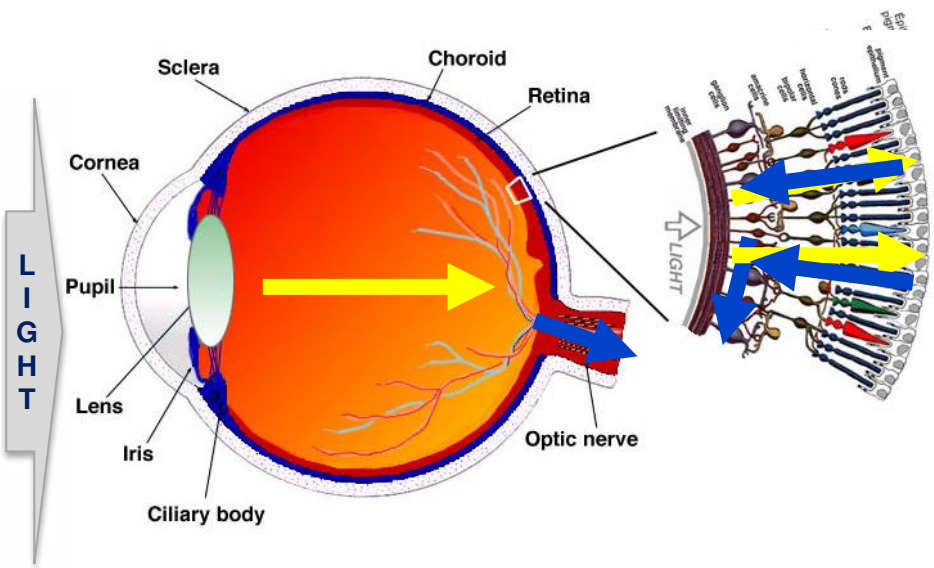
- Age-related disease
- Later blindness occurence: 70+ years old
- Worldwide prevalence: 12 to 15 million
- Prevalence in the US + EU: 4 million
- Incidence (US + EU): 350k - 400k patients annually

Retinitis Pigmentosa is Pixium Vision's initial target market

Source: 2012 World Health Organization – by 2020 there will be 75 million blind people in the world and 314 million partially-blind people
Fighting Blindness (UK) : 25K in UK and over 2M worldwide
CentralSight fact sheet End-Stage Age-related Macular Degeneration

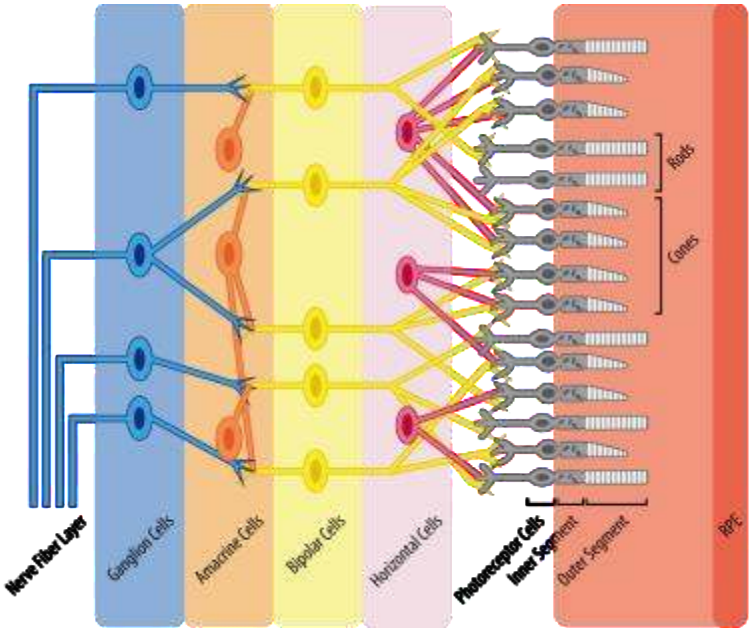
The loss of the photoreceptor function is a major cause of blindness

The eye transforms light into electric signals



- Photoreceptor cells convert light into signals
- The human retina contains 6 million cone cells responsible for central vision

Photoreceptor degeneration does not affect the rest of the retina



- RP and AMD are linked to photoreceptor degeneration
- **However, bipolar cells, ganglion cells and downstream visual pathways remain INTACT and FUNCTIONAL in the vast majority of RP and AMD patients**

With its technology, Pixium Vision is well positioned in the fast growing neuromodulation market

What is Neuromodulation?

- Induction of biological responses from electrical stimulation on nerves or zone where nerve activity is affected
- $5Bn+ market by 2018 implying a high double digit growth rate (around 15%)

Pathology/ Indication	Neuromodulation Technique	US prevalence
Deafness	Cochlear Implants	1,000,000
Parkinson's Disease	Deep Brain Stimulation	216,000
Depression	Vagas Nerve Stimulation Deep Brain Stimulation	775,000
Blindness Pixium vision	Vision Restoration Systems	~175,000 RP ~ 200,000 AMD

Source: NIH US Net prevalence patient data

Existing therapies

Emerging therapies

Pixium vision

Pixium Vision

The convergence of excellence

The Pixium Vision story relies on the convergence of technology and financing

Scientific Innovation → **Product Development and Clinical studies** → **First product launch**

IMI

Assets & Patents

c.€50m invested 2002-2012

INSTITUT DE LA VISION

Research & Patents

5-6 years of research

December 2011

Pixium vision

Patents

5-6 years of research

May 2012

Financial Round €24.5m 2 Tranches

Abingworth – investors in life sciences & healthcare; bpifrance; OMNES CAPITAL; Seventure; SOFINNOVA

