

Delivering on our strategy

FMP & Yellowstone Investor Evening 4th Nov 2019

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Allergy Therapeutics

Leading, fully integrated biopharmaceutical company based in the UK

Provide treatments that have potential to cure disease, not just symptoms. Focus on moderate to severe patients

PQ® Platform enabling ultrashort course treatment for grass, tree and ragweed allergies

Headquartered in Worthing, Sussex with about 500 employees

9% compound annual revenue growth achieved over the last 20 years

Leading provider of subcutaneous aluminium-free allergy vaccines

Spun out of Smith Kline Beecham in 1999

Market capitalisation of about £90m, AIM ticker LSE:AGY





Three Pillars to Growth: Advancing a Leading Allergy Immunotherapy Company



Three pillars to the business

01

Expanding in Europe

Strongly performing profitable business

Growing market share and additional product registrations



02

Strong pipeline

New technologies underpin pipeline breadth and depth

Investment strategy supported by growing revenue stream



03

Preparing for US entry

Significant opportunity in largest allergy market

Changing regulatory and reimbursement environment to drive market share towards Allergy's products

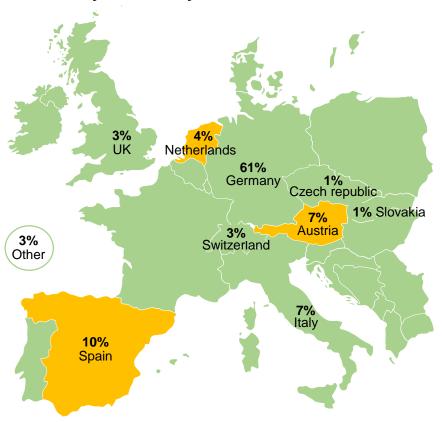




Sales breakdown for FY 2019



Sales by country

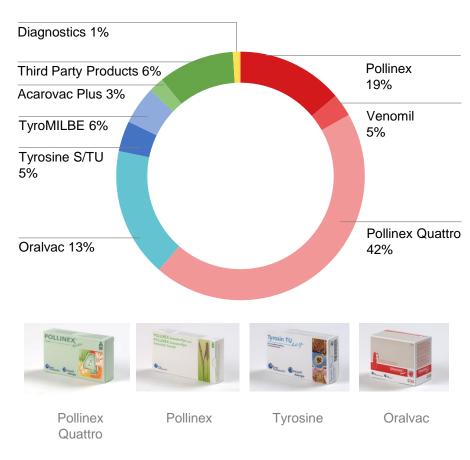


¹ Sales breakdown based on gross sales at budget exchange rates (before freight, rebates and exchange): £69.3 million.

After deducting rebates, freight charges and foreign exchange adjustments, total sales for FY2019 is £73.7 million

*Allergy Therapeutics currently has no products licensed for sale in the USA

Sales of £73.7m by product1*



Grass MATA MPL

Successful end of Phase II meetings with PEI and FDA

Product is one Phase III efficacy trial and completion of safety database away from filing in US

Draft protocol agreed including dosing for Phase III trial

Key product for US introduction – Ragweed would be product to follow

Timing of start of Phase III Grass trial autumn of 2020 subject to final adjustments



Birch MATA MPL

Primary end point of Phase III trial not met - Extensive work undertaken to understand results

Group in dialogue with PEI over results

Trial showed biomarker effect as well as expected safety profile

Secondary endpoint analyses of immunoglobulin markers including IgG and IgG4 showed highly statistically significant differences between active and placebo

Two previous Phase II dosing trials demonstrated efficacy and a classic dose response curve and a 32% reduction in symptoms

ATL will learn from the trial and put new ideas into the following field trials



Preclinical Pipeline: Polyvac peanut product

Single dose of virus like particle (VLP) combined with recombinant peanut allergen successfully protects against anaphylaxis when challenged with peanut Industrial scale-up progressing well with first in-human trial anticipated summer 2020

