FORM 7

MONTHLY PROGRESS REPORT

Name of Listed Issuer: <u>Algernon Pharmaceuticals Inc.</u> (the "**Issuer**" or the "**Company**").

Trading Symbol: AGN

Number of Outstanding Listed Securities: <u>15,775,757</u>

Date: September 8, 2023

Report on Business

1. Provide a general overview and discussion of the development of the Issuer's business and operations over the previous month. Where the Issuer was inactive disclose this fact.

The Company's main activities continue to include advancing its three core programs, including its N,N-dimethyltryptamine ("DMT") stroke and traumatic brain injury (TBI) research programs, its idiopathic pulmonary fibrosis ("IPF")/chronic cough program with Ifenprodil and its chronic kidney disease ("CKD") research program with Repirinast.

During August, the Company continued planning activities to advance its core programs focusing on the planning of a phase 2 clinical trial for DMT in stroke and a phase 2b clinical trial for Ifenprodil in chronic cough.

The Company completed a feasibility study and has finalized its clinical trial design for a 40 patient phase 2 DMT stroke which will study an intravenous sub-psychedelic dose of DMT in patients who are hospitalized after having suffered an acute ischemic stroke.

The Company, with the assistance of Maxim Group LLC. ("Maxim"), continues to identify and evaluate potential mergers and acquisition and strategic opportunities, including the potential spin-off of the Company's Ifenprodil chronic cough research program.

2. Provide a general overview and discussion of the activities of management.

The Company's main activities have included advancing its three core programs, including its DMT stroke research program, its IPF/chronic cough program with Ifenprodil and its CKD research program with Repirinast.

During August, the Company continued planning activities to advance its core programs focusing on the planning of a phase 2 clinical trial for DMT in stroke and a phase 2b clinical trial for Ifenprodil in chronic cough.

The Company completed a feasibility study and has finalized its clinical trial design for a 40 patient phase 2 DMT stroke which will study an intravenous sub-psychedelic dose of DMT in patients who are hospitalized after having suffered an acute ischemic stroke.

The Company, with the assistance of Maxim, continues to identify and evaluate potential mergers and acquisition and strategic opportunities, including the potential spin-off of the Company's Ifenprodil chronic cough research program.

3. Describe and provide details of any products or services that were discontinued. For resource companies, provide details of any drilling, exploration or production programs that have been amended or abandoned.

None

4. Describe any new business relationships entered into between the Issuer, the Issuer's affiliates or third parties including contracts to supply products or services, joint venture agreements and licensing agreements etc. State whether the relationship is with a Related Person of the Issuer and provide details of the relationship.

None

5. Describe the expiry or termination of any contracts or agreements between the Issuer, the Issuer's affiliates or third parties or cancellation of any financing arrangements that have been previously announced.

None

6. Describe any acquisitions by the Issuer or dispositions of the Issuer's assets that occurred during the preceding month. Provide details of the nature of the assets acquired or disposed of and provide details of the consideration paid or payable together with a schedule of payments if applicable, and of any valuation. State how the consideration was determined and whether the acquisition was from or the dispositions was to a Related Person of the Issuer and provide details of the relationship.

None.

7. Describe the acquisition of new customers or loss of customers.

None.

8. Describe any new developments or effects on intangible products such as brand names, circulation lists, copyrights, franchises, licenses, patents, software, subscription lists and trade-marks.

None.

9. Report on any employee hiring's, terminations or lay-offs with details of anticipated length of lay-offs.

None.

10. Report on any labour disputes and resolutions of those disputes if applicable.

None.

11. Describe and provide details of legal proceedings to which the Issuer became a party, including the name of the court or agency, the date instituted, the principal parties to the proceedings, the nature of the claim, the amount claimed, if any, if the proceedings are being contested, and the present status of the proceedings.

None.

12. Provide details of any indebtedness incurred or repaid by the Issuer together with the terms of such indebtedness.

None.

13. Provide details of any securities issued and options or warrants granted.

Security	Number Issued	Details of Issuance	Use of Proceeds
N/A	N/A	N/A	N/A

14. Provide details of any loans to or by Related Persons.

None.

15. Provide details of any changes in directors, officers or committee members.

None.

16. Discuss any trends which are likely to impact the Issuer including trends in the Issuer's market(s) or political/regulatory trends.

None

The Issuer's but inherent to the Business" section the year ended www.sedar.com	Issuer's indu on of the Issud d August 31	ıstry. Pleas er's manage	se to the " ement discu	Risks Relaussion and	ited To The analysis for

Certificate Of Compliance

The undersigned hereby certifies that:

- 1. The undersigned is a director and/or senior officer of the Issuer and has been duly authorized by a resolution of the board of directors of the Issuer to sign this Certificate of Compliance.
- 2. As of the date hereof there is no material information concerning the Issuer which has not been publicly disclosed.
- 3. The undersigned hereby certifies to the Exchange that the Issuer is in compliance with the requirements of applicable securities legislation (as such term is defined in National Instrument 14-101) and all Exchange Requirements (as defined in CNSX Policy 1).
- 4. All of the information in this Form 7 Monthly Progress Report is true.

Dated: September 8, 2023

James Kinley			
Name of Director or Senior			
Officer			
"James Kinley"			
Signature			
CFO			
Official Capacity			

Issuer Details Name of Issuer	For Month End	Date of Report YY/MM/DD		
Algernon Pharmaceuticals Inc.	August 31, 2023	2023/09/08		
Issuer Address Suite 400 – 601 West Broadway Street				
City/Province/Postal Code	Issuer Fax No.	Issuer Telephone No.		
Vancouver, BC V5Z 4C2	NA	(604) 398-4175 ext. 701		
Contact Name	Contact Position	Contact Telephone No.		
James Kinley	CFO	(604) 398-4175 ext. 701		