

Market Data

Aptorum Group

NASDAQ: APM

Fiscal Year	December
Industry	Biotechnology
Recent Price	\$1.00
Market Cap	\$35.7M
Shares Out.	35.7M
Float	9.1M
Avg. Volume (90-day)	124K
Revenue (ttm)	\$1.54M
Cash (mrq) ¹	\$8.13M
LT Debt (mrq) ¹	\$0.0M

As of May 10, 2022

¹ as of December 31, 2021

aptorumgroup.com

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Company Overview

Aptorum Group Limited (NASDAQ: APM) is a pharmaceutical company dedicated to developing and commercializing novel therapeutics to tackle unmet medical needs. Aptorum's current drug pipeline includes indications in orphan diseases, infectious diseases, and metabolic diseases. The Company launched two new clinical trials in 2021. Aptorum's Smart-ACT™ platform is designed to bring an average of three drug candidates for orphan diseases to clinical trials every 12-18 months. The Company is now preparing to launch a dietary supplement for women undergoing menopause and experiencing related symptoms, including osteoporosis. Targeting a global woman's health supplement market that is expected to reach \$17 billion in 2025, Aptorum is expected to generate near-term revenue with significant long-term growth potential.

Value Proposition

Aptorum's four core assets (RPIDD, SACT-1, ALS-4, NLS-2) target indications with a combined market value of more than \$250 billion. The company is building a diverse portfolio of over 12 projects primarily in therapeutics and diagnostics across a wide range of unmet disease areas. Aptorum translates early-stage discovery into proof-of-concept clinical stages and transforms clinical outcomes through licensing, co-development, and commercialization. The company's proprietary Smart-ACT™ platform combines a computational approach and wet lab validation to screen over 2,600 approved small molecule drugs. This process allows Aptorum to identify candidates that can be repurposed for selected orphan or unmet diseases. This platform's first proof-of-concept asset is SACT-1 for Neuroblastoma, a cancer mostly affecting children that develops from immature nerve cells. The Company announced positive data from its recently completed Phase 1 trial of SACT-1 for Neuroblastoma in May 2022. SACT-1 was also recently used to identify at least three small molecule drugs targeting COVID-19, which Aptorum is investigating in collaboration with Covar Pharmaceuticals and the University of Hong Kong's Microbiology Department.

Project	Candidate / Modality	Indication	Development Stage							
			Target Identification & Selection	Lead Discovery	Lead Optimization	IND (Or IND equivalent Enabling)	Clinical Trial Application Submission	Phase I	Phase II / III	
Article's Series										
ALS-4	Small molecule	Treatment of bacterial infections caused by <i>Staphylococcus aureus</i> including MRSA								
SACT's Series			Computational Discovery	In Vitro Validation	Existing PH/III Clinical Safety Data ¹	In Vivo Validation	IND Preparation & Submission	Phase I	Phase II / III	
SACT-1	Repurposed small molecule	Neuroblastoma and other potential cancer types								
RPIDD			Development and Experimentation		Product Optimization		Clinical validation & Pre-Commercialization preparation		Commercialization	
RPIDD	Liquid biopsy rapid pathogen diagnostics	Pathogen molecular diagnostics								
Project			Modality		Target Customer		Formulation		Commercialization and Distribution	
NativusWell® DOI (NLS-2) ²			Dietary Supplement		Women undergoing menopause					
Project	Candidate / Modality	Indication	Development Stage							
			Computational Discovery	In Vitro Validation	Existing PH/III Clinical Safety Data ¹	In Vivo Validation	IND Preparation & Submission	Phase I	Phase II / III	
SACT's Series										
SACT - COVID19	Repurposed small molecule	Coronavirus Disease 2019 (COVID-19)								
SACT-2	Repurposed small molecule	To be disclosed								
SACT-3	Repurposed small molecule	To be disclosed								
Article's Series			Target Identification & Selection	Lead Discovery	Lead Optimization	IND (Or IND equivalent Enabling)	Clinical Trial Application Submission	Phase I	Phase II / III	
ALS-1	Small molecule	Treatment of viral infections caused by influenza virus A								
ALS-2/3	Small molecule	Treatment of gram+ve bacterial infections								
Discovery Stage										
DLS-1+2	Small molecule & Degraders	NSCLC with mutation								
DLS-3	Small molecule	Autoimmune								

Current Discovery & Development Pipeline

Investment Highlights

- Four core assets showing significant progress:
 - **Rapid Pathogen Identification and Detection Diagnostics (“RPIDD”)** technology, a rapid, accurate, cost-effective and untargeted method to identify and detect existing or emerging unknown pathogens through liquid biopsy. Such unknown pathogens include DNA/RNA-based viruses such as coronavirus, antibiotic-resistant bacteria, fungi, etc.
 - **SACT-1**, a repurposed drug candidate for treating neuroblastoma discovered through its Smart-ACT™ drug discovery platform; received Orphan Drug Designation for SACT-1 in January 2022
 - **ALS-4**, a small drug molecule candidate for the treatment of infections caused by *Staphylococcus aureus* including *Methicillin-resistant Staphylococcus aureus* (MRSA)
 - **NLS-2**, a dietary supplement for the relief of menopausal symptoms.
- Clinical trials:
 - Recently completed Phase 1 trial SACT-1 for neuroblastoma treatment (global c. \$2.6 billion market); trial generated positive safety and bioavailability data to advance program
 - ALS-4 for *Staphylococcus aureus* (incl. MRSA) saw positive interim results for Phase 1 trial (global c.\$3.0 billion market) with no Serious Adverse Events (SAE) or clinically relevant changes in vital signs
- Near-term commercialization of dietary supplement for woman’s menopausal health (global c. \$17 billion supplement market)
- Three additional SACT programs underway, including investigation of at least three repurposed drug candidates for coronavirus (COVID-19) in collaboration with Covar Pharmaceuticals and The University of Hong Kong
- 12+ therapeutic candidates under development in areas including infectious diseases, gastrointestinal microbiome, and drug repurposing for orphan diseases; representing a combined over \$8 billion+ global market opportunity
- Expanding IP protections; two new patents granted in Q1 2022
- Over 50+ staff, clinical advisors, and consultants with vast experience in drug development and clinical trials, including US FDA, EMA, and NMPA purposes.