Those vaccinated with candidate vaccine exhibited no symptoms compared to placebo, when challenged with peanut

Peanut represents a new opportunity into \$8bn* worldwide food allergy market

Safety profile of product evaluated and found **not** to induce anaphylaxis

Successful meeting with Swissmedic and PEI on outline protocol for first inhuman trial

Allergy Therapeutics PLC Positive results achieved from preclinical research of Polyvac Peanut

^{*}The Journal of Allergy and Clinical Immunology 2016. 1% of US population. EACCI Food Allergy and Anaphylaxis Guidelines Group 2016 0.2% of Western European Population. Management assumption of annual treatment of \$2k

US allergy immunotherapy market represents a significant and attractive commercial opportunity



\$2bn**

estimated allergy immunotherapy market

\$300-400m**

Estimated peak grass sales

~100 injections

Current treatment: up to 100 injections over 3-5 years***

16%

Some adherence levels as low as 16%*

None

Currently no registered injected products

4+1products

Grass, Tree, Ragweed and Mite plus Peanut (new area of market)



^{*}Hankin CS, Cox L, Lang D, et al 2007 JACI

^{**}Internal estimate

^{***}Professor Lawrence DuBuske MD

Allergy Therapeutics PLC Summary and Outlook

2019 highlights



PQ Grass

Phase III

Progression in US in H2 2020

PQ Birch

Phase III

Primary endpoint not met – comprehensive review of dataset underway

Increase in market share over 12 months to June 2019

0.5 point increase in market share

increase in revenue to

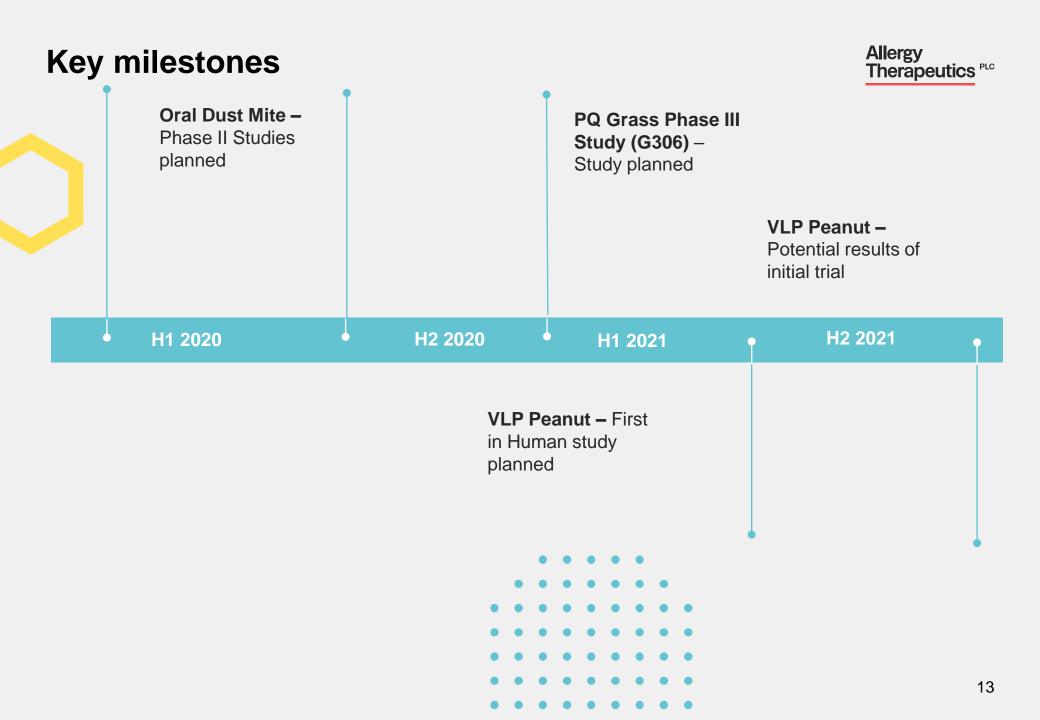
£73.7m (2018 £68.3m)

