August 4, 2021

Chris Ehrlich Chief Executive Officer Locust Walk Acquisition Corp. 200 Clarendon Street, 51st Floor Boston, MA 02116

Re: Locust Walk

Acquisition Corp.

Amendment No. 1 to

Registration Statement on Form S-4

Filed July 19, 2021 File No. 333-257091

Dear Mr. Ehrlich:

We have reviewed your amended registration statement and have the following

comments. In some of our comments, we may ask you to provide us with information so we

may better understand your disclosure.

Please respond to this letter by amending your registration statement and providing the

requested information. If you do not believe our comments apply to your facts and

circumstances or do not believe an amendment is appropriate, please tell us why in your

response.

After reviewing any amendment to your registration statement and the information you

provide in response to these comments, we may have additional comments. Unless we note

otherwise, our references to prior comments are to comments in our July 13, 2021 letter.

Amendment No. 1 to Registration Statement on Form S-4 filed July 19, 2021

Risk Factors

Risks Related to the Discovery, Development and Regulatory Approval of our Product

Candidates

Any difficulties or delays in the commencement or completion, or any terminations or

suspensions, of our current or planned clinical..., page 51

We note your response to our prior comment 15. Please also include disclosure

that pembrolizumab for the treatment of frontline NSCLC and other indications is owned

by Merck the first time

it is discussed in your risk factor disclosure, where appropriate.

We also refer to your

disclosure that your zotatifin product candidate is being evaluated in combination with other

FDA-approved inhibitors, such as fulvestrant and herceptin.

Chris Ehrlich

Locust Walk Acquisition Corp.

August 4, 2021

Page 2

Please revise your disclosure throughout the prospectus to clarify that these inhibitors are

therapies owned and developed by third parties.

Use of our product candidates could be associated with side effect, adverse events..., page 54

We note your response to our prior comment 18. Please revise your disclosure on page 55

further to clarify that certain treatment-emergent adverse events observed during your

Phase 2a trial of tomivosertib combined with Anti-PD-(L)1 agents were

Grade 3 in

severity, as referenced on page 172. Background of the Business Combination, page 114

3. We note your response to prior comment 13. Please expand your disclosure to discuss the $\ensuremath{\text{S}}$

valuations of comparable public companies and the analysis provided by Locust Walk

Partners that were considered by the LWAC board or tell us why you believe such $% \left(1\right) =\left(1\right) +\left(1\right) +\left($

disclosure would not be material to investors.

You may contact Michael Fay at 202-551-3812 or Angela Connell at 202-551-3426 if

you have questions regarding comments on the financial statements and related matters. Please $\,$

contact Jane Park at 202-551-7439 or Christine Westbrook at 202-551-5019 with any other questions.

Sincerely,

FirstName LastNameChris Ehrlich

Division of

Corporation Finance Comapany NameLocust Walk Acquisition Corp.

Office of Life

Sciences

August 4, 2021 Page 2

cc: Cheston J. Larson, Esq.

FirstName LastName