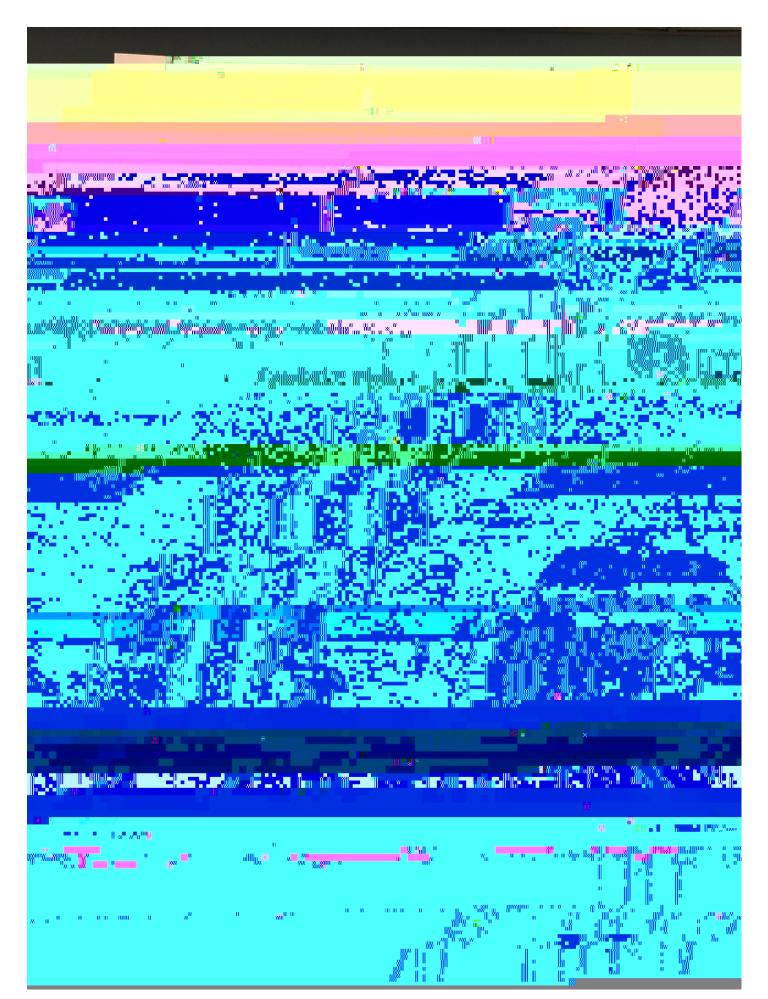
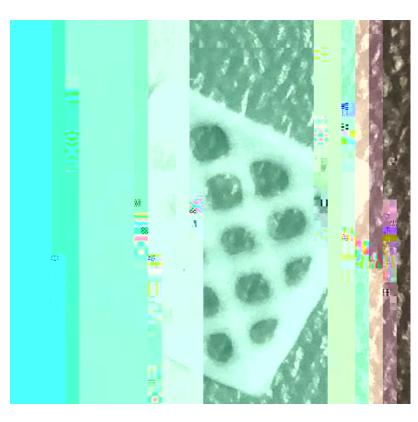
### Abstract

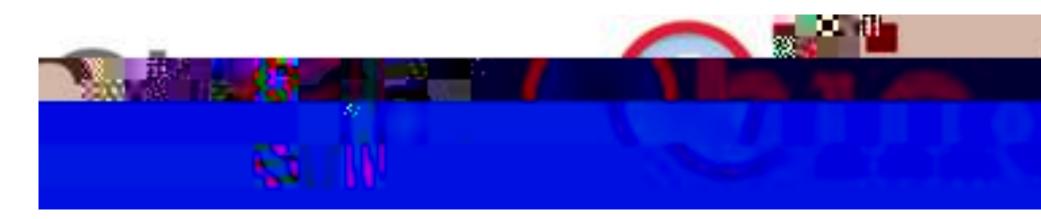
Myoblasts, Myogenic stem cells, are capable of differentiating into functioning contractile myocytes (muscle fibers). In skeletal muscle, myocytes are aligned in parallel arrays as fascicles. This allows for contraction in only one direction, all myocytes working together cooa [stTQ40 G[Mentor: Dr)54(. Gary )18(W 0 g0 G 0.06 Tc[. )]TETQ EMC /P & MCID 26 & 0.06 Tc[. ]]TETQ EMC /P



M1: 3D Printing System



M2: Printed tissue grid



# **3D Bio-Printing of Muscle Tissue**

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## Methods

Results



	Nucleic
Sample ID	Cone
3D 400 1.2	65.1
3D 400 2.2	4.6
3D 200 3.2	5.8
3D 200 4.2	4.4

2.5 0.01 RNA | ng/µl Figure 2: This is an analytical table showing the spectral data (Nanodrop®) verifying that cells survived the printing and subsequent culture phase of tissue construction.

ng/µl

ng/µl

To further confirm the viability of the cells embedded in the tissue future experiments will use the fluorescent DAN stain, DAPI, to confirm normal nuclear structure. Also, in future printing runs, the distribution and careful quantitation of each sample must be much more carefully controlled and recorded. Lastly, before quantitative PCR(qPCR) analysis, the contamination

quantitative Polymerase chain reactions can be performed using select myo-specific primers.

### Figure 1 Data

Figure 1: R1: These measurements were made using stage micrometer. What is indicated in the table is average width in millimeters of each image. The average bar width is 1.12mm with a standard deviation of .22mm

0.01 RNA

#### Acid Sample Unit Type IC. 1.73 0.13 RNA ng/µl 2.66 0.01 RNA

4.02

### Figure 2 RNA Data

### Discussion

