Innovations powered by XBIO<sup>®</sup> technology

# Next Science Limited Investor Presentation

SCIENCE

NEXT

ASX: NXS

May 2019

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# **NEXT SCIENCE**<sup>®</sup>

An established R&D company commercialising anti-biofilm solutions

Biofilms represent 90% of all bacteria and pose a threat to humans, animals and the environment

XBIO is the only nontoxic solution to deconstruct Biofilm and destroy the bacteria Founded 2012 Listed on ASX April 2019 Market cap ~\$498m

80,000 Patients treated with XBIO technology Multiple FDA clearances with multiple international regulatory approvals pending

Strong Revenue Growth Established global distribution agreements

Extensive pipeline across Medical Devices, OTC Drugs and Pharmaceuticals

**17 Patents** 49 Patent applications

# **Corporate Overview**

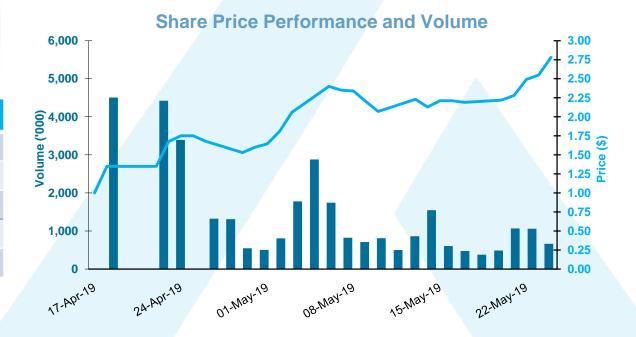
Stock Overview	
ASX code	NXS
Share Price (24 May 2019)	\$2.78
Market capitalisation @ \$2.78	\$498M
Total Shares on Issue	179.2m
Listed Shares (tradable)	65.7m
Escrowed Shares	113.4m
Options	10.7m
Shareholders (1,555 at listing)	3,265
Trading range (since listing)	\$1.24 to \$2.78
Average daily volume (since listing)	1.4m Shares

Substantial Shareholders	
Auckland Trust Company Ltd*	25.96%
Walker Group Holdings Pty Ltd*	16.46%
Matthew Myntti (Founder & CTO)	11.53%
Judith Mitchell (Managing Director)	2.64%
Total Board & Management Shareholdings	15.4%

\* Entities related to Lang Walker

#### **Escrowed Securities**

Shares escrowed until 26 June 2019	0.71m
Shares escrowed until 25 September 2019	0.07m
Shares escrowed until 18 April 2020	39.17m
Partly Paid Shares escrowed until 18 April 2020	0.65m
Shares escrowed until 18 April 2021	72.85m
Total Shares Escrowed	113.4m
Options escrowed until 18 April 2021	5.85m



# Biofilm is a global healthcare problem

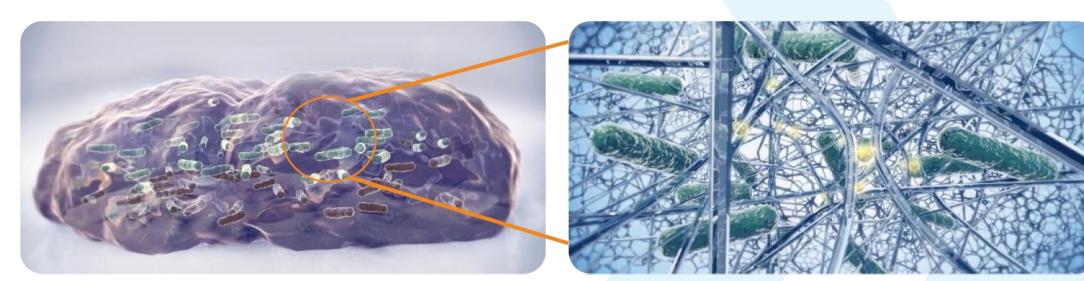


••• **90%** OF BACTERIA EXISTS IN BIOFILMS STRUCTURE Bacteria in BIOFILMS can become up to 1000-fold more resistant to antibiotics and biocides when compared to planktonic counterparts.

..........

# Biofilm is a three-dimensional problem

- **1) PLANKTONIC PATHOGENS**
- 2) **BIOFILM-ENCASED PATHOGENS**
- **3) ENCASING STRUCTURE**



Polymicrobial community of organisms

Slimy tangles of protective polymer fibers linked with metallic bonds

# The problem with Biofilm and encased pathogens

# BIOFILM

## **BLOCKING EFFECTIVE TREATMENT**

- Large molecules such as large-molecule antimicrobials, antibodies, and inflammatory cells
- » Biofilm matrixes act as diffusion barrier to small molecules like antibiotics

## **HYBERNATION** (quiescent bacteria)

 Biofilm matrixes have developed a mechanism for a subpopulation to become metabolically quiescent (i.e., to hibernate)

# **ENCASED PATHOGENS**

## **MUTUAL PROTECTION**

- » Exhibit cooperative protective effects through RNA and DNA transfer
- Some species can assist other species to attach and incorporate into the biofilm (quorum sensing)

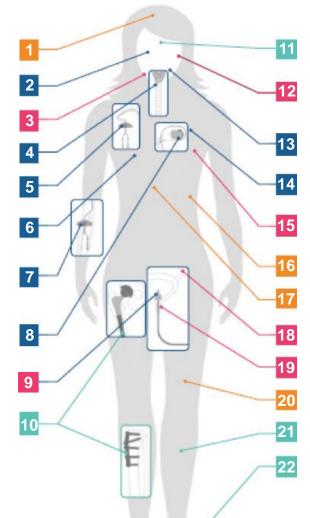
## SEEDING

 Biofilm colonies can detach from the main colony and recolonize individually or in clumps

# Biofilms pose a far-reaching threat to humans, animals and the environment

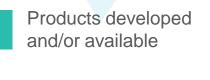
#### **DEVICE-RELATED INFECTIONS**

- 1) Ventricular derivations
- 2) Contact lens
- 3) Mouthwash
- 4) Endotracheal tubes
- 5) Vascular central catheters
- 6) Tissue fillers, breast implants
- 7) Peripheral vascular catheters
- Prosthetic cardiac valves, pacemakers and vascular grafts
- 9) Urinary catheters
- 10) Orthopedic implants and prosthetic joints



#### TISSUE INFECTIONS

- 11) Acne
- 12) Chronic otitis media, chronic sinusitis
- 13) Chronic tonsillitis, dental plaque, chronic laryngitis
- 14) Endocarditis
- 15) Lung infection in cystic fibrosis
- 16) Kidney stones
- 17) Biliary tract infection
- 18) Urinary tract infection
- 19) Vaginosis
- 20) Osteomyelitis
- 21) Surgical site infections
- 22) Chronic wounds



In development

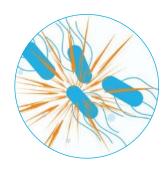
Areas for research

No research at this time

# Xbio<sup>TM</sup> - The Solution to Biofilm

Xbio<sup>™</sup> formulations leverage a patented composition-of-matter and method of action which encompass the non-toxic technology designed to physically break down the biofilm's protective structures. This exposes and eradicates bacteria that are enveloped within the technology and provides targeted therapy with no known antimicrobial resistance.

# The Solution – Xbio<sup>™</sup> TECHNOLOGY



## DECONSTRUCT THE BACTERIAL BIOFILM BARRIER

Next Science's Xbio breaks the ionic bonds that hold the biofilm together. The polymers are then pulled into solution, effectively dissolving the biofilm barrier.

## DESTROY BACTERIA WITHIN THROUGH CELL LYSIS<sup>1</sup>

With the barrier dissolved, bacteria are exposed and more vulnerable to attack. Bacteria enveloped by Xbio technology experience cell lysis and are destroyed. Cell lysis is non-discriminatory destroying gram-positive and gram-negative bacteria, persister cells, and spores. There is no known resistance mechanism to cell lysis.

## **DEFEND** FROM RECOLONISATION

The periodic release of bacteria from biofilms has been linked to chronic relapsing Infections.<sup>2</sup> Disrupting and destroying the biofilm barrier can reduce the rate of biofilm recurrence by up to 1,000 times, effectively defending against recolonization.<sup>3</sup> Unlike other agents that claim to destroy biofilms, there is no known evidence of bacterial resistance to the Xbio technology.

Xbio<sup>™</sup> is the only non-toxic solution to deconstruct the bacteria's protective barrier. We've applied material science innovation to physically deconstruct the bacteria's protective structures, exposing and then eradicating bacteria through cell lysis<sup>1</sup>, rather than using toxic or resistance building ingredients.

 Lysis: disintegration by rupturing the cell membrane.
 Costerton JW et al.
 Potera C :antibiotic resistanc: biofilm dispersing agent rejuvenates older antibiotics" Environmental Health Perspectives 118 (7) 228.

