Array/atch

MULTI-AWARD WINNING INNOVATIVE DRUG DELIVERY TECHNOLOGY











ADVANCING NEW THERAPEUTICS TO TREAT MILLIONS WORLDWIDE



DerMap - patent protected dissolving microneedle drug delivery platform



Attractive therapeutic applications identified targeting ~\$70bn markets



Breakthrough – dissolving microneedles with 100% drug loading



Multiple exit routes for significant ROI potential



Low development risk – novel delivery of approved medicines



Experienced team assembled

KEY MILESTONES TO DATE

SPUN OUT OF UCC 2024 - €1.34M GRANTS INVESTED PRECLINICAL POC IN ATTRACTIVE LEAD APPLICATIONS

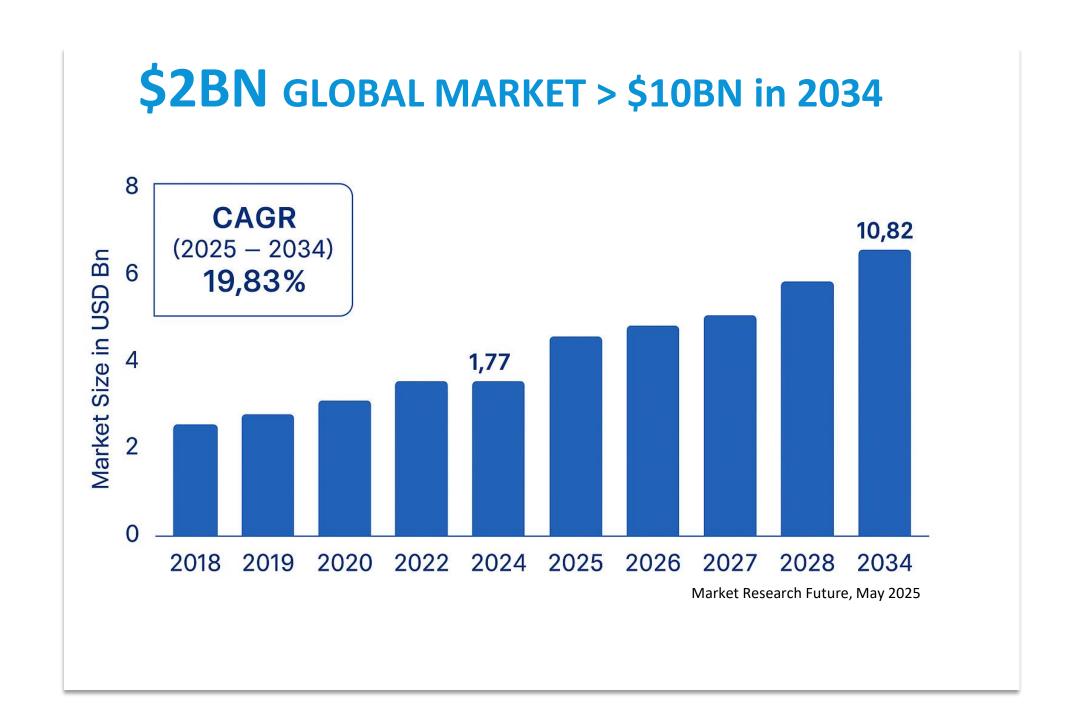
RECOGNISED MULTIPLE AWARDS
ACHIEVED

EXPERIENCED
TEAM AND SAB
ASSEMBLED

RAISING €3M SEED TO ADVANCE TO CLINICAL TRIALS



MICRONEEDLE PATCH DRUG DELIVERY MARKET OVERVIEW

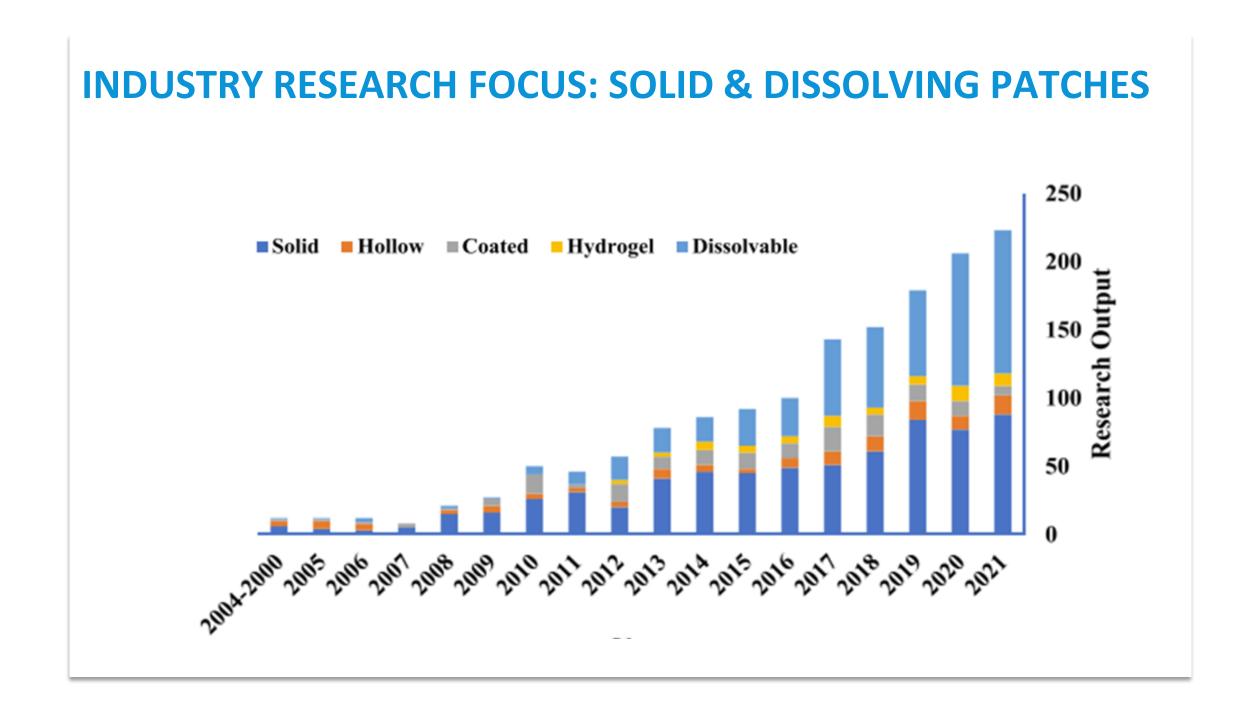


Rationale

- Fear of needles; reduce sharp waste
- Healthcare saving (self administration)
- Accessibility: eliminate of need for cold chain storage

Growth Drivers

- Increase in chronic diseases
- Need to target delivery for greater efficacy and safety
- Cosmetic use; vaccines



Microneedle Types

- Solid microneedles lead the market but are expected to decline due to their inefficient two-step application process
- Dissolving patches offer the advantage of no additional manipulation following insertion

Problem – Need for Polymers

Drug delivery success to date hampered due to requirement for polymers - limits on drug loading; drug incompatibility; lack of validated analytical procedures; safety concerns.



Array DISSOLVING MICRONEEDLE PATCH – COMPETITIVE LANDSCAPE

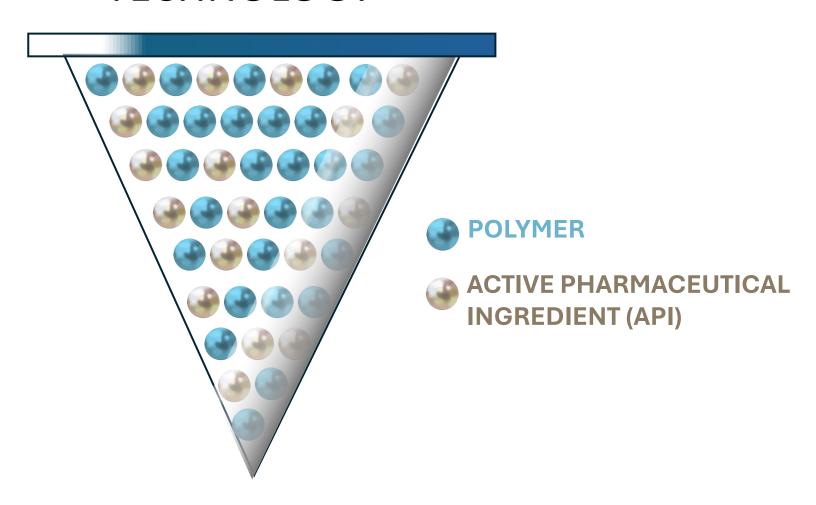
COMPANY	TECHNOLOGY	STAGE	FOCUS	DISSOLVING	POLYMER FREE
Micron	Microarray	Clinical – IIb	Vaccines		X
Vaxess Technologies	MIMIX TM	Clinical – I	Vaccines		X
CosMed Pharmaceuticals	Soluble microneedles	R&D	Cosmetics		X
Vaxxas	HD-MAP	Clinical – I	Vaccines		X
ArrayPatch	DerMap TM	Pre-clinical	Small molecules		

- DerMapTM is highly differentiated the only technology with microneedles made entirely from drug
- Competitors mostly focusing on vaccines
- DerMapTM is a unique platform technology with applications across multiple therapeutic areas already in development including nail fungal infection, skin cancer, migraine and small peptides.



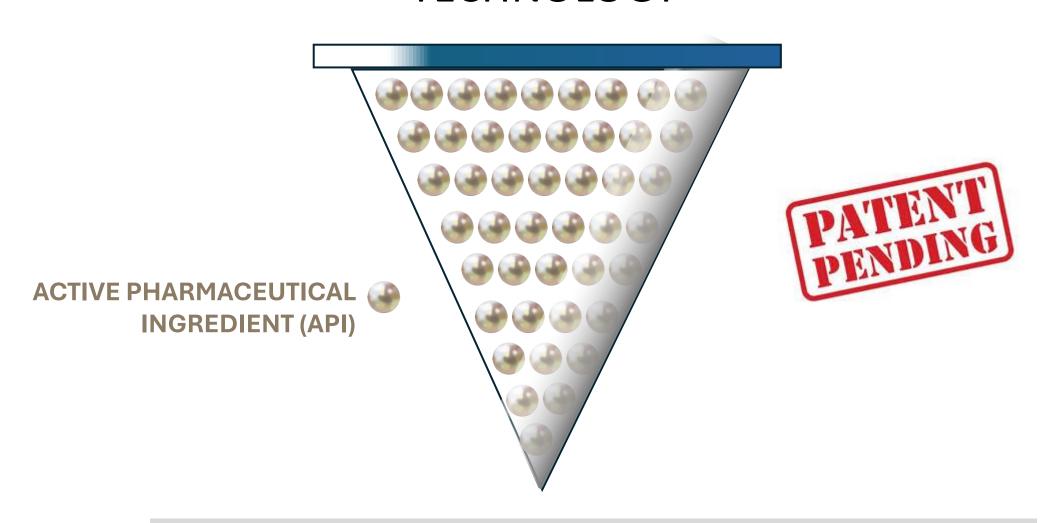
DERMAPTM – UNLOCKING THE POENTIAL OF THE MICRONEEDLE PATCH

COMPETITOR DISSOLVING MICRONEEDLE PATCH TECHNOLOGY



- Up to 80% of microneedle made from polymer to ensure structural integrity
- As little as 20% room left for API
 - X Reduces drug loading potential
 - X Reduces efficacy and dosing flexibility
 - X Potentially increases safety risk

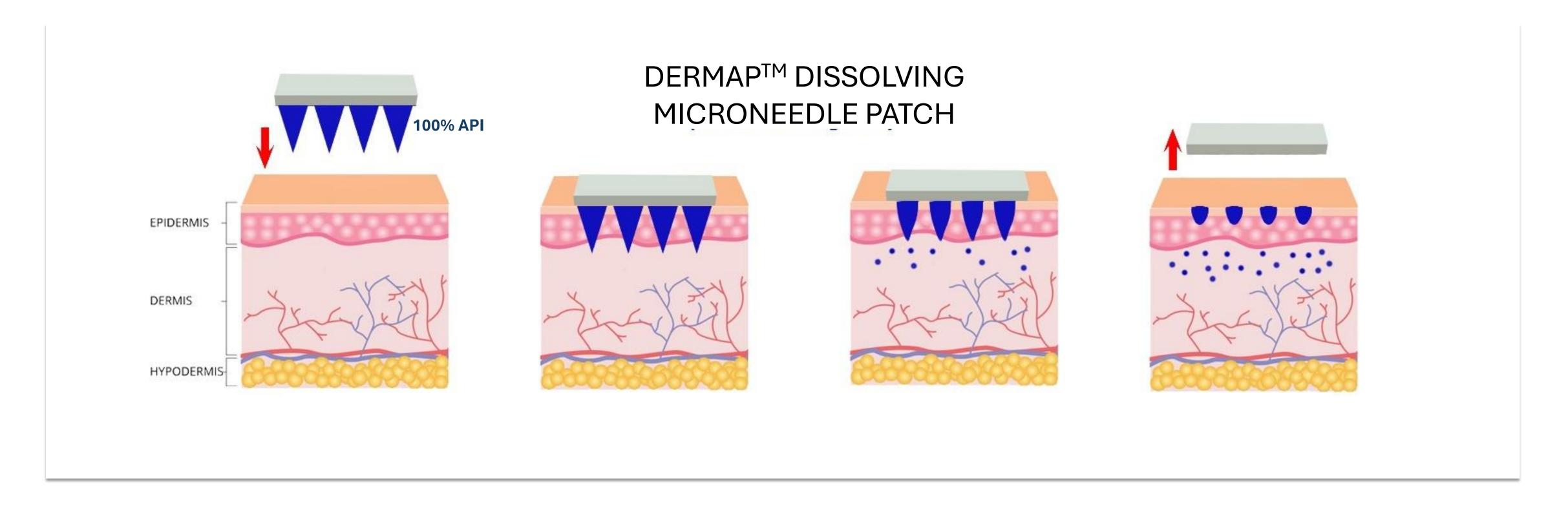
DERMAPTM DISSOLVING MICRONEEDLE PATCH TECHNOLOGY



- Uniquely, up to 100% of microneedle made from API; no polymer
- Basis of ArrayPatch patent¹
 - Maximum drug loading potential
- Increases efficacy and dosing flexibility
- Potentially reduces safety risk



DERMAPTM – UNLOCKING THE POENTIAL OF THE MICRONEEDLE PATCH



- DerMapTM patch is pressed onto the skin painless as only penetrates the outer skin (stratum corneum)
- Applied overnight, removed in morning
- Microneedles dissolve, allowing delivery of drug payload
- Platform: pipeline of small molecules identified with specific properties suitable for formulation into microneedles



Array DERMAP DELIVERY PLATFORM — Multiple Products

Product	Research	Preclinical	Phase I	Phase II/III	Market
ITZ-DerMap Nail fungus					\$6BN
VES-Deriviap Basal Cell Carc					\$10BN
BET-DerMap Psoriasis					\$34BN
ZOL-DerMap Migraine					\$6BN
Others 14 molecules					

- Reduced development risk all molecules being delivered are already approved
- Strong IP DerMapTM Technology patent filed US/EU/S Korea, expiry 2041; additional IP planned
- Additional pipeline opportunities approaches from 3rd parties for research collaborations

1 in 4 people can be affected with <u>nail</u> fungal infection at any given time





Array THE PROBLEM

Onychomycosis (nail fungal infections)









Current treatments (oral/topical) up to 1 year – only 1 out of 3 show cure response