Operating profit pre R&D up 22%

Net profit of £3.5m (2018: £7.5m loss)

Successful completion of Litigation with £6m settlement

Cash balance of £27.4m (2018: £15.5m)



House dust mite product

Primary endpoint of safety and tolerability achieved in Phase I trial

Phase II dosing trial anticipated in 2020

Market opportunity of

\$3-4bn* worldwide with only

Europe partly tapped already

Acarovac product without MPL growing well in Spain and Austria

Potential of 8 injection model compared to 12-15 average of competitors and once a day for 3 years oral treatment Potential additional product in US portfolio following two Phase III trials

Allergy Therapeutics PLC **Short-course** product with global potential

Innovative, Broad Pipeline and Marketed Products



	Pre-clinical		Phase I	Phase II	Phase III	Market/Registered	Also available as a Named Patient Product
Grass MATA		Shor	t-course SCIT				,
Tree MATA		Shor	t-course SCIT				
Ragweed MATA	*	Shor	t-course SCIT				·
Bee Venom SCIT		Shor	t-course SCIT				·
Wasp Venom SCIT		Shor	t-course SCIT		,		,
Grass MATA MPL		Shoi	t-course Grass SC	T with MPL		>	
Birch MATA MPL		Shor	t-course Birch SCI	T with MPL		>	
Ragweed MATA MPL		Shor	t-course Ragweed	SCIT with MPL		>	
Trees MATA MPL		Shor	t-course Tree SCIT	with MPL			
Oral Grass, Trees & House Dust Mite			otherapy with flexib	ble-dosing			
Modified Mite Platform	Short-course HDM SCIT +	mod MPL	ified Allergen				
Peanut SCIT	Short-course Peanut SCIT						

Cutting-edge Platform Technologies

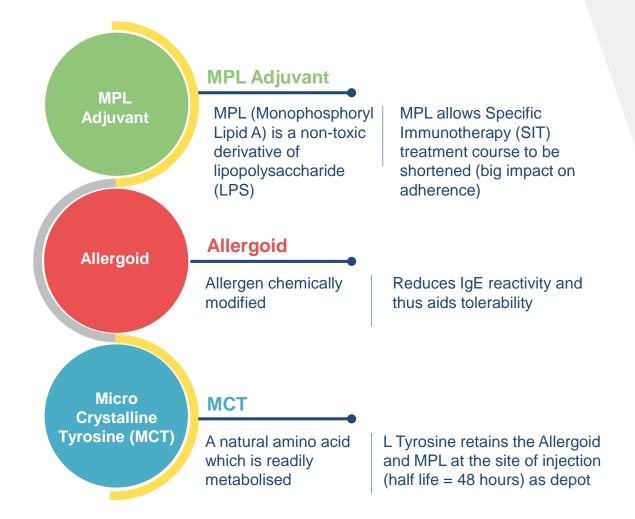


	Modified Allergen (Allergoid)	Native Allergen	Recombinant Allergen	Microcrystalline Tyrosine (MCT)	Monophosphoryl Lipid A (MPL)	Virus-Like Particles (VLP)
MATA						
MATA MPL (PQ)	⊘			Ø	Ø	
Sublingual						
Mite SCIT	⊘					
Mite SCIT + MPL	Ø			Ø	Ø	
Venom SCIT						
Peanut*				Ø		Ø

^{*} Product under pre-clinical investigation, full product profile yet to be determined

PQ: Differentiated platform approach enhances compliance, leads to higher efficacy and successful outcomes

Allergy Therapeutics PLC



Allergy Therapeutics PLC

Allergy Therapeutics: Company with Solid Sales and Global presence

Sales and marketing network comprising c.140 European sales force