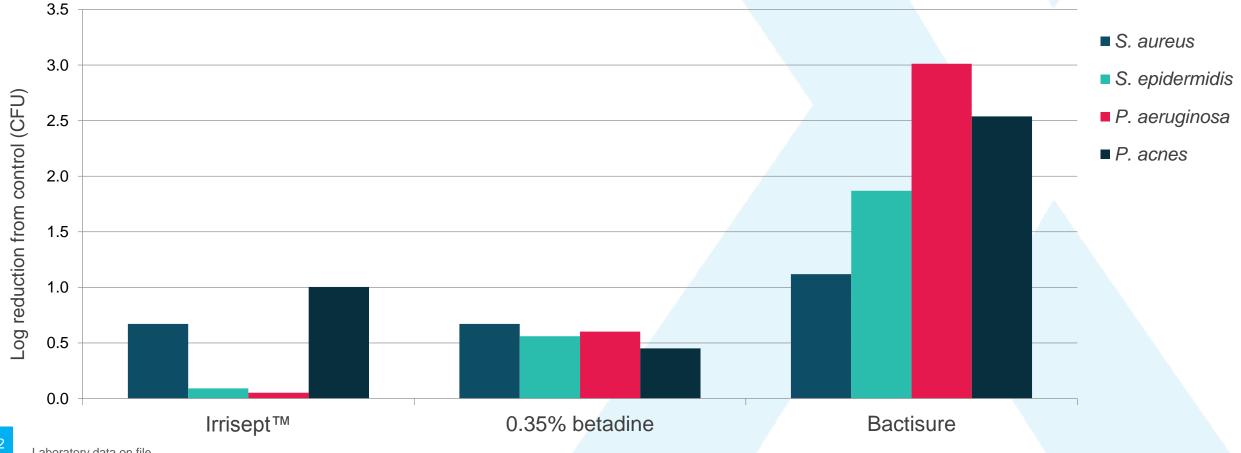
# Xbio<sup>™</sup> Family – Current Products

4 FDA CLEARED PRODUCTS CURRENTLY IN THE US MARKET, WITH SALES OF ACNE GEL TO COMMENCE IN AUSTRALIA IN 2019.



# Xbio<sup>™</sup> Bactisure Surgical Lavage Biofilm efficacy versus current technologies

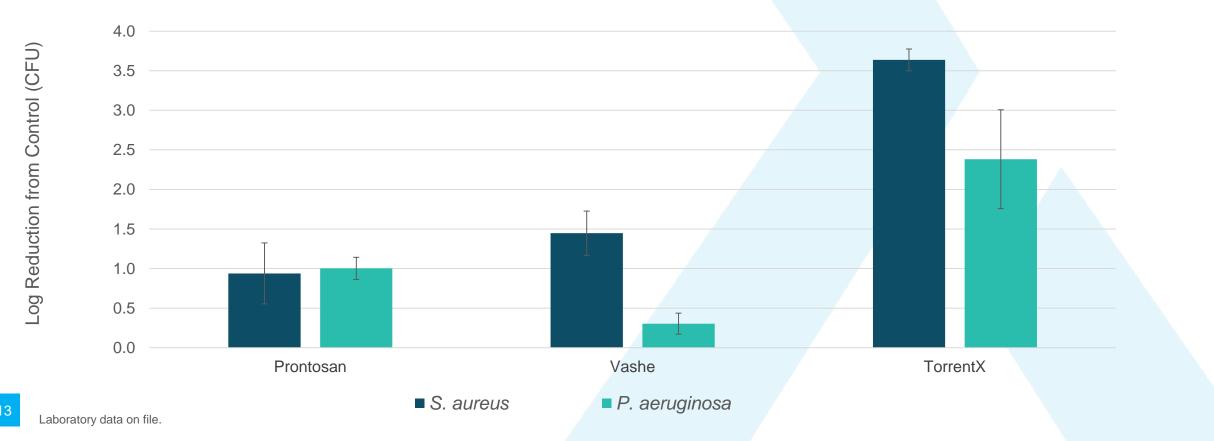
Montana State University (MSU) BIOFILM EFFICACIES OF SURGICAL LAVAGES-1 MINUTE STATIC 72 HOUR BIOFILM, DRIP FLOW REACTOR MODEL



# Xbio<sup>™</sup> Wound Wash (TorrentX) Biofilm efficacy MSU drip flow reactor model (3 day Biofilm)

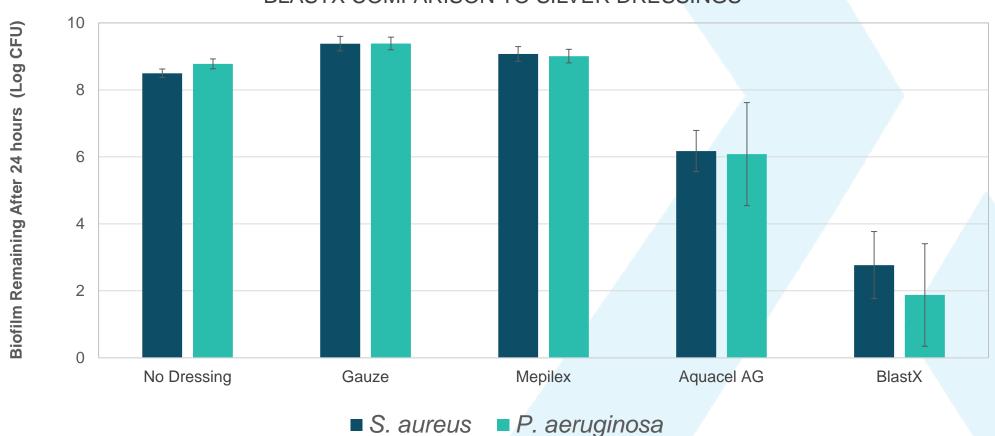
MSU 20-SECOND FLOW + 1 MINUTE SOAK BIOFILM TESTING RESULTS SHOW TORRENTX EXHIBITS UP TO *8 TIMES MORE* BACTERIAL BIOFILM LOG REDUCTION COMPARED TO COMPETITORS