June 2013

Start of clinical trials for IRIS®

June 2014

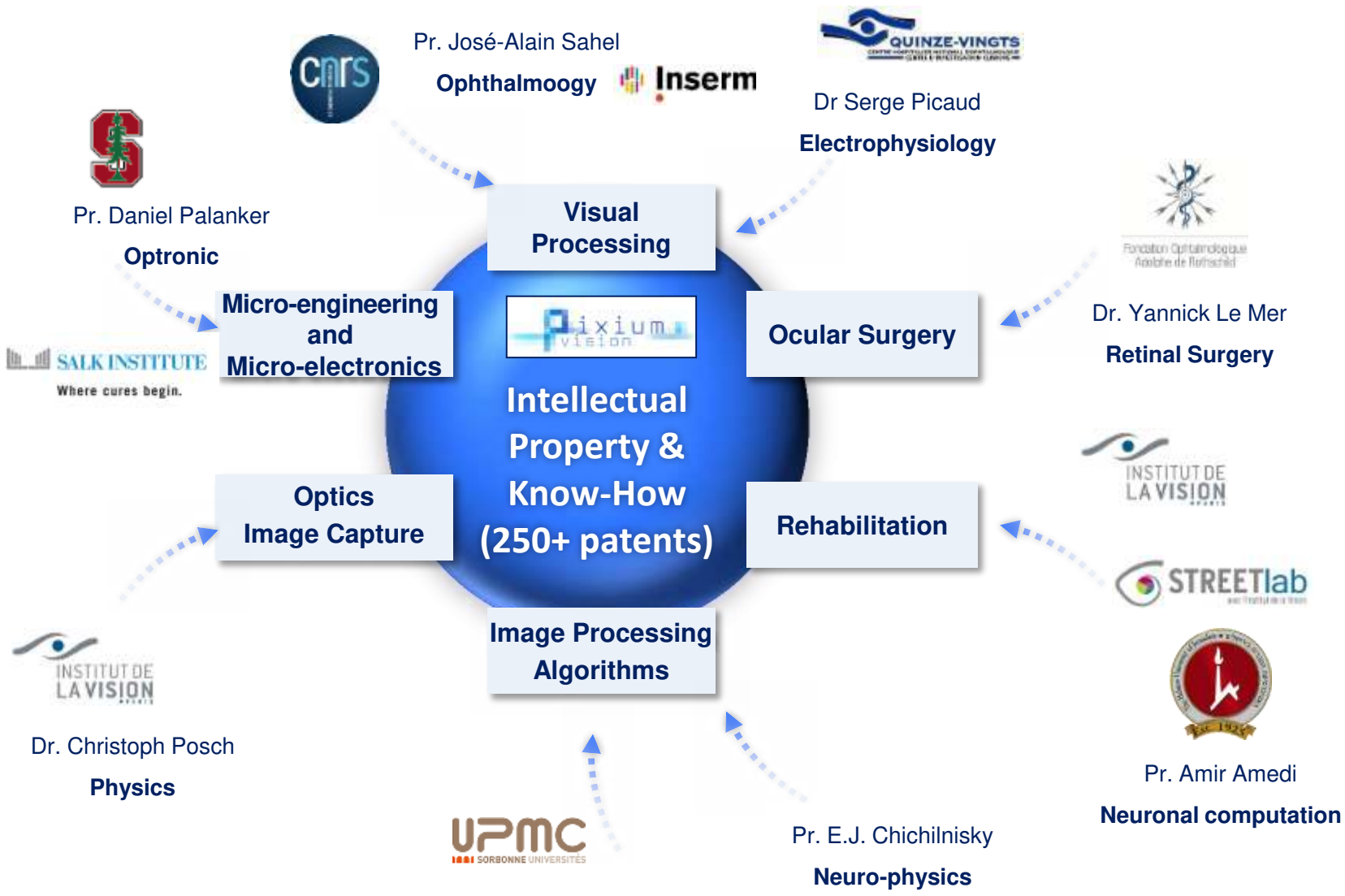
IPO

Gross proceeds of €39.5m

2015

CE Mark & First Sales

Pixium Vision is supported by French excellence and global expertise

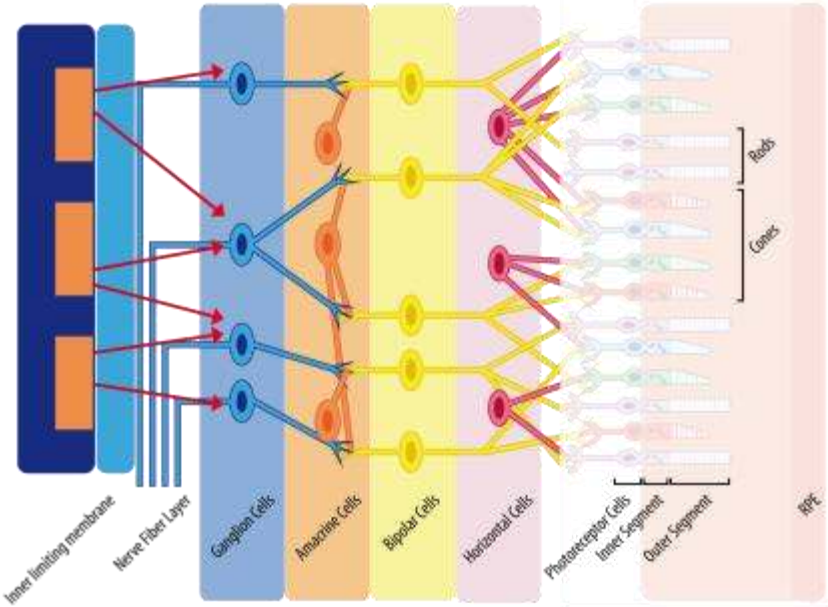


IRIS®

A state of the art Vision Restoration System

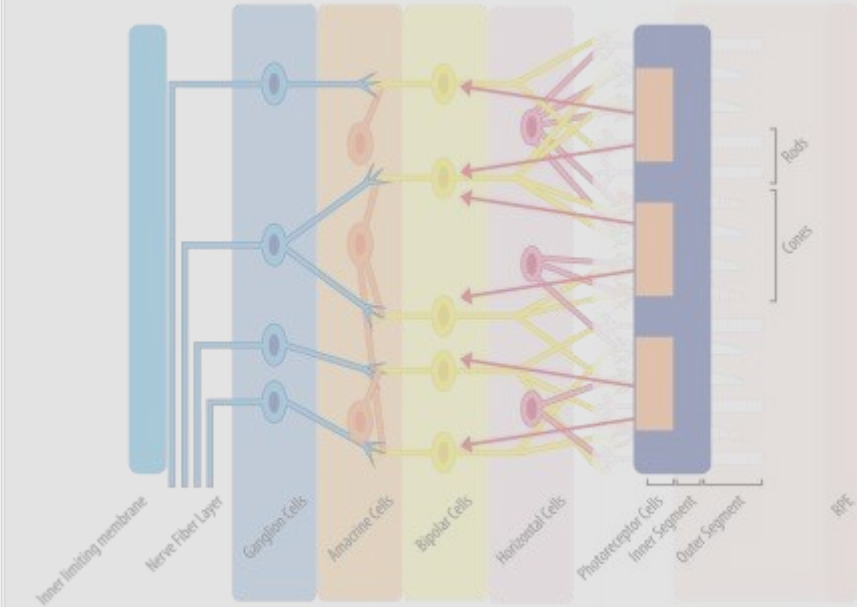
Pixium Vision is developing two differentiated Vision Restoration Systems

