Grant E. Pickering President and Chief Executive Officer SutroVax, Inc. 353 Hatch Drive Foster City, California 94404

Re: SutroVax, Inc.

Amendment No. 1 to Draft Registration Statement on Form S-1 $\,$

Submitted December 31, 2019

CIK No. 0001649094

Dear Mr. Pickering:

We have reviewed your amended draft registration statement and have the following $% \left(1\right) =\left(1\right) +\left(1\right) +$

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 providing the requested information and either submitting $\ensuremath{\mathsf{E}}$

an amended draft registration statement or publicly filing your registration statement on $% \left(1\right) =\left(1\right) \left(1\right$

 ${\tt EDGAR.}$ 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 the information you provide in response to these comments and your $% \left(1\right) =\left(1\right) \left(1\right) +\left(1\right) \left(1\right) \left(1\right) +\left(1\right) \left(1\right) \left($

amended draft registration statement or filed registration statement, we may have additional comments.

Amendment No. 1 to Draft Registration Statement on Form S-1 $\,$

1. We note your response to our prior comment 1 and continue to object to

your characterization of SVX-24 as having "the potential to become the

standard of care." The qualifier, "if approved," does not address our concerns that this

language continues to imply that the product will be effective and will replace the current

standard of care before a competing vaccine does, neither of which is appropriate at this stage of development.

Please revise your registration statement to remove this language.

Grant E. Pickering

FirstName Inc.

SutroVax, LastNameGrant E. Pickering

Comapany NameSutroVax, Inc.

Prospectus Summary, page 1

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FirstName LastName

Risk Factors Summary, page 5

2. We note your response to our prior comment 5 and reissue. The risk that your intended

approach will not be sufficient for regulatory approval or that regulators will require field

efficacy trials or longer trials with more participants than you currently anticipate exists $% \left(1\right) =\left(1\right) +\left(1\right$

now. Please add a separate bullet point in this section to discuss that risk.

questions regarding comments on the financial statements and related matters. Please contact

Ada D. Sarmento at 202-551-3798 or Mary Beth Breslin at 202-551-3625 with any other

questions.

Sciences cc: Robert W. Phillips, Esq.