MSU BIOFILM TESTING - 20 SECOND SPRAY + 1 MINUTE SOAK



# Xbio<sup>™</sup> Gel Products Biofilm efficacy versus current technologies

MONTANA STATE UNIVERSITY - CENTER FOR BIOFILM ENGINEERING RESULTS 72 HOUR BIOFILM, DRIP FLOW REACTOR, 24 HOUR TREATMENT, ~8 LOG CONTROL



BLASTX COMPARISON TO SILVER DRESSINGS

# **BLASTX Chronic Wound Care – Case Studies**

### 67 YEAR OLD DIABETIC FEMALE WITH A HISTORY OF TRAUMA TO THE LOWER LEG



84 Y/O MALE, TYPE II DIABETES, KIDNEY FAILURE, HYPERTENSION, MULTIPLE CANCERS – RIGHT FOREARM LOOP GRAFT BECAME INFECTED



4.7 cm wound, unresponsive to standard care, surgery and antibiotics for >8 months



Patient's wife applied BlastX with dressing changes every 2 days. Bridged to 2 smaller wounds after 1 week



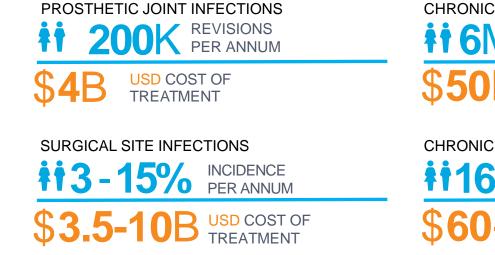
Continued therapy and returned 2 weeks later, wound healed

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# Large Market Opportunity

## COST AND TREATMENT IN THE US

Next Science products touch a breadth of applications across many medical specialties as they address the issues of chronic infection.



CHRONIC WOUNDS PATIENTS PER ANNUM

50B USD COST OF

CHRONIC SINUSITIS **16-32** WASHOUT PATIENTS PER ANNUM **560-65** USD COST OF TREATMENT FUNGAL NAIL TREATMENTS PATIENTS PER ANNUM

\$2.6B USD COST OF TREATMENT

CHRONIC OTITIS MEDIA (MIDDLE EAR INFECTIONS)