EPI-RETINAL STIMULATION



IRIS®

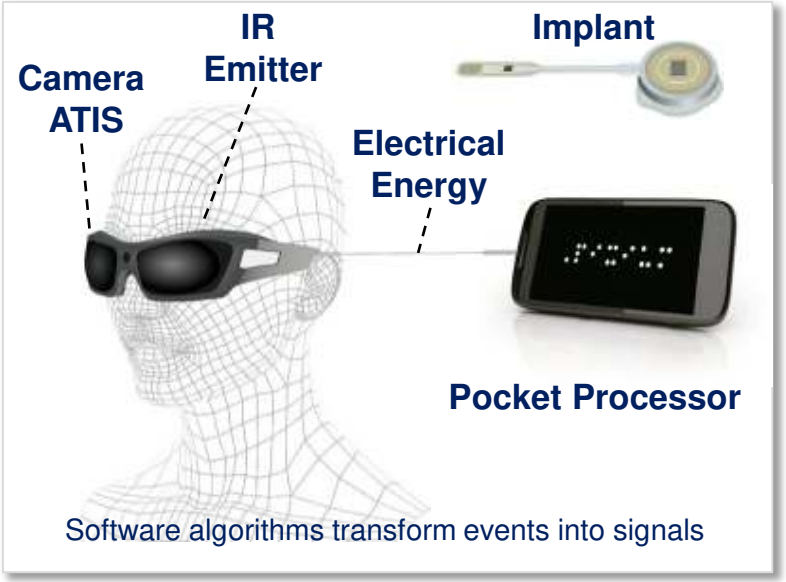
SUB-RETINAL STIMULATION



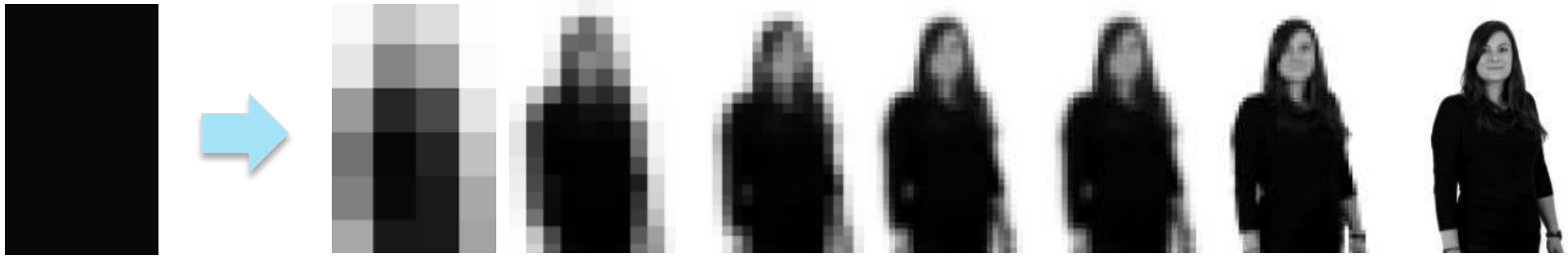
PRIMA

The IRIS® Vision Restoration System

A technically advanced system designed to deliver important clinical benefits

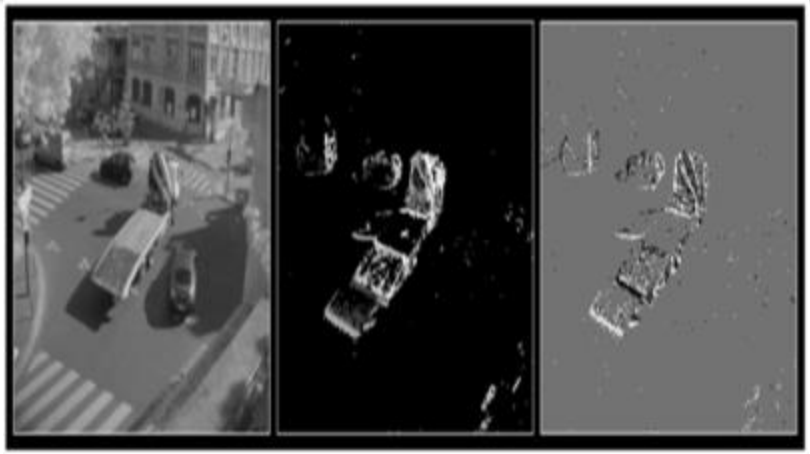


Initial goal is to deliver light and shape perception, and to localize objects giving the patient the ability to negotiate an unfamiliar environment



A unique proprietary *event-based* camera, functioning like the human eye

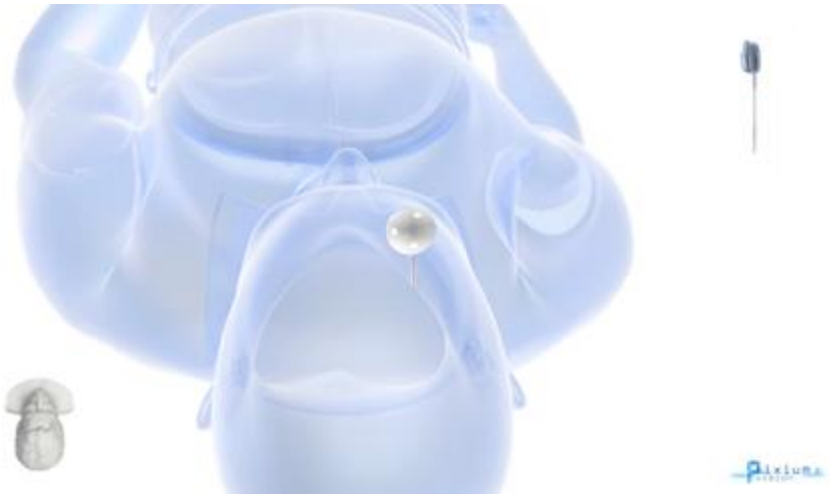
Our eyes only see changes in our environment (contrast, intensity, luminance, etc.)
Photoreceptors are activated independently



Camera Features	Benefits
• Breakthrough bio-mimetic camera	• Replicates normal vision in real time
• Neuromorphic – asynchronous, event-based: light is encoded into asynchronous impulses (-1,0,+1)	• Reduces energy consumption and brandwith
• Output relates directly to signals observed in the corresponding levels of biological retinas	• Visual information can be directly understood by the visual cortex

IRIS®, a technically advanced epi-retinal implant

- Epi-retinal implant
- 150 electrodes in the commercial version
- Electrical power by induction
- Simple surgical procedure
- Compatible with
  - Next-generation vision sensors
  - Signal-processing algorithms
- Well tolerated by patients so far