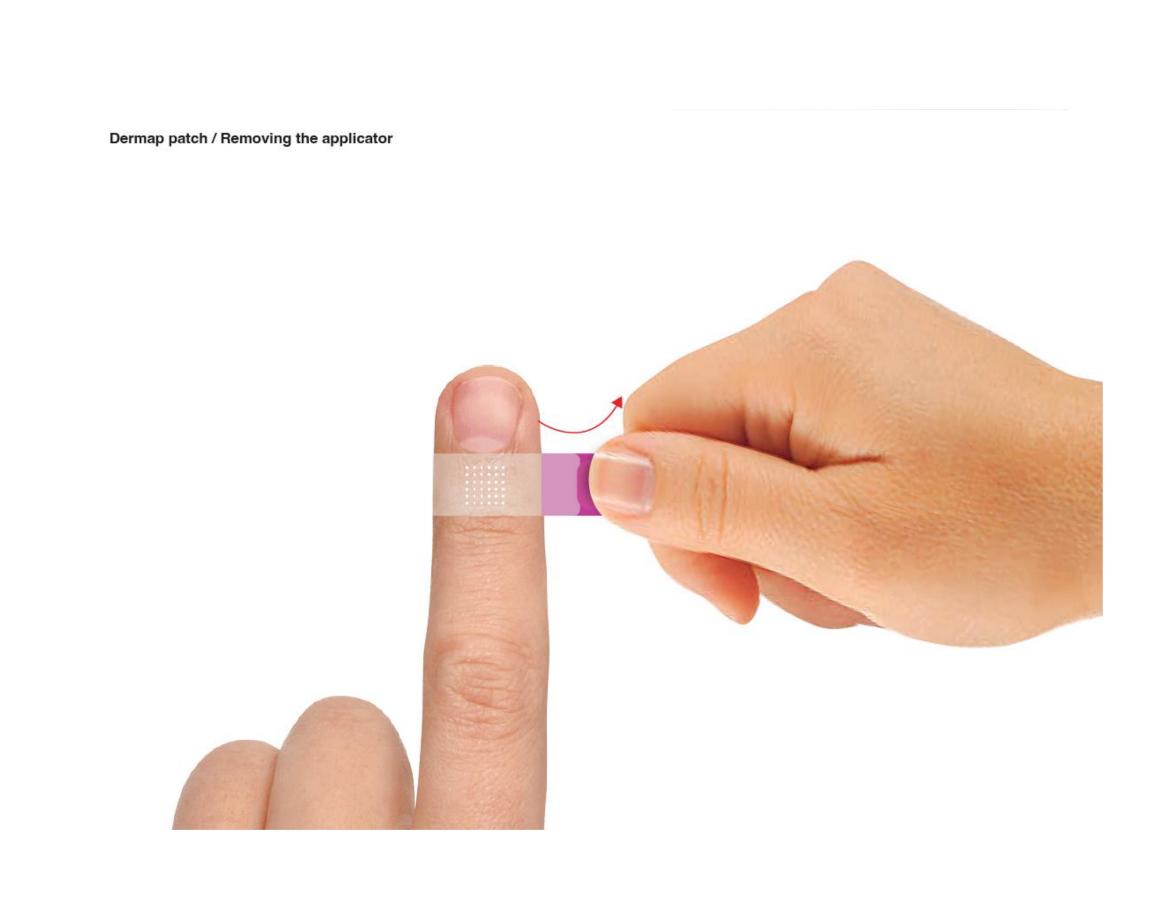
Severe liver toxicity with current oral treatments



Array ITZ-DerMapTM

Delivery of previously approved nail fungal drug itraconazole via DerMap microneedle patch







Array ITZ-DerMapTM

Delivery of previously approved nail fungal drug itraconazole via DerMap microneedle patch