**3** US SURGERIES PER ANNUM

\$24B USD COST OF TREATMENT

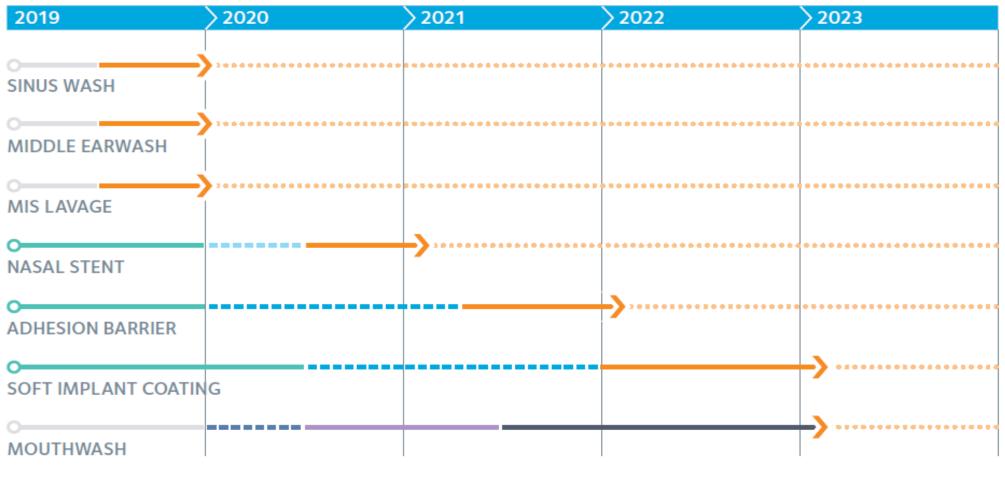
# Products Currently in Market

PRODUCT	INDICATIONS	MARKET SIZE	COMMERCIALISATION
Bactisure Surgical Lavage	Removes biofilm and bacteria from any open surgery Shown to be effective in treating Prosthetic Joint Infection (PJI) Other surgical fields to be explored in 2019	Overall market ~100M procedures a year ~Target 1M procedures p.a. 2019 Minimum 46,000 procedures	<ul> <li>Zimmer Biomet – Global distribution agreement</li> <li>Market Leader in Joint Replacements (US\$7.8B Annual Revenue), with a global sales force</li> <li>Contract includes take or pay minimum purchases for 20 years</li> <li>April 2017, started marketing in Revision surgeries</li> <li>June 2018 - moved to Primary Joint Surgeries using the US Reconstruction sales network</li> <li>Expanding to all Open surgeries in US 2019, and to Europe Q4, 2019</li> </ul>
BlastX Antimicrobial Wound Gel	Any topical wound, or 1st or 2nd degree burn Effective in the treatment of chronic wounds (Diabetic Foot Ulcers, Venous Leg Ulcers) and topical wounds	Global patient population 10M per annum US Expenditure \$50B	Global distribution with <b>3M</b> commenced in 2019 In 2018 BlastX was sold by a Next Science direct sales force Contract includes take or pay minimum purchases for 3 + 3 years
ACNE Cream (OTC Product)	Acne – Problem Skin OTC / Cosmeceutical Market	Globally \$5B market	Exclusive distribution through <b>Advanced Skin Technology's</b> 3,000+ clinic network, and non-exclusive online sales in Australia/NZ to start in H2 2019 Combination of other OEMs for distribution and royalty models under discussion
TorrentX Wound Wash (OTC Product)	Wound wash for all wounds/ non sterile environment	~10M Chronic Wounds ~Accident and Emergency	Distribution agreements in discussion Product released in USA in December 2018 via internal sales team
SurgX – Antimicrobial Sterile Wound Gel (Medical Device)	Added as a topical treatment when closing a surgical site to prevent surgical site infection (SSI)	~200M surgeries a year	Distribution agreements in discussion Product released in USA in October 2018 via internal sales team and commission based distributors

# 2019 Product Submissions Medical Devices

PRODUCT	INDICATIONS	MARKET SIZE	COMMERCIALISATION
Sinus Wash	Chronic Sinusitis treated with Functional Endoscopic Sinus Surgery (FESS) or Washouts	~1M FESS Surgeries a year	Distribution agreement with Grace Medical (slide 27)
			Being submitted for approvals: FDA CE, Canada & Australia
MIS Lavage	Antimicrobial wash out for	~100M Procedures/ Surgeries a year	Distribution agreements in discussion
(Minimally Invasive Surgery)	minimally invasive surgical procedures, endoscopies, colonoscopies, arthroscopies		Being submitted for approvals: FDA & CE
Middle Ear Wash	A wash for the Middle Ear during a tympanoplasty procedure. For patients with chronic ear infections, that have chosen to have grommets and tubes inserted to try to resolve the infection	Chronic active or suppurative otitis media affects 65 to 330 million people worldwide	Distribution agreements in discussion
			Being submitted for approvals: FDA & CE

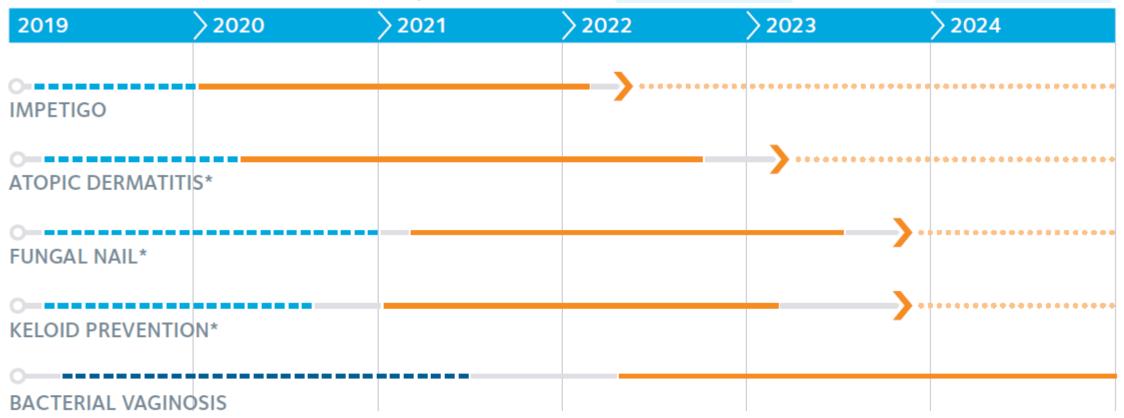
# **Development Plan – Medical Device Products**



Development
 Safety Trials and validation
 Pre market approval trials
 Clinical trial
 Contract Negotiations
 Manufacturing validation and verification
 Regulatory review
 Market Launch

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# Pharmaceutical Developments – Pharmaceutical Products



Concurrent Trials 🔲 Phase II and III Clinical Trials 📕 Submission and Approval 🗦 Market Launch

\*These products will be submitted through the 505 2(b) Drug Approval Pathway.

