

AFT PHARMACEUTICALS

Investor Presentation May 2016

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PRESENTING TODAY . . .



Dr Hartley Atkinson

Founder and Chief Executive Officer

Founded AFT in 1997 along with his wife, Marree

Previously Sales & Marketing Director, Medical Director, Product Manager and Medical Manager at Swiss multinational pharmaceutical company, Roche



Malcolm Tubby

Chief Financial Officer

Involved with AFT since its establishment in 1997, providing financial, operational and governance expertise

Experience in senior finance positions in public and private companies in the pharmaceuticals (Allergan), fast-moving consumer goods (Frucor Beverages), insurance and healthcare industries

FY2016 HIGHLIGHTS

98 countries that *Maxigesic* is licensed in

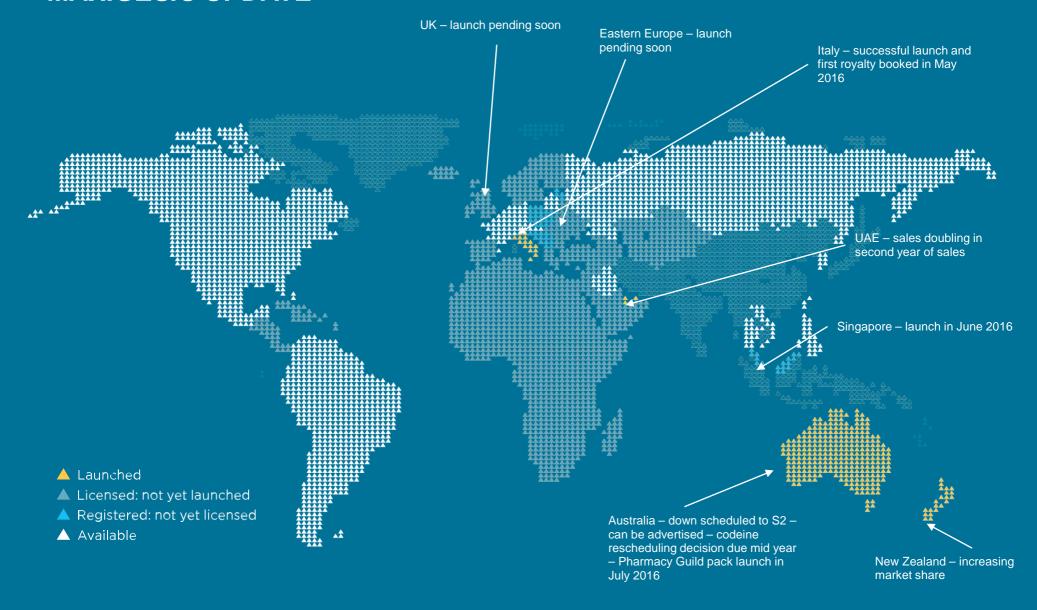
countries that *Maxigesic* is launched and sold in

number of clinical studies AFT will have running in FY2017

\$65.8m total income for FY2016*

\$28.1 m available cash as at 31 March 2016

MAXIGESIC UPDATE



MAXIGESIC HIGHLIGHTS

Additional out-licensing and distribution agreements for *Maxigesic* oral dose forms have been secured to increase the number of countries to 98 as at 31 March 2016.

Clinical Trials for *Maxigesic* oral dose forms are well underway with studies running in Amman, Jordan; various centres in New Zealand; Cardiff, Wales; Melbourne, Australia; Guadalajara; Mexico.

An IND (Investigational New Drug) Application has been successfully opened with the FDA for *Maxigesic* IV and the first clinical study under the IND is underway in Christchurch, New Zealand. The next study is planned to start in 2016 in the United States (Texas and Maryland).

Regulatory applications for the first additional *Maxigesic* oral dose forms to be filed from this year.

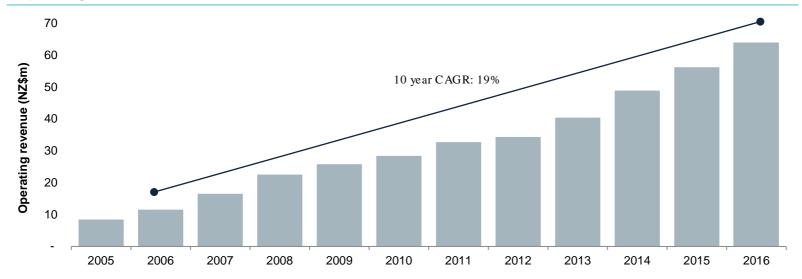
Additional out-licensing and distribution agreements for *Maxigesic IV* have also been secured to now reach 69 countries.