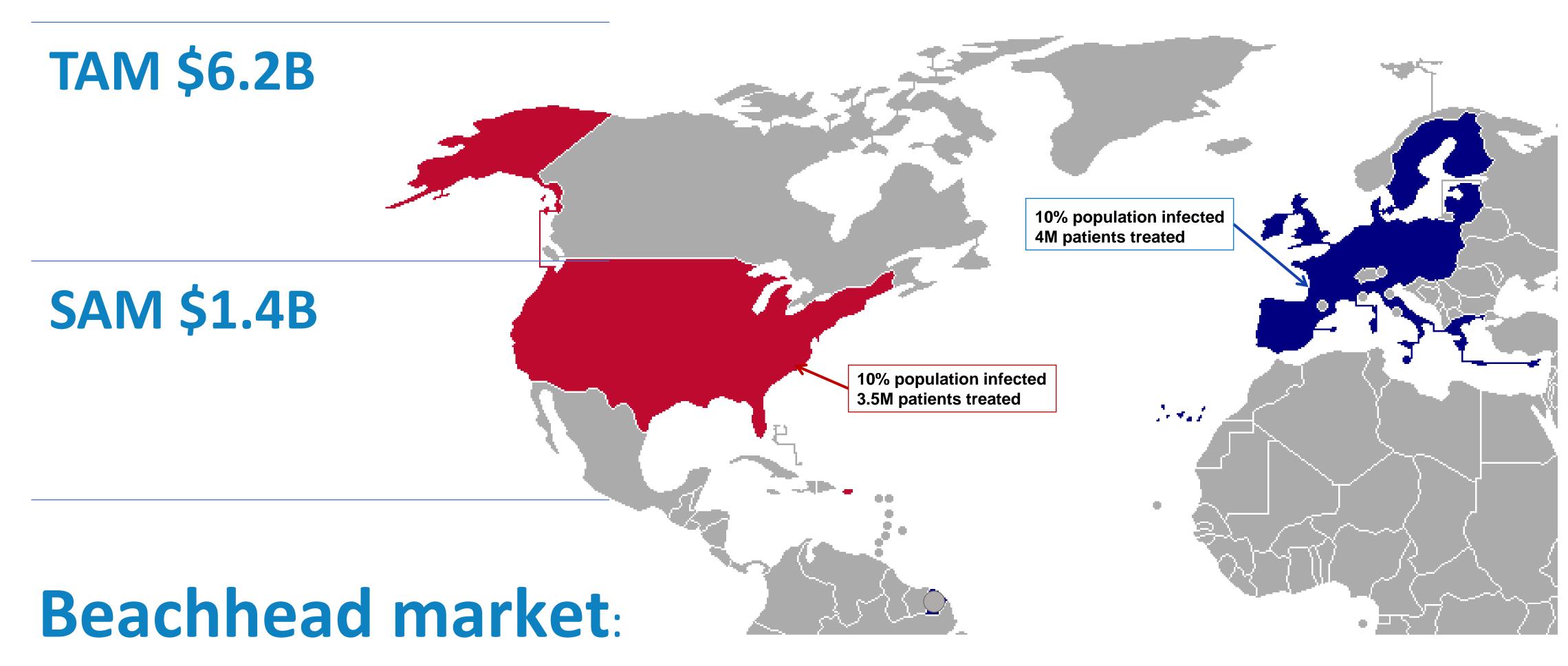




https://www.youtube.com/watch?v=F6jPA9liUMc



Array NAIL FUNGUS - MARKET OPPORTUNITY



Diabetics > 150m suffer from nail infection



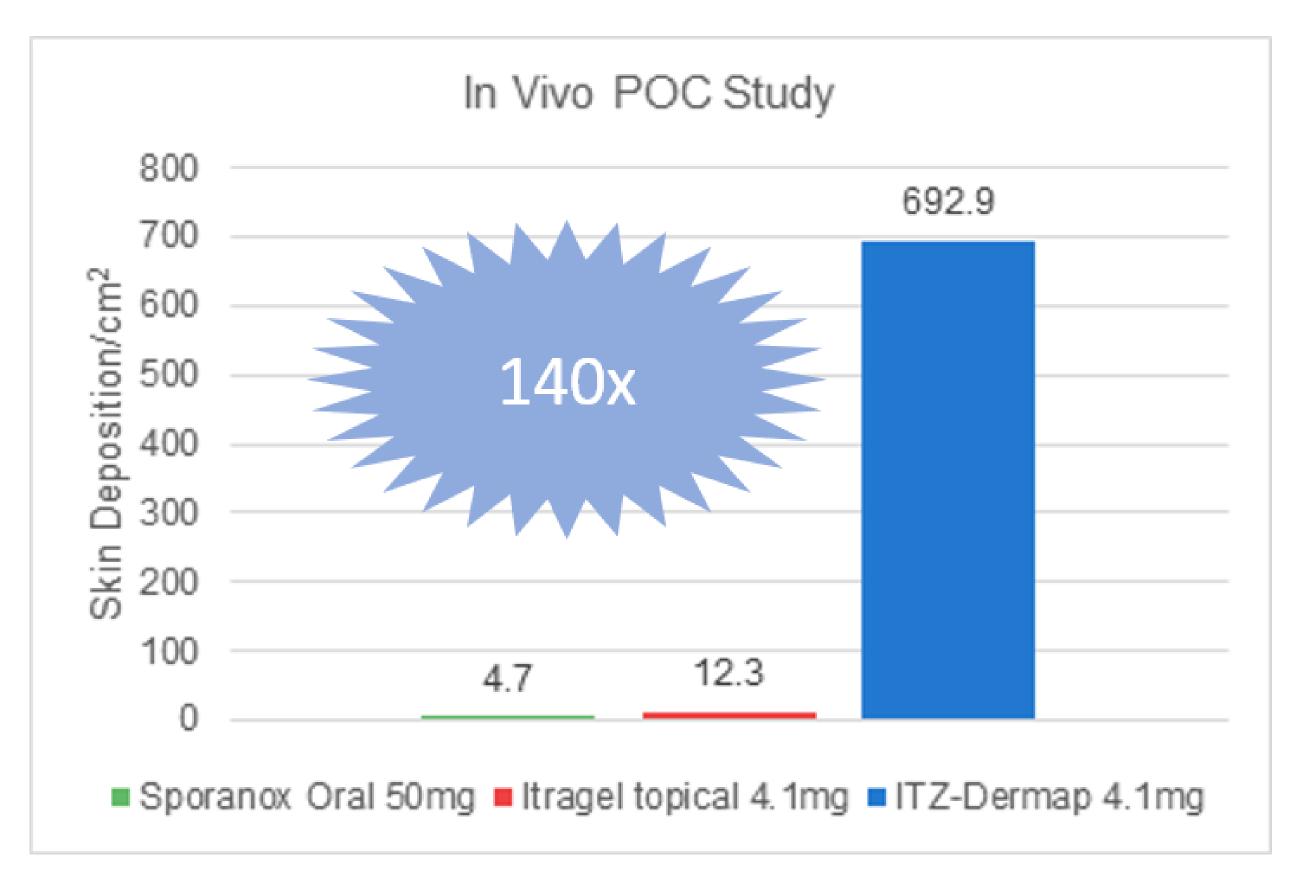
Array ITZ-DerMapTM DIFFERENTIATION Vs COMPETITORS