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# **Board of Directors**



George Savvides has 30 years of experience in the Australian & NZ healthcare sector, including sitting on a number of company Boards. He was CEO of two successful IPO listings on the ASX, Sigma in 1999 and Medibank Private in 2014. He served as Medibank CEO for 14 years.



Dan Spira is the CEO of iNova Pharmaceuticals. Previously Vice President at Valeant Pharmaceuticals / Bausch + Lomb from 2011 to 2015. Prior to Valeant, Mr. Spira spent over 15 years at Johnson & Johnson progressing through a range of sales, marketing and general management roles.



Judith Mitchell joined Next Science as CEO in October 2017. Previously Ms. Mitchell was President, DePuy Synthes Asia Pacific, the Orthopaedics Division of Johnson and Johnson, holding the same role at Synthes GmbH prior to acquisition. Prior to that, Ms. Mitchell held various executive management roles at Cochlear and GE Medical Systems.

> AILEEN STOCKBURGER NON EXECUTIVE DIRECTOR

Aileen Stockburger is a highly accomplished Strategic Planning and Corporate Transaction Executive with proven expertise backed by an extensive portfolio of successful business development deals. Ms. Stockburger currently sits on various corporate boards and advises companies in areas of business development and corporate transactions. Previous she had held multiple business development and finance roles within Johnson and Johnson where she was responsible for numerous M&A, licensing agreements and divestitures across the corporation.



Bruce Hancox is currently a director of a number of listed and private Australian and New Zealand companies. He previously held a number of senior roles at Brierley Investments Ltd. in New Zealand as General Manager, Group CEO and Chairman, as well as serving on the board of a number of their subsidiaries.



Mark Compton is an experienced Non-Executive Director and CEO in healthcare and life sciences organisations including Australian Securities Exchange (ASX) listed companies. Mark is Chairman and Non-Executive Director of Sonic Healthcare Limited (ASX: SHL), a global medical diagnostics and healthcare organisation. Sonic is a Top 50 ASX listed company. Mr Compton was a non-executive director on the board of MQ Health and is Chairman of St Luke's Care and has held various Chief Executive Officer/Managing Director roles including of St Luke's Care, Immune System Therapeutics Limited, Royal Flying Doctor Service of Australia, SciGen Limited and Alpha Healthcare Ltd.

## NEXT SCIENCE

## **Experienced Management team**



### JUDITH MITCHELL MANAGING DIRECTOR

Judith joined Next Science as CEO in October 2017. She is a seasoned executive in the Healthcare Sector with an enviable track record of success. Prior to joining Next Science, Judith was President DePuy Synthes, Asia Pacific and her executive experience further includes Senior roles with Cochlear and GE Medical Systems.



#### DR. MATTHEW MYNTTI CHIEF TECHNOLOGY OFFICER

Dr. Myntti is the founder and CTO of Next Science where he leads the product development, research and innovation activities for the company. Dr. Myntti received his Master's and Doctoral degrees in Materials Science and Engineering from the University of Dayton and has in excess of 25 granted US patents. Prior to founding Next Science, Dr. Myntti led the biomaterials group at Medtronic Surgical Technologies.



#### JON SWANSON CHIEF OPERATING OFFICER

Jon oversees the Quality, Operations and Regulatory teams to deliver on development timelines and meet the goals set with our commercial partners. Jon joined Next Science from McKinsey, where he worked with various Fortune 500 companies to improve their product development and operations capabilities. Prior to McKinsey Jon was with Medtronic in a variety of senior leadership roles.



#### JACQUELINE BUTLER CHIEF FINANCIAL OFFICER

Jacqueline is a qualified Chartered Accountant with broad European and Australian experience in a variety of financial roles. Prior to Joining Next Science, Jacqueline worked as CFO and Company Secretary at Avira Resources Limited where she supported a successful ASX listing.



#### BYRON DARROCH PARTNERSHIPS

Byron leads the Partnership team for Next Science and is an accomplished business executive. He has a track record of delivering accelerated growth through implementation of innovative models and strategies. Prior to joining Next Science, Byron was General Manager at Atomo Diagnostics where he helped bring largescale HIV self-testing to the global market.