IRIS® : A technically advanced and differentiated VRS

	Device Features	IRIS®	Main Competitor
Technology	Camera	Event Based	Frame Based
	Patient Programming - Tuneability	Yes	No
	Number of Electrodes	150 electrodes	60 electrodes
Surgery	Surgical Procedure	2.5 hours	Up to 4 hours
	Explant and Replacement	Yes	Replacement not proven

IRIS®: A clear path to market

Aiming for a leading market position

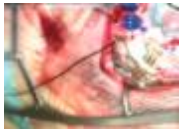
IRIS® : Continue to build evidence for CE Mark

Ongoing Clinical Trial

1. Incidence, severity and duration of all ***adverse events*** at 4, 6, 9, 12 and 18 months
2. Assessment of the capability of patients to ***perform visual tasks*** with and without the device at 4, 6, 9, 12 and 18 months

Regulatory Path

Implantation Surgery



Implant Activation (after 4 weeks)

Rehabilitation Program

CE Mark 2015

CE

Clinical Centers

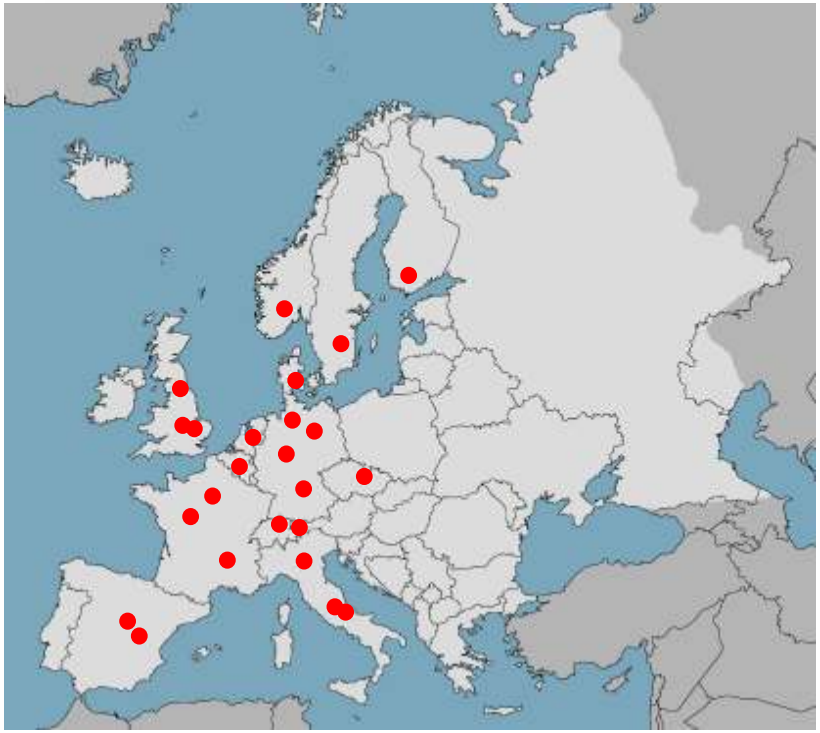
- **Paris & Nantes**
- **Graz**
- **Hamburg**

Rehabilitation Program

- Programs tailored for each patient
- Rehabilitation programs will enable further software improvements
- Patients' vision improves during the course of their rehabilitation program

A lean and specialized commercial organization

25 to 30 key ophthalmic surgery centers in Europe



These centres give access to ~80% of qualifying patients*

Market development process

Ongoing:

- KOL engagement
- Discussions with patient associations
- Participation in major scientific and medical conferences

Country/market assessments to select and prioritize centers

Recruitment of a lean internal technical/clinical specialist sales team focused on:

- Commercial & educational activities
- Training & support of orthoptists

Sales team to reach a peak of 2 to 3 team members per country & sales admin employees

* Company Estimate

Staged launch planned for IRIS®

1

European IRIS® Launch

Wave 1

IRIS® CE Mark 2015

First European Sales Launch: 2015

Launch Accounts

- IRIS® clinical sites subject to reimbursement
- Expand across Wave 1 launch countries

2

European IRIS® Launch

Wave 2

Second European Launch: 2016

Launch Accounts

- Subject to reimbursement
- Expand across Wave 2 countries

3

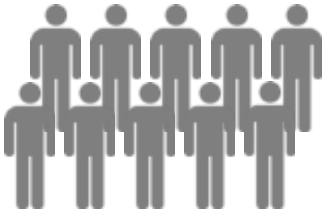
US IRIS® Launch

US Launch: early 2018

IRIS® path to the US market

1

Gather results from European clinical trial



2

File an Investigational Device Exemption (IDE)

- Planned for early 2015
- Pixium Vision anticipates that FDA will require clinical results from at least 30 patients with 2 years follow-up

3

Obtain Pre-Market Approval (PMA)

- US launch of IRIS® to start early 2018

IRIS® pricing and reimbursement drivers

Cost of blindness

Competitor reimbursement benchmark

€78,000	€100,000
€96,000	$144,000

Quality of clinical data

Comparative MedTech therapies

and software improvement

Pricing & Reimbursement Drivers

Software upgrades and support services will generate an additional revenue stream

Software & support services sales

Illustration with key role of software evolution in Cochlear implant market

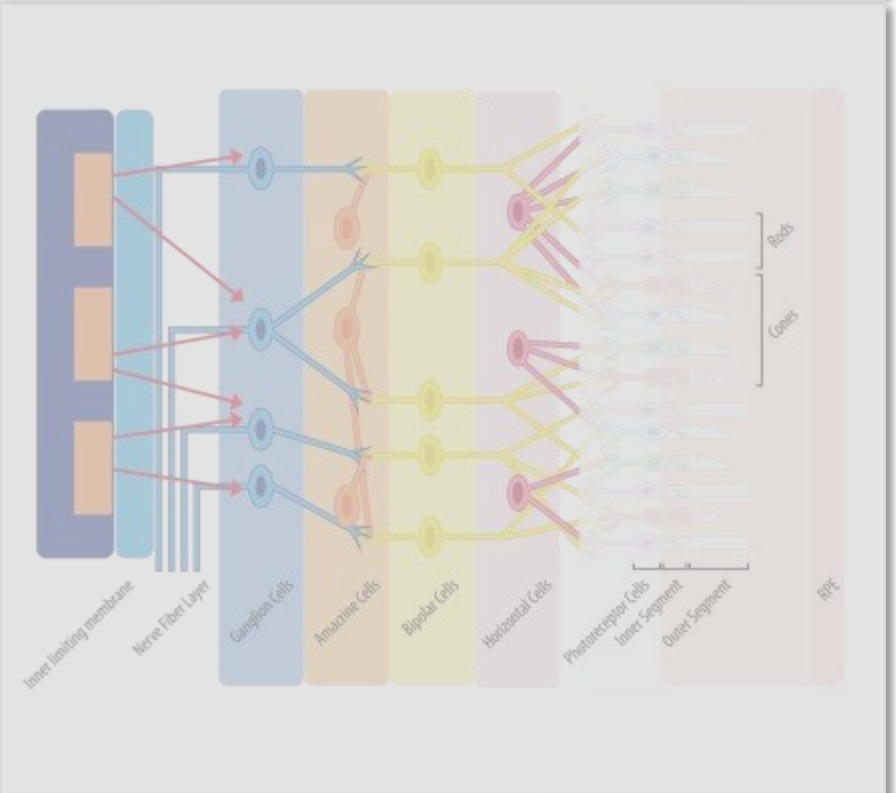
- **Software improvement needed to:**
  - Improve performance and patient benefit
  - Enhance product life cycle management
- **Software improvement to leverage implant clinical utility:**
  - Develop GPS, reading and other applications