		Dose	Skin deposition	Liver toxicity	Dose Frequency
DerMap 12 Patches	The Patch Creshable In Danker International Control of the Market	Low 1/10	High 140x	Low	Once/week
Lamisil® 1% w/w cream (GSK)	Terbinafine hydrochloride CAMISIL® To Gream Broad-spectrum antifungal Cream	High	Low	Low	Twice/day Output
Lamisil® 250mg (GSK) 14 tablet	NOVARTIS Lamisil® Tablets Lamisine Terbinafine 250 mg 28 tablets on a Calendar pack	High	Low	Yes	Twice/day Output
SPORANOX® 100mg (Janssen-Cilag Ltd) 28 capsules	Sporanox® Itraconazole 100 mg per capsule 15 oral capsules Janssen	High	Low	Yes	Twice/day

Array ITZ-DerMapTM DEVELOPMENT STATUS

KEY DEVELOPMENT HIGHLIGHTS

- ✓ 6 month stability studies
- ✓ Penetrative ability studies
- Bioavailability studies
- ✓ In vivo proof of concept studies (versus current therapy)
- ✓ Manufacturing scale up designed
- Prototypes
- ✓ HPRA/EMA/BfArM engagement advice on clinical plan received



- 140 times greater skin deposition of active medicine (itraconazole) versus oral delivery of the same medicine
- Provides basis for more effective and safer treatment



Array atch ITZ-DerMapTM DEVELOPMENT PLAN TO APPROVAL

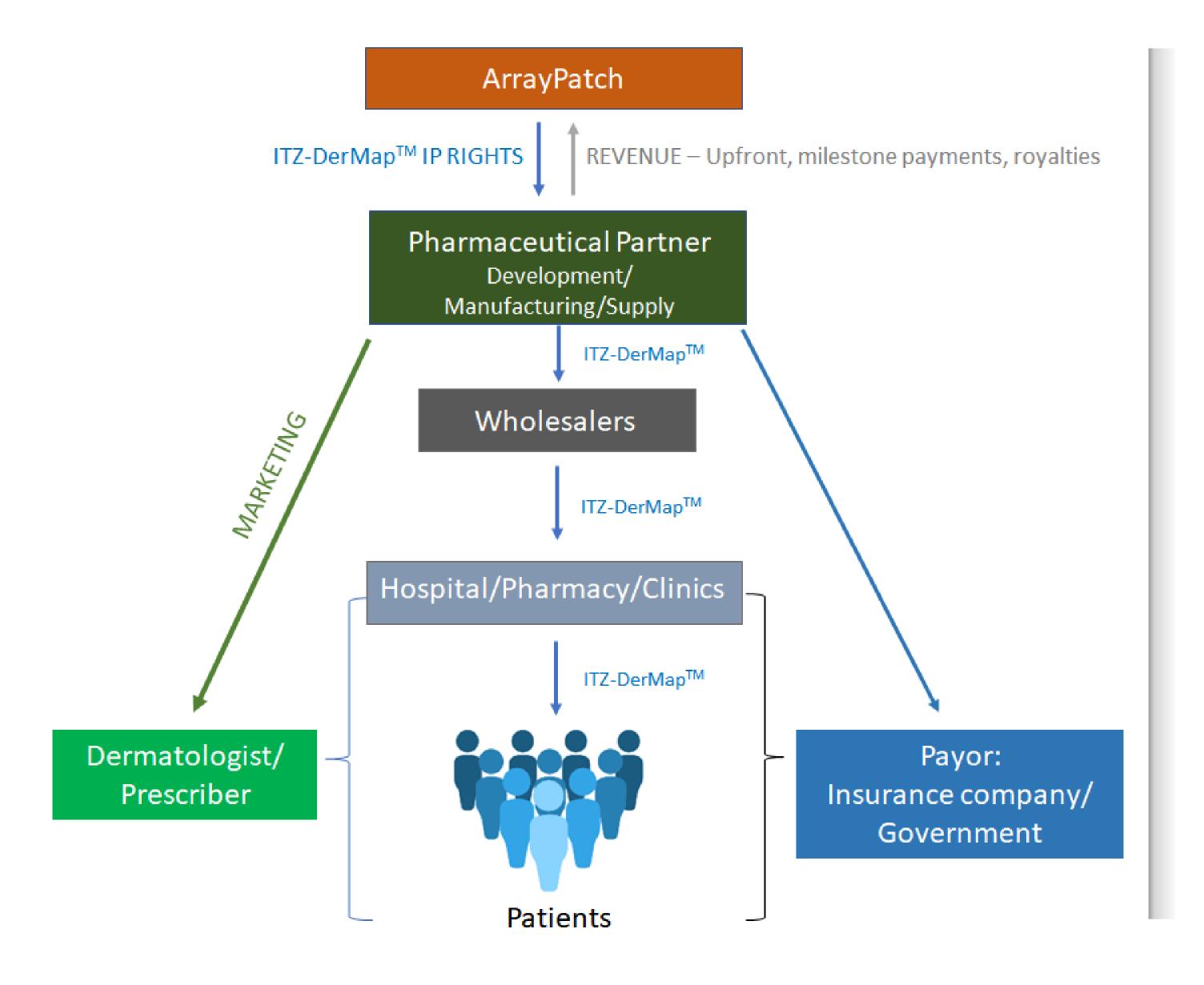
	TIMEFRAME FROM SEED FUNDING:																			
	Year 1			Year 2				Year 3				Year 4			Year 5					
	Q1	Q2	Q3	Q4	Q1	Q2	Q3 (Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Pre-clinical/CMC		•					•			•										
Animal tolerability study																				
Phase I synthesis scale up clinical supply																				
Stability																				
Phase 2/3 clinical supply																				
Clinical																				
Tolerance study (no API involved)							POT	ΓENTΙ	AL PA	ARTNE	RING									
Phase I (12 healthy vlts, open label)																				
Phase 2/3 Pivotal (150 pts, open label)																				