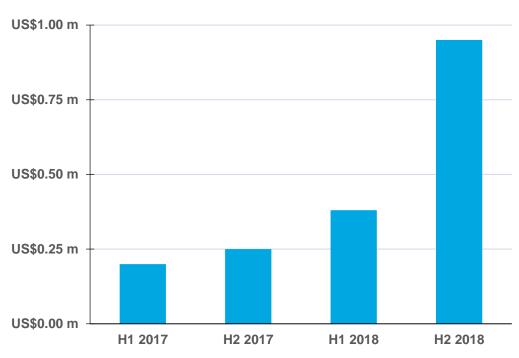
# **Business Update**

# FY 2019 Year to Date

- Q1 FY19 Sales
- New Scientific findings
- Regulatory update
- Commercial update
- Patents update
- Outlook

# **Rapid Market Penetration**

## SALES REVENUE SINCE PRODUCT LAUNCH

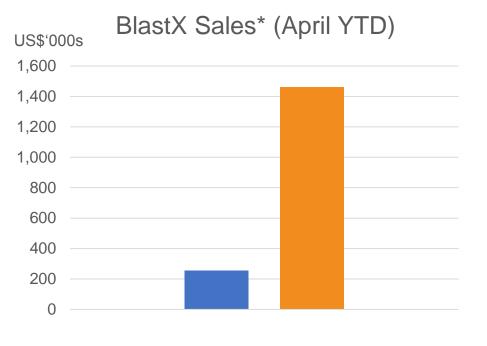


### **BACTISURE SALES REVENUE**

#### **BLASTX SALES REVENUE**

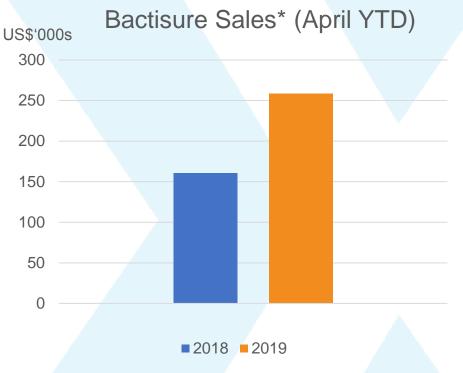


# Strong Sales Growth continues in 2019



#### 2018 2019

- BlastX distribution transitioned to 3M sales force in Q1 2019
- 3M sales training occurred in February
- Maintained top 24 BlastX customers and grew customer base
- 10 person partner support team to support the KCI Advanced Wound Care sales team, post the 3M acquisition of Acelity (KCI wound care products)



- 61% sales growth vs prior YTD
- Growing use of Bactisure outside of orthopaedics eg Breast reconstruction

# **Recent Scientific Presentations**

## In vitro Evaluation of Biofilm Disrupting Agents (BDA) against Candida auris and other Candida species

Authors: Jose A Vazquez1, Sushama Wakade1, Matt Myntti2, Elias Manavathu1

Medical College of Georgia at Augusta University, Augusta, GA, 2.Next Science, Jacksonville, FL., USA

Products tested: BlastX, TorrentX and Next Science Surface Disinfectant

Presented ECCMID Meeting April 13-16, 2019 Amsterdam, Netherlands

#### **Conclusion:**

The use of these novel BDAs with excellent antimicrobial and antifungal activity make them very valuable in eradicating surface and wound colonization of *Candida* sub species, including the MDR-*C. auris*, and thus possibly decrease the spread of this *Candida* sub species.

## A novel disruptive agent influences the wound healing process (animal study)

Author: Kayla Bounds

Texas Tech University Medical Health Centre, Lubbock TX, USA

Presented SAWC May 7 – 10, 2019 San Antonio TX

Conclusions: (In mouse studies)

- BlastX prevents overexuberant inflammation in a clean wound by reducing the level of pro-inflammatory cytokines while promoting the appropriate formation of blood vessels by increasing CXCL10 on Day 1
- » BlastX accelerates wound healing by enhancing the numbers of M2 macrophages on day 3
- » BlastX advances re-epithelialization on day 7 increasing levels of C/C involved in keratinocyte hyperplasia

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# **Commercial Development: Appointment of Sinus Partner**

Grace Medical We Listen.

# Partner, Deal & Product

#### Grace Medical (GM)

- » US-based Grace Medical, is a recognized global market leader in the ear, nose, throat (ENT) marketplace.
- It offers an extensive range of products for use in ENT procedures.
   Deal
- » Grace Medical appointed as exclusive distributor of Next Science Sinus Lavage in USA, Europe, Australia & other key markets.
- » Grace Medical has committed to meeting stipulated performance criteria, NXS has maintained flexibility should Grace Medical not meet these measures.
- The 5 year distribution agreement commences upon the date that NXS first obtains regulatory approval in any country within the territory granted to Grace Medical. It will automatically renew thereafter for successive 5 year periods subject to either party having a right to terminate on 6 months' written notice before the end of the initial or any subsequent terms and various other events which entitle a party to terminate.

#### Product

- » NXS's Sinus Lavage, using Xbio<sup>™</sup> Technology, is designed to be used as an adjunct to both FESS and balloon sinuplasty procedures to aid in the washing away and removal of debris, including microorganisms from the sinus cavities.
- NXS expects regulatory submissions to be filed in the US and Europe Q3, 2019.