PRIMA Vision Restoration System

Building on IRIS® leading market position

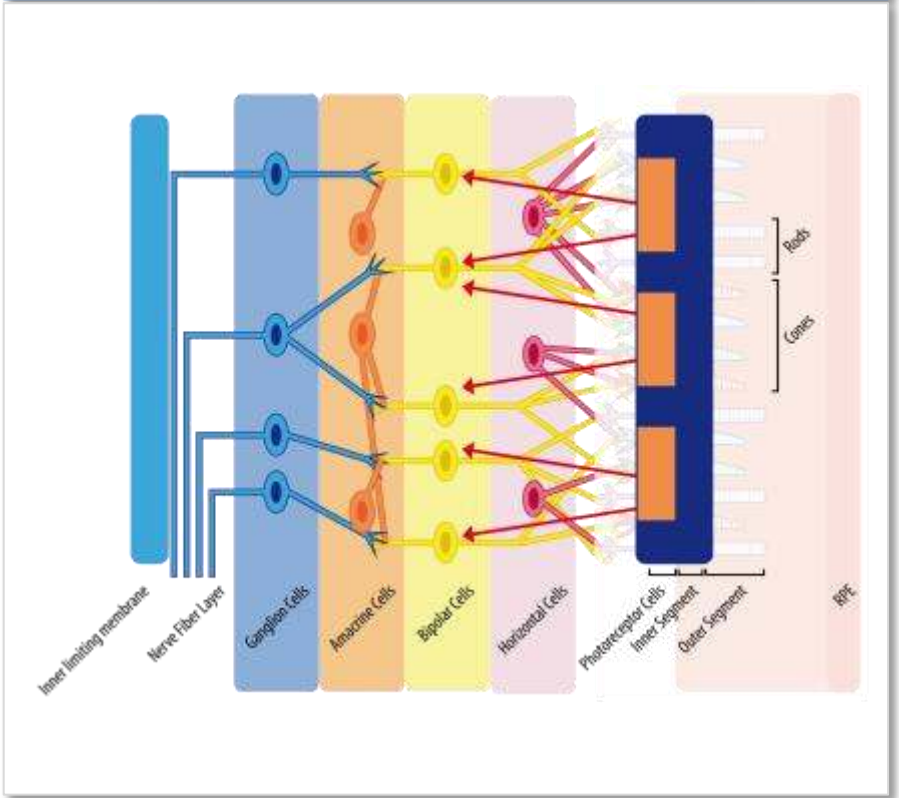
Pixium Vision is developing two differentiated Vision Restoration Systems

EPI-RETINAL STIMULATION



IRIS®

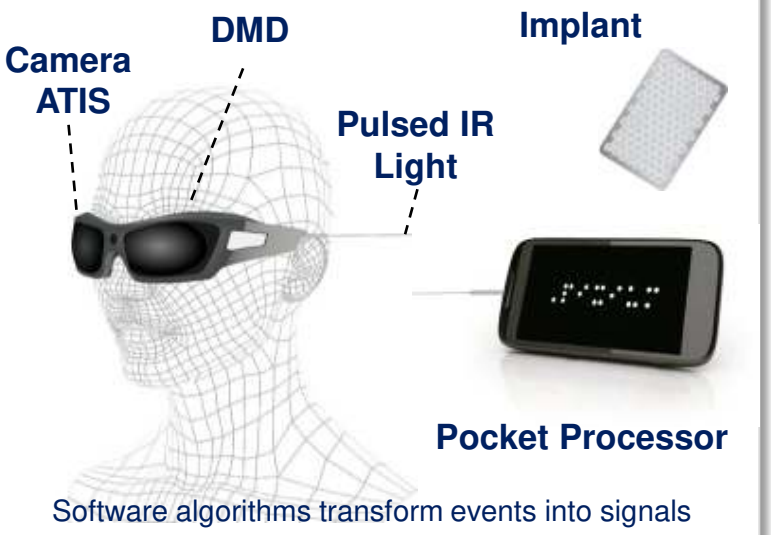
SUB-RETINAL STIMULATION



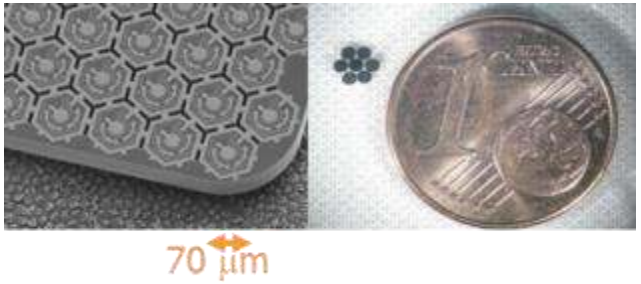
PRIMA

The PRIMA Vision Restoration System

A technically advanced system designed to deliver further clinical benefits

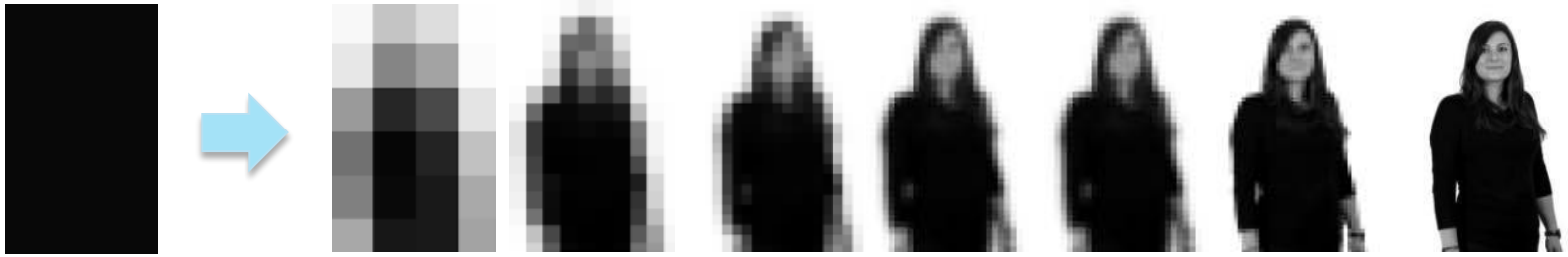


Software algorithms transform events into signals



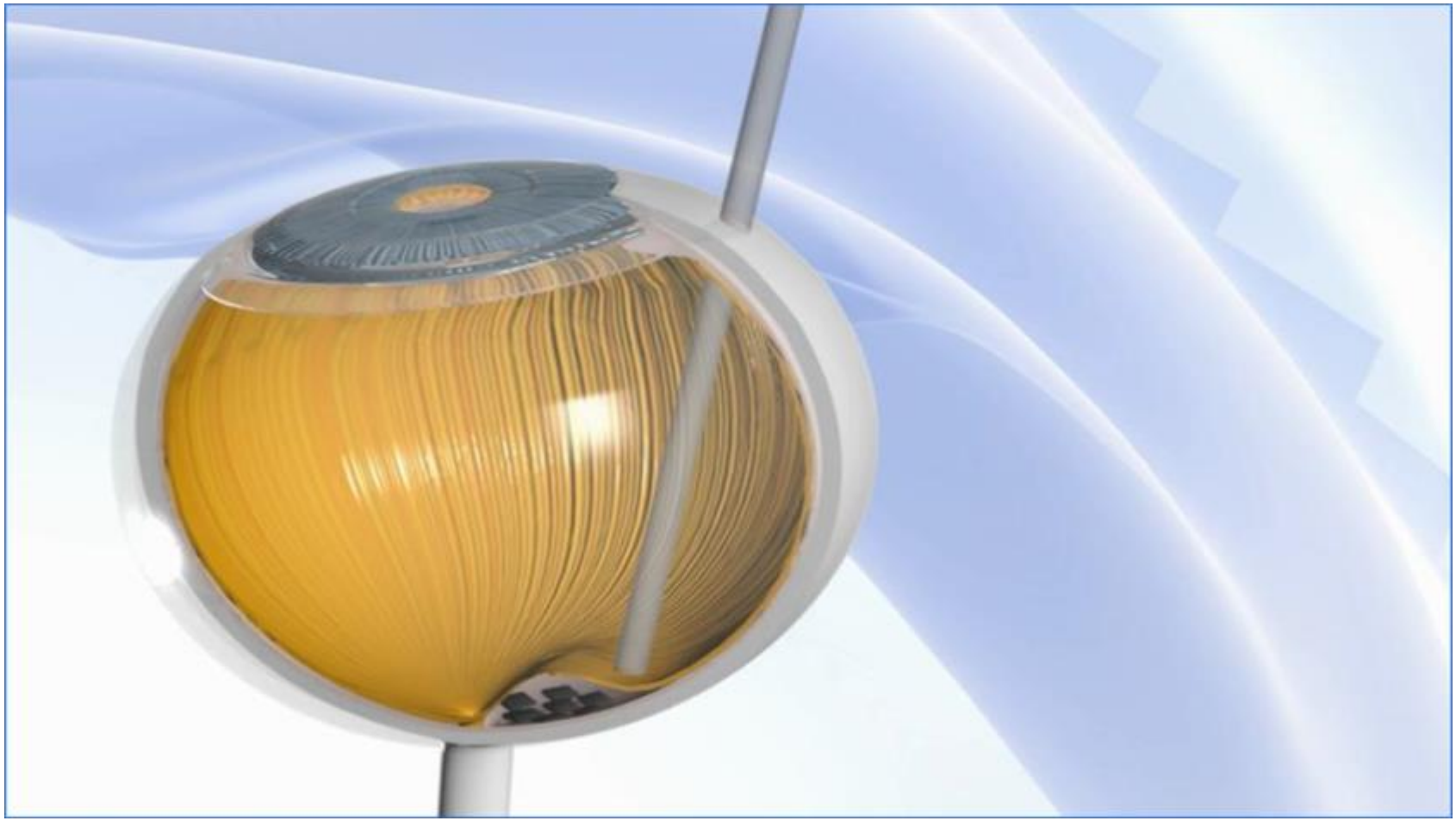
- Physiological signal processing
- Simpler and shorter surgical procedure
- Retinal chips in modules up to 5,000 electrodes
- Advanced processing algorithms
- Reduced energy requirements enabling miniaturization of components

Goal is to deliver improved visual perception to the level of direct facial recognition

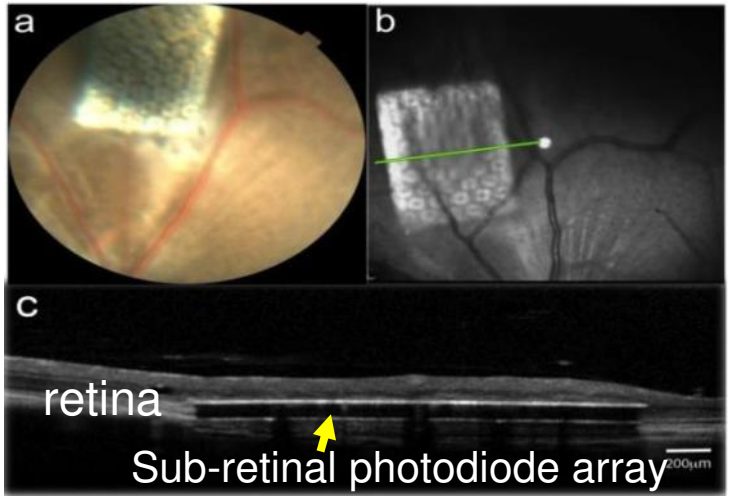


PRIMA, a sub-retinal implant

Sub-retinal implants directly stimulate the retinal cells that used to be connected to the photoreceptors

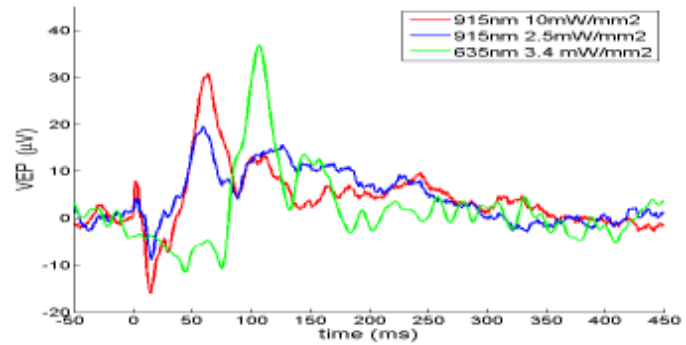


PRIMA: Validated in pre-clinical models

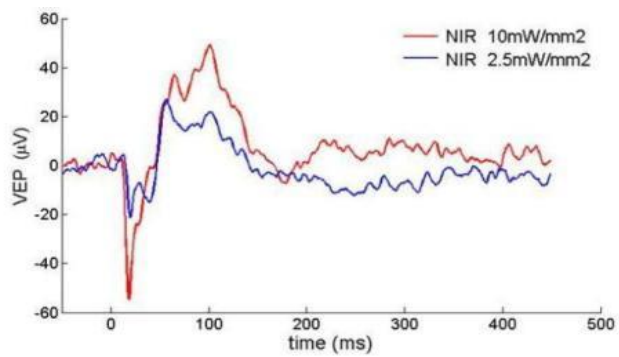


- Proof of concept achieved in rats
- Safety demonstrated in rats and pigs
- Scale-up of manufacturing process ongoing
- First in man expected in 2016

Visual Evoked Potential: Normal rats *



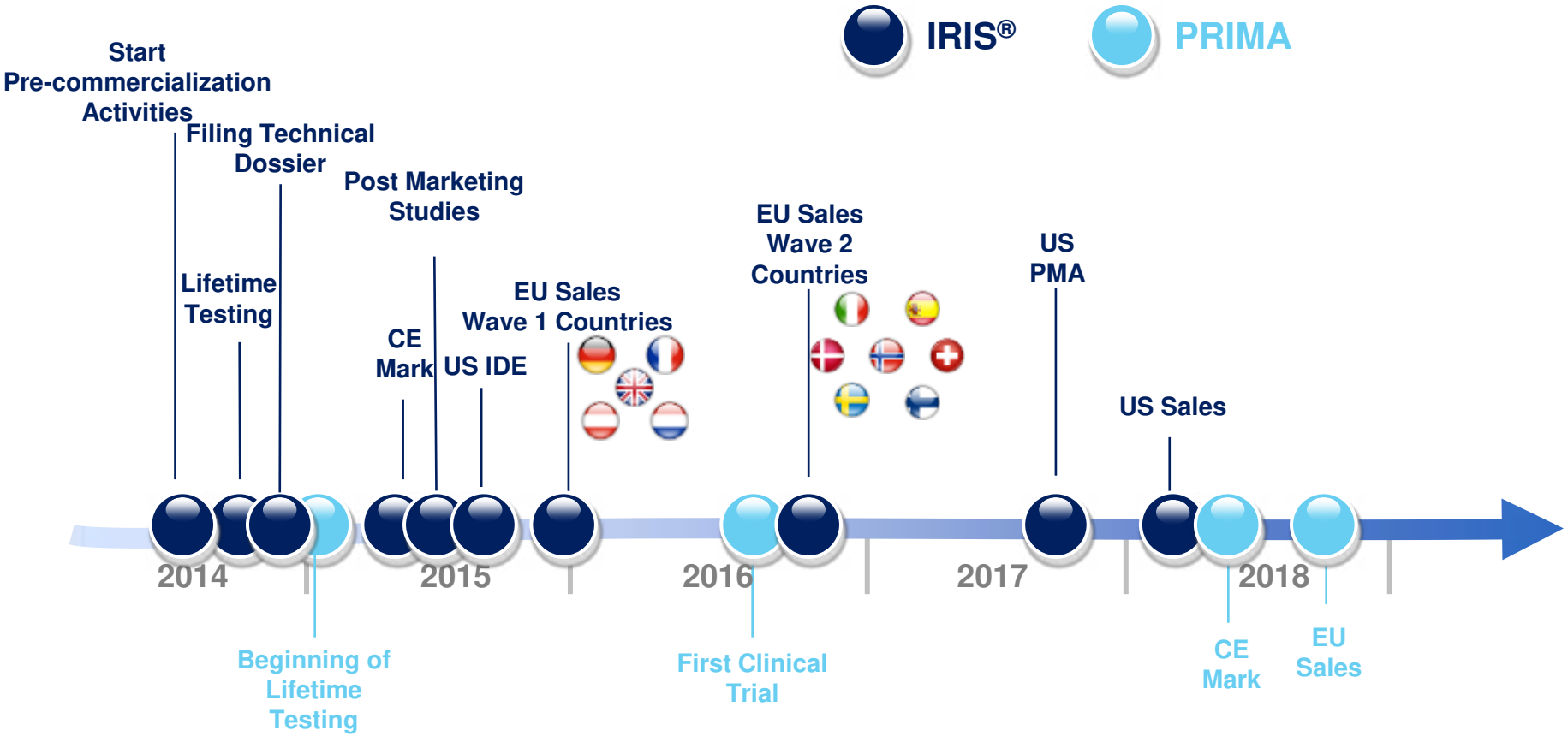
Visual Evoked Potential: Blind rats *



* Ref: *J. Neural. Eng* 9: 046014(2012)

A rich upcoming newsflow

Pixium to complete IRIS® clinical trial and prepare for commercial launch in 2015; Prima to follow



Multiple major value-creation milestones anticipated in the next 2 years

H1 2014 Financials

Reminder IPO use of proceeds

1

IRIS®

Completion of clinical trial and commercialization of IRIS® in Europe

Clinical trial and commercialization of IRIS® in the USA

2

PRIMA

Development of PRIMA: from clinical trials to the obtention of regulatory approvals for commercialization in Europe and the US

