Prof. Ugur Sahin, M.D. Chief Executive Officer BioNTech SE An der Goldgrube 12 D-55131 Mainz Germany

Re: BioNTech SE

Registration Statement on Form F-1

Filed September 9, 2019 File No. 333-233688

Dear Dr. Sahin:

We have reviewed your 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 $% \left(1\right) =\left(1\right) +\left(1\right$

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 August 20, 2019 letter.

Registration Statement on Form F-1

Dilution, page 103

1. Please explain why the pro forma net tangible book value per share does not include the $\,$

receipt of proceeds of 190.2 million relating to ordinary shares registered before June 30,

 $\,$ 2019 under the Series B private placement that is included in the proforma

. . .

Capitalization.

Prof. Ugur Sahin, M.D.

FirstName SE

BioNTech LastNameProf. Ugur Sahin, M.D.

Comapany 19, 2019

September NameBioNTech SE

Page 2

September 19, 2019 Page 2

FirstName LastName

Management's Discussion and Analysis of Financial Condition and Results of Operations

Financial Operations Overview

Revenue, page 109

2. Please clarify in your disclosure that the significant increase in collaboration revenue from $\,$

the Sanofi agreement in 2018 was due to a reimbursement of 50% of Cellscript sublicense

costs pursuant to a separate Sub-sublicense agreement.

Biotech Business Unit

Comparison of the Six Months Ended June 30, 2019 and 2018

Revenue, page 115

3. You disclose that the increase in revenue in your Clinical segment from the six months ${\bf x}$

ended June 30, 2018 to the six months ended June 30, 2019 was predominantly due to the $\,$

progress of your collaboration agreement with Sanofi into the clinical stage from the

 $\bar{\ }$ research stage. However, the table on page 110 shows a decrease in revenue from the

Sanofi agreement for those periods. Please clarify.

Business

Eli Lilly TCR Therapy Collaboration, page 206

We note your reference here to "low double-digit percentages." Please revise your

disclosure to narrow the royalty range to no more than ten percentage points (for example,

between twenty and thirty percent).

Penn Agreement, page 239

We note your response to our prior comment 4 and reissue in part.

Please quantify more

specifically the maximum aggregate milestone payments under the Penn Agreement, as

opposed to providing a wide range of potential milestone payments of "up to an eight-

figure dollar amount."

4 Revenue from contracts with customers, page F-44

We note your response to prior comment 5 and your revised disclosure. Please clarify in

the disclosure that the reimbursement is for 50% of Cellscript sublicense costs as you

stated in the response. Please also disclose that the Sub-sublicense Agreement is dated

December 22, 2018.

Prof. Ugur Sahin, M.D.

FirstName SE

BioNTech LastNameProf. Ugur Sahin, M.D.

Comapany 19, 2019

September NameBioNTech SE

Page 3

September 19, 2019 Page 3 FirstName LastName

You may contact Vanessa Robertson at (202) 551-3649 or Lisa Vanjoske at

3614 if you have questions regarding comments on the financial statements and related

matters. Please contact Tonya K. Aldave at (202) 551-3601 or Justin Dobbie, Legal Branch

Chief, at (202) 551-3469 with any other questions.

Sincerely,

Division of

Corporation Finance

Office of Healthcare &

Insurance

Eric Blanchard, Esq. cc: