

ASX ANNOUNCEMENT

Results of Annual General Meeting

Sydney 27 November 2020. Actinogen Medical ASX:ACW ('ACW' or 'the Company') advises that its Annual General Meeting of Shareholders was held today at 10.30 am AEDT.

In accordance with Listing Rule 3.13.2 and section 251AA(2) of the Corporations Act 2001 (Cth), the Company advises that the details of the resolutions and the proxies received in respect of each resolution are set out in the attached proxy summary.

ENDS

Actinogen Medical

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Announcement authorised by the Board of Directors of Actinogen Medica About Actinogen Medical

Actinogen Medical (ASX:ACW) is an ASX-listed biotechnology company developing novel therapies for neurological, psychiatric and metabolic diseases associated with chronically elevated cortisol. The company is currently developing its lead compound, Xanamem, as a promising new therapy for Alzheimer's disease, Fragile X syndrome, schizophrenia and diabetes. The cognitive dysfunction, behavioural abnormalities, and neuropsychological burden associated with these conditions is significantly debilitating for patients, and there is a substantial unmet medical need for new and improved treatments.

About Xanamem™

Xanamem's novel mechanism of action works by blocking the production of intracellular cortisol – the stress hormone – through the inhibition of the 11β -HSD1 enzyme in the brain. There is a strong association between persistent stress and the production of excess cortisol that leads to detrimental changes in the brain, affecting memory, cognitive function and behaviour and neuropsychological symptoms. The 11β -HSD1 enzyme is particularly highly concentrated in the hippocampus and frontal cortex, areas of the brain impacted by a number of diseases and disorders, including Alzheimer's disease, Fragile X syndrome, schizophrenia, diabetes and other conditions associated with chronically raised cortisol.

The Company's XanaHES Phase I trial exploring the safety and tolerability of Xanamem 20mg once daily in healthy elderly volunteers, confirmed the drug's safety profile with no treatment-related serious adverse events. Additionally, the trial demonstrated that Xanamem produced a statistically significant improvement in cognition over placebo, which, along with other recently generated data, confirms 11β -HSD1 inhibition by Xanamem as a promising potential treatment for cognitive impairment associated with raised cortisol.

The Company plans to initiate Phase II studies of Xanamem in various disease areas in 2021, including MCI due to Alzheimer's disease, and Fragile X syndrome.

Xanamem is an investigational product and is not approved for use outside of a clinical trial by the FDA or by any global regulatory authority.

Xanamem™ is a trademark of Actinogen Medical.

Disclaimer

This announcement and attachments may contain certain forward-looking statements that are based on subjective estimates and assumptions and relate to circumstances and events that have not taken place and may not take place. Such forward looking statements involve known and unknown risks, uncertainties, and other factors (such as significant business, economic and competitive uncertainties and contingencies, and regulatory and clinical development risks and uncertainties) which may cause the actual results or the performance of Actinogen Medical to be materially different from the results or performance expressed or implied by such forward looking statements. Past performance is not a reliable indicator of future performance. There can be no assurance that any forward-looking statements will be realised. Actinogen Medical does not make any representation or give any warranty as to the likelihood of achievement or reasonableness of any forward-looking statements.

Actinogen Medical encourages all current investors to go paperless by registering their details with the designated registry service provider, Link Market Services.



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ACTINOGEN MEDICAL LIMITED

RESULT OF ANNUAL GENERAL MEETING (ASX REPORT)

ANNUAL GENERAL MEETING Friday, 27 November, 2020

As required by section 251AA(2) of the Corporations Act 2001 (Commonwealth) the following statistics are provided in respect of each resolution on the agenda.

Resolution Voted on at the meeting			Proxy Votes (as at proxy close)				Poll (Manner in which votes were cast in person or by proxy on a poll (where applicable) on a poll at the meeting			
No	Short Description	Strike Y/N/NA	For	Against	Discretionary (open votes)	Abstain	For	Against	Abstain **	Result
1	ADOPTION OF REMUNERATION REPORT	N	285,982,749 82.53%	11,224,116 3.24%	49,302,088 14.23%	2,231,087	338,088,927 96.79%	11,224,116 3.21%	2,231,087	Carried
2	RE-ELECTION OF DR GEORGE MORSTYN	NA	307,793,338 86.03%	610,000 0.17%	49,362,088 13.80%	3,146,087	359,959,516 99.83%	610,000 0.17%	3,146,087	Carried
3	ADOPTION OF NEW CONSTITUTION	NA	305,634,940 85.56%	2,223,398 0.62%	49,362,088 13.82%	3,691,087	357,801,118 99.38%	2,223,398 0.62%	3,691,087	Carried
4	APPROVAL OF INCREASED PLACEMENT CAPACITY	NA	300,193,832 83.57%	9,677,506 2.69%	49,362,088 13.74%	1,678,087	352,360,010 97.33%	9,677,506 2.67%	1,678,087	Carried
5	RATIFICATION OF PRIOR ISSUE OF PLACEMENT SHARES ISSUED UNDER LISTING RULE 7.1	NA	53,546,319 52.16%	605,944 0.59%	48,508,359 47.25%	3,400,000	104,858,768 99.43%	605,944 0.57%	3,400,000	Carried
6	RATIFICATION OF PRIOR ISSUE OF PLACEMENT SHARES ISSUED UNDER LISTING RULE 7.1A	NA	53,546,319 52.16%	605,944 0.59%	48,508,359 47.25%	3,400,000	104,858,768 99.43%	605,944 0.57%	3,400,000	Carried

^{** -} Note that votes relating to a person who abstains on an item are not counted in determining whether or not the required majority of votes were cast for or against that item