Reminder IPO use of proceeds

P&L summary

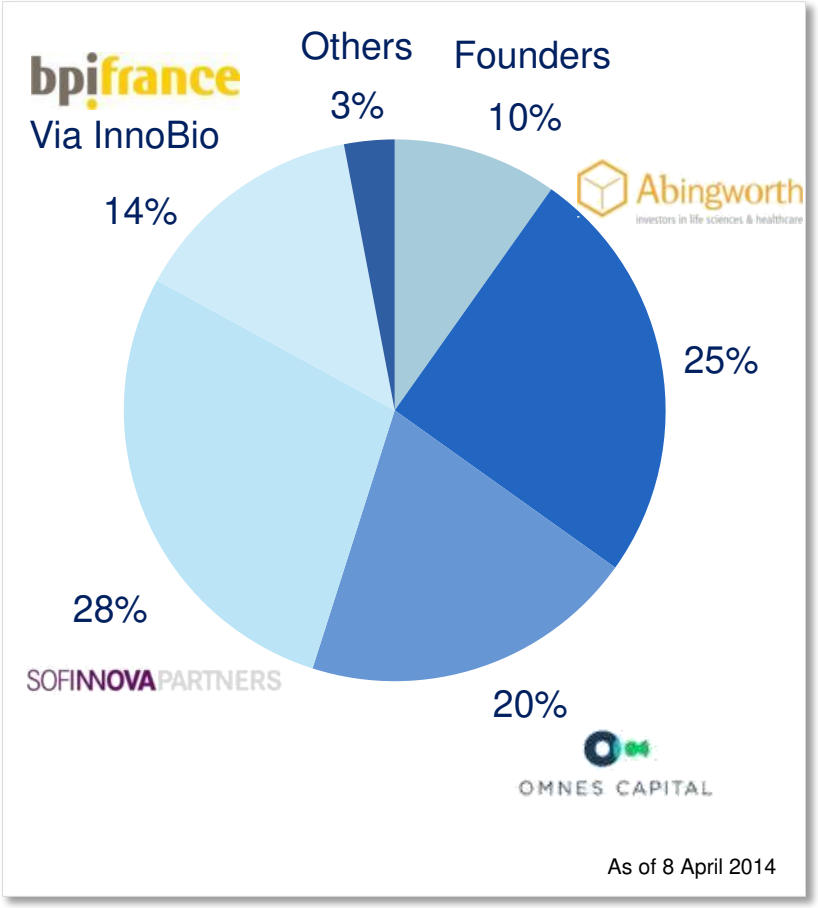
in thousand euros	**S1 2014**	**S1 2013**
Operating income / other income	1 104,1	574,4
Research and Development	4 510,6	2 285,0
General and Administrative	930,1	498,6
Operating income	(4 336,5)	(2 209,2)
Net profit	(4 325,7)	(2 211,3)
Earnings per share	€ (0,62)	€ (0,09)

Cash flow statement summary

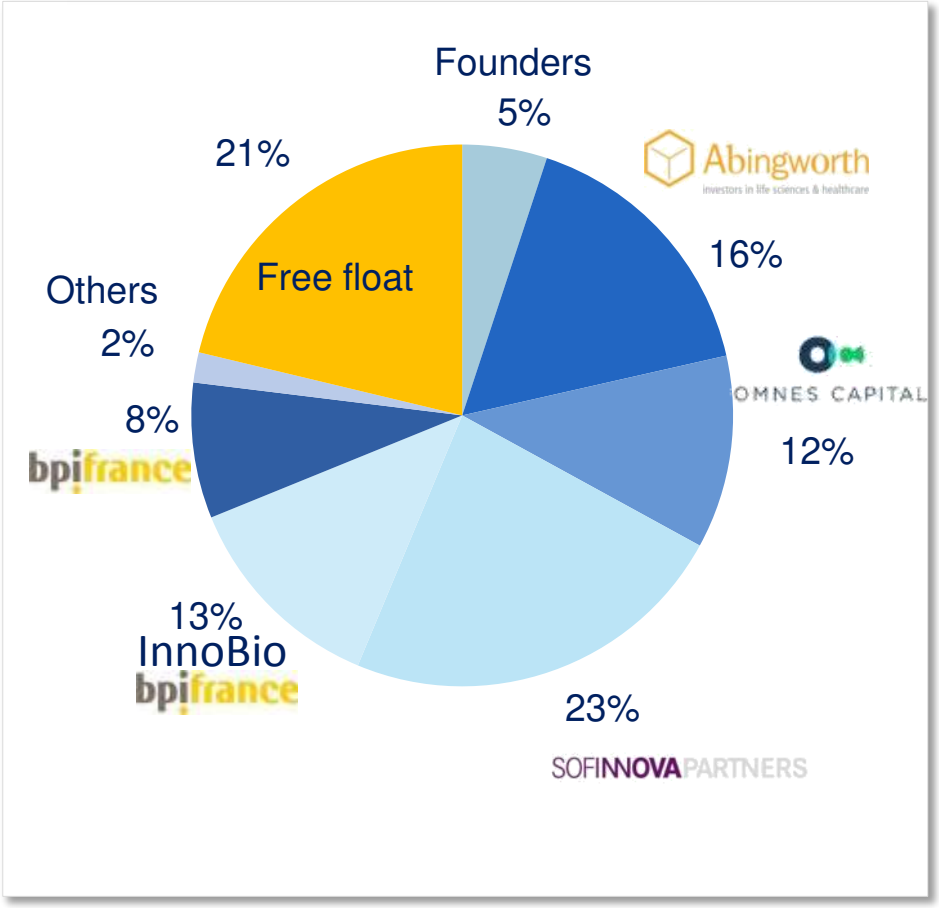
in thousand euros	**S1 2014**	**S1 2013**
Opening cash and cash equivalents	9 420,2	3 088,6
(Decrease) / Increase in cash position	32 383,4	(217,1)
O/W net cash flows from operating activities	(4 530,1)	(2 618,4)
Closing cash and cash equivalents	41 803,6	2 871,5

Shareholder structure

Pre-IPO shareholder structure
On a non-diluted basis



Post-IPO shareholder structure
On a non-diluted basis
(with full exercise of the extension clause and overallotment option exercised at 95.8%)



Competitive landscape

System	Number of Electrodes	Epi-Retinal Or Sub-Retinal	Features & Benefits	Clinical Results	Regulatory Status
Pixium vision	• IRIS® : 50 to 150 • PRIMA : up to 5000	• IRIS®: Epi-Retinal • PRIMA: Sub-retinal	• 2h surgery • Explantable • Neuromorphic Camera • Tunable software	• Short term study on 19 patients • 10 Patients CE mark study ongoing	• CE Mark filing end of 2014
Second Sight	• Argus II : 60 electrodes	• Epi-retinal	• CMOS camera	• Argus I: 6 patients • Argus II: IDE on 30 patients	• Argus-II CE Mark Feb 2011 • FDA HDE Feb 2013
retina implant	• Alpha IMS	• Sub-Retinal	• Visual field of 12° • Non explantable	• 11 patients from 2005 to 2009 • 30 patients CE mark	• CE Mark July 2013
NanoRetina	• 500 electrodes	• Insufficient data	• No camera	• Launch scheduled for 2016	• Pre-clinical phase