Additional out-licensing agreements are expected to be announced over the coming FY2017 financial year. There are currently 109 countries out-licensed for *Maxigesic* oral dose forms and 80 for *Maxigesic IV*.



FINANCIAL PERFORMANCE - REVENUE GROWTH

Operating revenue, FY2000 - FY2016



FY2015 Operating revenue by region

FY2016 Operating revenue by region



Southeast Asia

Singapore

10 approved products12 more in registrationJune launches of 4 products

Malaysia

4 approved products 13 more in registration

Rest of World

Increasing launches 7 countries in Q2-3 FY2017

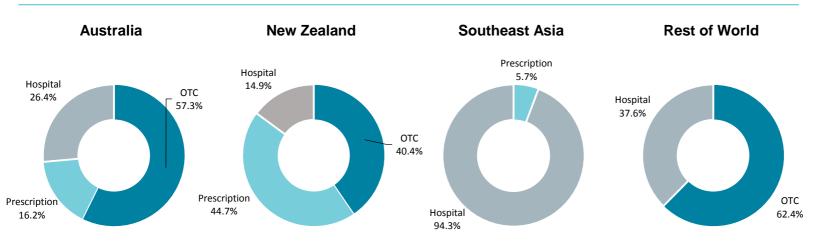


FINANCIAL PERFORMANCE – REVENUE BY REGION AND CHANNEL

Operating revenue by region, FY2014 - FY2016

NZ\$000's, year ended 31 March	2014	% of total	2015	% of total	2016	% of total
Australia YoY Growth	20,035 -	40.9%	26,324 31.4%	46.8%	31,224 18.6%	48.8%
New Zealand YoY Growth	28,790 -	58.8%	29,398 2.1%	52.3%	31,135 <i>6.0%</i>	48.7%
Southeast Asia YoY Growth	-	-	161 -	0.3%	648 <i>302.5%</i>	1.0%
Rest of World YoY Growth	114 -	0.2%	358 214.0%	0.6%	1,008 <i>181.6%</i>	1.6%
Total Operating Revenue YoY Growth	48,939 -	100.0%	56,241 <i>14</i> .9%	100.0%	64,014 13.9%	100.0%

Operating revenue by channel by region, FY2016



FINANCIAL PERFORMANCE - SUMMARY P&L

NZ\$000's, year ended 31 March	2016	% of revenue	2015	% of revenue
Revenue	64,014		56,241	
Cost of sales	(40,435)	63.2%	(35,083)	62.4%
Gross profit	23,579	36.8%	21,158	37.6%
Gross profit %				
Other income	2,295	3.6%	1,270	2.3%
Selling and distribution expenses	(19,634)	30.7%	(17,157)	30.5%
General and administrative expenses	(6,804)	10.6%	(5,475)	9.7%
Research and development expenses	(8,092)	12.6%	(5,761)	10.2%
Equity accounted loss of joint venture entity	(302)	0.5%	-	-
Operating loss	(8,958)		(5,965)	
Finance income	291		35	
Finance costs	(2,019)		(7,225)	
IPO, listing and capital raising costs	(2,623)		-	
Loss before tax	(13,309)		(13,155)	
Tax benefit / (expense)	42		282	
Loss after tax	(13,267)		(12,873)	

FINANCIAL PERFORMANCE – SUMMARY BALANCE SHEET

NZ\$000's, as at 31 March	2016	2015
ASSETS		
Current assets		
Inventories	17,686	14,686
Trade and other receivables	16,288	11,251
Cash and other equivalents	28,055	4,700
Current income tax asset	26	88
Total current assets	62,055	30,725
Non-current assets		
Property, plant and equipment	407	411
Intangible assets	2,111	1,669
Deferred income tax assets	546	408
Investment in joint venture entity	185	
Total assets	65,304	33,213
LIABILITIES		
Current liabilities		
Trade and other payables	10,428	8,258
Provisions	3,083	1,890
Derivative liabilities	402	305
Total current liabilities	13,913	10,453
Non-current liabilities		
Interest bearing liabilities	23,161	20,739
Total liabilities	37,074	31,192
Equity		
Share capital	53,902	12,892
Retained earnings	(25,637)	(10,732)
Cash flow hedge reserve	· - 1	(305)
Share options reserve	65	· -
Foreign currency translation reserve	(100)	166
Total equity	28,230	2,021
Total liabilities and equity	65,304	33,213



