



DIVISION OF  
CORPORATION FINANCE

UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
WASHINGTON, D.C. 20549

September 14, 2021

Branislav Vajdic  
Chief Executive Officer  
HeartBeam, Inc.  
2118 Walsh Avenue, Suite 210  
Santa Clara, CA 95050

**Re: HeartBeam, Inc.**  
**Registration Statement on Form S-1**  
**Filed September 7, 2021**  
**File No. 333-259358**

Dear Dr. Vajdic:

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 21, 2021 letter.

Registration Statement on Form S-1, Filed September 7, 2021

Dilution, page 35

1. We note your response to comment 4 and your revised disclosure in the dilution section. Please reconcile the statement that, "The amount of Common Stock outstanding excludes Common Stock issuable on a Qualified Financing such as this IPO, at a conversion price equal to seventy (70)% of the price per share in the public offering" and your disclosure that, "... and after the conversion of the Convertible Notes (and interest) through this Qualified Financing (as defined in the Convertible Notes), the Company's pro forma as adjusted net tangible book value as of [redacted], 2021 would have been \$ [redacted] or \$ [redacted] per share" or advise.

Branislav Vajdic  
HeartBeam, Inc.  
September 14, 2021  
Page 2

Business

Overview

FDA Regulatory Path, page 48

2. On page 48 you state: “We have finished an internal pilot study on 40 subjects and the results indicate that our synthesized 12-lead ECG is a better match to the standard 12-lead ECG recording than the FDA cleared synthesized 12-lead ECG device that is in the market.” Please revise to provide the details of this study.

You may contact Mary Mast at 202-551-361 or Kevin Kuhar at 202-551-3662 if you have questions regarding comments on the financial statements and related matters. Please contact Margaret Schwartz at 202-551-7153 or Tom Kluck at 202-551-3233 with any other questions.

Sincerely,

Division of Corporation Finance  
Office of Life Sciences

cc: Scott Linsky, Esq.