APPROVAL

- Development for US and Europe
- Leverage shortened 505(b)(2) US development path given itraconazole already approved
- CRO input and proposals received
- Plan subject for further regulatory guidance



Array ITZ-DerMapTM GO TO MARKET STRATEGY



- License to larger commercial partner post initial clinical trials
- Partner responsible for final development studies, launch and sales
- Reduces financial, clinical and execution risk
- Provides validation for technology.
- Partnering revenues to fund pipeline development
- Positive partnering discussions already underway



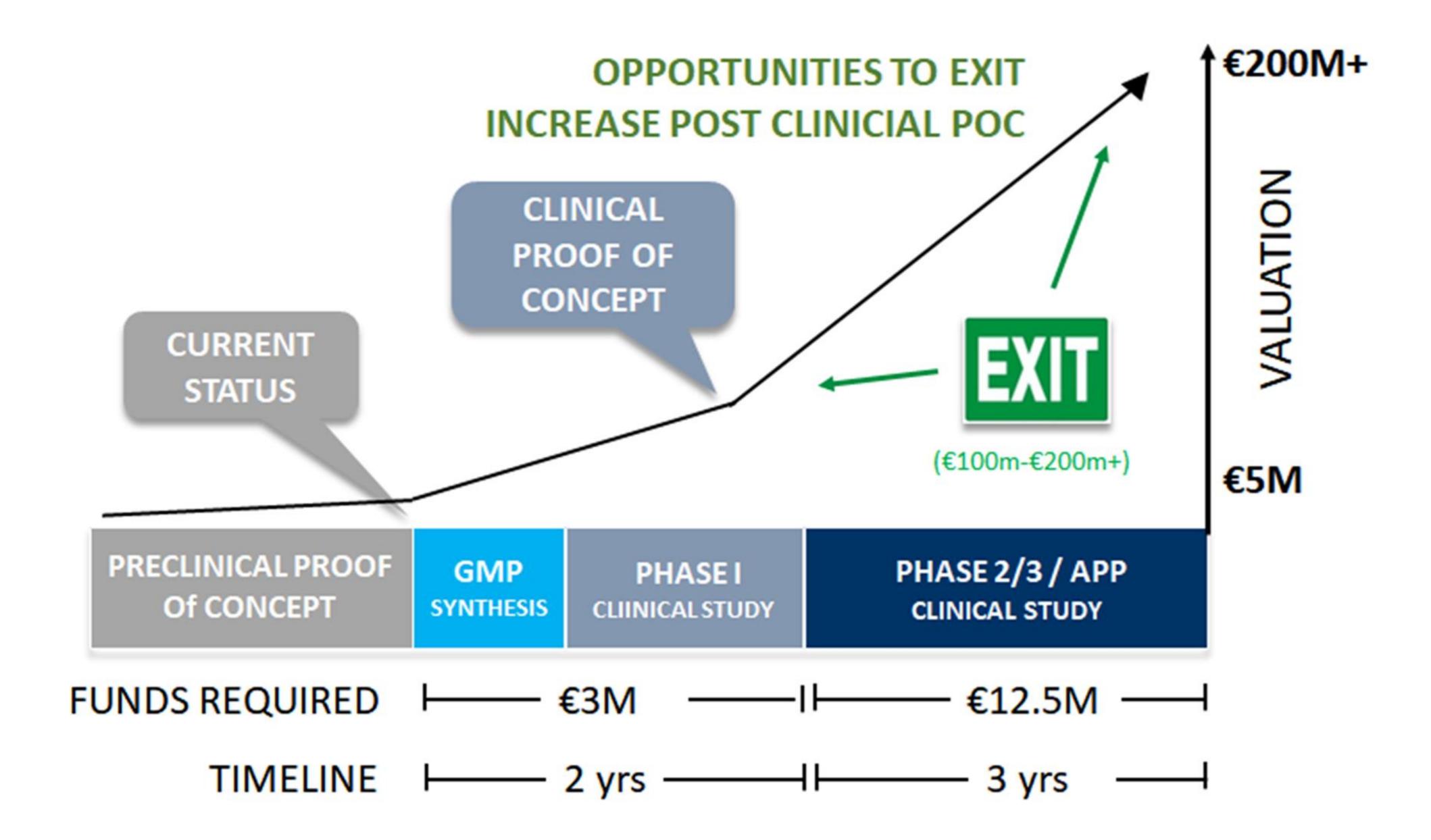
Array ITZ-DerMapTM PEAK REVENUE POTENTIAL (US)