FINANCIAL PERFORMANCE – SUMMARY CASHFLOW STATEMENT

NZ\$000's, year ended 31 March	2016	2015
Net cash (used in)/generated from operating activities	(14,429)	(13,105)
Net cash (used in) investing activities	(694)	(483)
Net cash generated from financing activities	36,705	17,135
Net increase in cash	21,582	3,547
Impact of foreign exchange on cash and cash equivalents	1,773	(95)
Opening cash and cash equivalents	4,700	1,248
Closing cash and cash equivalents	28,055	4,700

CLINICAL STUDY PROGRAM

- 12 studies across the AFT portfolio
- 1500+ patients
- 7 countries Australia, Jordan, Mexico, New Zealand, Russia, United Kingdom, United States
- Clinical Trials for *Maxigesic* oral dose forms are well underway with studies running in Amman,
 Jordan; various centres in New Zealand; Cardiff, Wales; Melbourne, Australia; Guadalajara; Mexico.
- An IND (Investigational New Drug) Application has been successfully opened with the FDA for
 Maxigesic IV and the first clinical study under the IND is underway in Christchurch, New Zealand. The
 next study is planned to start in 2016 in the United States (Texas and Maryland).
- NasoSURF Device Clinical Studies to start in 2016

NASOSURF NEBULISER DRUG DELIVERY AND TREATMENT SINUS CONDITIONS

Product description	A handheld ultrasonic nasal mesh nebuliser for the intranasal delivery of medication and treatment of chronic sinusitis
Rationale for investment in product	 To expand our existing allergy and hospital product ranges locally Significant global potential
Current status	 Under development Pilot scale production underway (May 20 units; July 100 units; August 300 units)
Our near term plans	 FDA pre-IND meeting in preparation (Aug-Sep 2016) Distribution studies – healthy volunteers (Q3 2016) patients (Q3-4 2016) Conscious sedation PK studies (Q3-4 2016) Conscious sedation Clinical Studies (Q4 2016 – Q1 2017) Register a Class I medical device (Q1 2017) Post Sinus Surgery studies (Q4 2016 – Q2 2017) Drug delivery use sales (Class IIa) late 2017 early 2018

The Naso SURF Nebuliser has desirable features over currently marketed nebulisers, which are not approved for delivery of specific drugs intranasally and do not possess a number of the advantages of the Naso SURF Nebuliser



Sales will be generated from

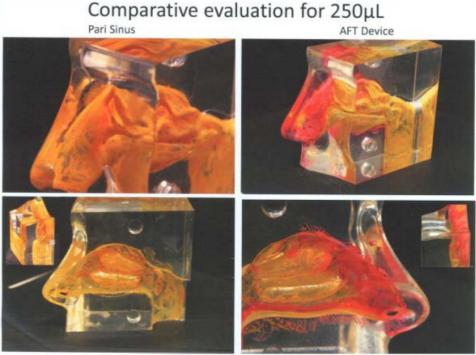
- 1) device sales,
- 2) a per use charge administered through RFID (radio frequency identifier) cards, and
- 3) consumables

NASOSURF VERSUS GERMAN GOLD STANDARD A F Tpharmaceuticals

NasoSURF vs Pari Sinus

IntraNasal Distribution





SUMMARY OF NEAR TERM PLANS



Launch Maxigesic in new countries



Advance *Maxigesic* registrations in North America Further licensing agreements



Clinical studies of key innovative products



NasoSURF clinical trials with pilot devices Achieve first registrations for NasoSURF as a Medical Device



Build revenues of new OTC products launched in Australia Build *Maxigesic* market share post scheduling changes



Launch OTC products in Southeast Asia Complete further registrations and launches