Qures Group Ltd

Richard G. Stead CEO Qures Group Ltd Nov 2021

How Tears can take us from ABR to \$100m

Tears protect eyes from pathogen ingress



Our Product

We have a unique technology, and means of delivery, bringing efficacies lacking in existing products Product Hypothiocyanite Created at point of use

Development

Enhanced stability pre and post creation

Proof Successful small study in URTI

The Problem

Deaths from AMR infections are rising

Forecast of >10mn worldwide by 2050*



What payers want from a treatment for infections

Governments and Insurers

- Effective
- Fast acting avoiding time in hospital
- Simple to administer non-medical carers and at home
- Resistance avoided

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WHAT CARERS WANT FROM A TREATMENT FOR INFECTIONS

Doctors, GPs, Nurses, Health workers

- Effective
- Fast acting
- Broad spectrum effectiveness
- •Simple to administer (<u>At home</u>)
- <u>Resistance</u> avoided
- •Bio/Nature-identical
- Not harmful to friendly bacteria
- No adverse effects
- Cost-effective
- •Complements immune system

Target Market Value

ABR Prescription is \$15bn*

AB Prescription >\$57bn Of which sales to distributors for ABR \$5bn*. Our target market.

*Source: Grand View Research

If we can achieve 1% = \$0.05bn t/o with gp >\$0.025bn

Growth

1. First to market for Indication A (gram-ve)

2. Secure FDA approval, thereby securing market protection

Business Models

1. Developer to Distribution to users

2. Licences to others – create our "competitors"





Do we have what they want from a treatment for infections??



... and more.

QURES

ANTI-MICROBIALS

Can we meet the demands for ABR treatment?

Effective	yes/no	yes
Fast acting	no	yes
 In-vivo efficacy 	yes	yes
 Simple to administer 	yes/no	yes
<u>Resistance avoided</u>	no	yes
 Bio-identical 	no	yes
 Not harmful to commensal bacteria 	no	yes
 Absence of adverse effects 	no/yes	yes
Cost-effective	yes/no	yes/no
 Complements immune system 	no	yes

Advantages/Benefits of Qures ABR treatment

- Fast acting
- Simple to administer
- Resistance avoided

Reduced time in hospital Reduction in lost working time Can be handled by semi-skilled Home treatment possible Huge savings

- Not harmful to commensal bacteria Avoid later problems
- Absence of adverse effects

Patient satisfaction

Financial Programme

Years	Activity
1 -2	Investment \$8
2-3-4	Clinical Trials
4-5	Sales T/O \$100m
	GP \$50m

Team

Richard Stead	CEO	Founder
Dr Kevin Pritchard	Executive	Science
Dr Paul R Clayton	Non Exec	Medical Research
Dr Diana Garnham	Non Exec	Chair
Prof Michael Ashby	Non Exec	
Prof Neil Williams	Non Exec	
Lucy Futter	Exec Gov	vernance and Marketing

Risks

- Slow to be granted clinical trial go-ahead.
- FDA etc do not grant marketing approvals.
- In-direct Competitors appear.
- Big pharma infringe our patents.

Next Steps

2021

Secure funding

2022 - 2023

- Spending on research and personnel
- Regulatory programme begins

2023 - 2024

• Clinical trials

2024 /25

- Marketing approvals
- Sales start

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THANK YOU

Richard G. Stead CEO