	Topical	Oral	OTC	Total
TRx/pts (M)	2.1	2.4	1.0	5.5
Peak Mkt Share	10%	5%	1%	
AWP Price \$	1,500	1,500	1,500	
Deductions	50%	50%	50%	
Peak Net Revenues \$M	159	89	7	255

Sources: Cleveland Clinic; Wilsons analyst reports on Hexima; Moberg Pharma; TRx - total prescriptions; pts = patients.

- US represents main target market peak US revenues \$200-300M
- Upside penetration of ~30M currently untreated patients in the US; strong clinical efficacy/safety
- Lower EU revenue potential given prevalence of OTC treatments; lower pricing available

Array ITZ-DerMapTM PRODUCT/FUNDING ROADMAP



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EXIT – BENCHMARK TRANSACTIONS

ALTARIS

2020: Acquired microneedle drug delivery business Kindeva

\$650M



2019: Licensed European rights to topical nail fungus treatment from Moberg

€50M plus royalties

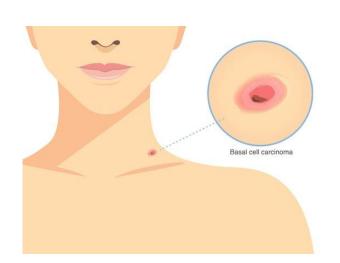


2017: Licensed European rights to novel anti-fungal therapeutic from Basilea

\$500M plus royalties

- Likely partners for ITZ-DerMapTM identified: Bausch, Karo, Almirall, Sandoz, Sanofi, Galderma
- Likely CDMO technology acquirors: LTS Lohman, Kindeva, Hovione, Becton Dickinson, Merck, J&J

Array Patch PIPELINE OPPORTUNITIES - PRECLINICAL



VES-DERMAPTM

Basal Cell Carcinoma

- Certain areas of body not suitable for surgery – e.g. Face, elbow, knee
- Oral medication difficult to tolerate
- Ves-DerMapTM offers potential solution applying patch adjacent to BCC



BET-DERMAPTM

Psoriasis

- Thicker skin around elbows and needs makes targeting plaque more difficult
- BET-DerMapTM is designed to target medicine into the dermis for more effective outcomes



ZOL-DERMAPTM

Migraine

- Current treatment can take >2hrs to work
- Patients need faster treatment
- ZOL-DerMapTM offers potential solution applying patch to forehead

\$10BN

\$34BN

\$6BN

SEED USE OF PROCEEDS

ITZ-DerMap [™] development costs	€1.7M
Personnel costs	€1.2M
Other (net of tax credits)	€0.1M
Total	€3.0M

SEED MILESTONES TO BE ACHIEVED

- ITZ-DerMapTM GMP manufacturing scale up (for clinical)
- ITZ-DerMapTM Phase I safety data with biomarker readouts supporting Phase 2/3 design
- On-going strategic partner engagement.
- ~2 year runway

CURRENT CAP TABLE*

Founders	10,000	75%
UCC	2,000	15%
Option Pool	1,333	10%
Total	100%	100%

^{*€235}k 5% Convertible loan notes issued to date - will convert at a **20**% discount to SEED round.



Lead investor, Lakeside Capital

> €1.8M commitments











Array EXECUTIVE TEAM



CEO/Founder Waleed Faisal

- Led several El projects
- Raise > 1.3m funding
- Lead inventor DerMap
- QA Xeolas Pharmaceutics





CFO Darren Cunningham

- >25 yrs industry experience
- Co-founder/ former CEO Inflection Biosciences & Mysthera Therapeutics
- Fundraising > \$100M
- Global licensing >\$200M





CSO Tord Laboda

- >15 yrs industry experience
- Extensive knowledge of the full value chain
- Ex VP and Head of Global Clinical Dev. LEO Pharma





COO **Ryan Bamsey**

- >25 years developing and launch of microneedle products
- Design and implement QMS aligned with ISO9001
- >4 years product development















Array ADVISORY BOARD



Scientific Advisor Prof Abina Crean

- Prof Pharmaceutics, UCC
- >6 years industrial R&D
- >13 years microneedle R&D
- Funded PI in SSPC, SFI





Clinical Advisor Mary McGonagle

- **Board Certified** Dermatologist and Dermatopathologist
- >4 years Biotechnology Consultant
- Adjunct Prof





Clinical Advisory Alan Irvine

- Dermatologist
- Prof Dermatology Trinity College
- > 300 Publications
- Chief Investigator Trials





Regulatory Advisor Stephen Liggett

- >10 years Regulatory/Product **Development Consultant**
- Director of Regulatory Affairs at Icon Ltd
- PhD Pharmaceutics

















INVEST?

Unique microneedle technology platform to improve safety and efficacy

Delivering approved drugs - short and low risk development pathway

Platform solution, with multiple exit opportunities

Market size of 1st application > \$6B

Experienced team

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