# **Opportunity**

### Opportunity

- An estimated 40 million people in the USA and 35 million people in Western Europe have Chronic Rhinosinusitis (CRS).
- In the US, ~1.3 Million people annually seek out an ENT specialist for treatment, with about 550,000 opting for Functional Endoscopic Sinus Surgery (FESS) typically once medical management of CRS has failed.
- » A relatively new procedure, balloon sinuplasty, which is a less invasive procedure, is also gaining in popularity.
- These procedures, although effective, do not address the problem of biofilm in CRS. Microbial biofilms have been implicated as a cause of recalcitrant CRS.
- In the case of CRS patients, biofilms may also be a significant factor to a stronger inflammatory response that perpetuates the sinusitis symptoms.
- Independent research has shown that biofilms are one of the leading causes of antibiotic failure in CRS in patients.
- In several studies, biofilms have been shown to be prevalent in anywhere from 45-100% of CRS patients.

# **Product Regulatory Timelines**

BlastX	CE Mark Submitted, Canada Health – July, Australia Submission – H2 2019
Bactisure	CE Mark under review, Canada Health – July, Australia Submission – H2 2019
TorrentX	<ul> <li>510 (k) submitted expanding claims. As a 510 (k) device, TorrentX can be sold through the same channels into the hospital as existing Next Science products with expanded indications</li> </ul>
Hard Surface Disinfectant	<ul> <li>Submitted to EPA with request for Biofilm effectiveness claim Q2 2019. Expected to be approved H1 2020</li> </ul>
Sinus	Submissions CE Mark, FDA 510 (k) – H2 2019, Canada and Australia – H2 2019
Middle Ear	<ul> <li>Submissions CE Mark, FDA 510 (k) – H2 2019</li> </ul>
MIS Lavage	<ul> <li>Submissions CE Mark, FDA 510 (k) – H2 2019</li> </ul>
Audit	<ul> <li>Qualified the business through the MDSAP audit, allowing the company to file previously FDA approved or CE Mark approved products in Canada, Australia, Japan and Brazil</li> </ul>

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# **Expansion of Patent Portfolio**

## **17** PATENTS **49** APPLICATIONS GRANTED **49** PENDING TO DATE

Next Science Intellectual Property covers its Xbio<sup>TM</sup> platform technology and provides broad protection for the solution parameters of the given formulations. The patents cover the types of the ingredients (such as surfactants) not the specific ingredients.

Patents also cover the method of action as well as composition of matter. Specific patents cover claims in wound gel, acne treatment, oral rinse, post surgical disinfection and solid forms. Patent coverage now in China.



PATENTS GRANTED: USA • EUROPE • JAPAN • AUSTRALIA PATENTS PENDING: BRAZIL • CANADA • CHINA • INDIA

## Outlook 2019 and beyond

- Sales growth expected to continue through compounding growth drivers
  - Increased market penetration of existing products in the US market
  - Further focus from 3M in wound care post acquisition of Acelity (global wound care products)
  - Geographic expansion of sales outside the US (Canada Q3, Australia Q4, Europe CE submitted)
  - Development and commercialisation of new products and applications in human health
  - Extension of technology into applications outside human health
- Acne product to launch in Australia in H2 2019
  - Additional products to enter the Australian market: BlastX, Bactisure H2 2019
- ForrentX and SurgX distribution agreements under negotiation
- Strong R&D pipeline with new scientific findings continues to expand the applications for Xbio
  - Wider adoption of the technology by different medical specialties grows the clinical experience with the technology and expands the research base on the applications for the technology
- New Regulatory approvals expected in the US, Europe, Canada and Australia

# **Investment Summary**

- Unique, non-toxic technology with proven efficacy in eradicating both biofilm and bacteria
  - Solves a clear unmet medical need and avoids creating antibacterial resistance
  - Proven in over 80,000 patient treatments, multiple FDA Clearances with broad IP protection
- Proven commercial demand
  - Validated by global distribution agreements with industry leaders (Zimmer Biomet, 3M and Grace Medical)
- 5 products in market and 4 products submitted for clearance by end 2019 targeting high value market segments with unmet needs
  - Extensive product pipeline across medical devices, OTC drugs and prescription pharmaceuticals
  - Key markets include Chronic Wounds US\$50b, Surgical Site infections US\$3.5-US\$10b, Prosthetic Joint infections US\$4b, Chronic Sinusitis US\$60b and many others
- Continuing sales growth outlook by leveraging distribution partners, new product and new market entry, and geographic expansion
- High margin (>80%) and highly scalable production via multiple contract manufacturers
- Strong management team and clear growth strategy
- Fully funded to accelerate commercialisation and product development
- **NEXT SCIENCE**<sup>®</sup> Investor Presentation May 2019

# **NEXT SCIENCE**<sup>®</sup> Break through biofilm.

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Additional biofilm education can be found at: biofilm.healthcare