

A functional human motor unit platform engineered from human embryonic stem cells and immortalized skeletal myoblasts

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Background: Although considerable research on neuromuscular junctions (NMJs) has been conducted, the prospect of in vivo NMJ studies is limited and these studies are challenging to implement. Therefore, there is a clear unmet need to develop a feasible, robust, and physiologically relevant in vitro NMJ model.

Objective: We aimed to establish a novel functional human NMJs platform, which is serum and neural complex media/neural growth factor-free, using human immortalized myoblasts and human embryonic stem cells (hESCs)-derived neural progenitor cells (NPCs) that can be used to understand the mechanisms of NMJ development and degeneration.

Methods: Immortalized human myoblasts were co-cultured with hESCs derived committed NPCs. Over the course of the 7 days myoblasts differentiated into myotubes and NPCs differentiated into motor neurons.

Results: Neuronal axon sprouting branched to form multiple NMJ innervation sites along the myotubes and the myotubes showed extensive, spontaneous contractile activity. Choline acetyltransferase and β III-tubulin immunostaining confirmed that the NPCs had matured into cholinergic motor neurons. Postsynaptic site of NMJs was further characterized by staining dihydropyridine receptors, ryanodine receptors, and acetylcholine receptors by α -bungarotoxin.

Conclusion: We established a functional human motor unit platform for in vitro investigations. Thus, this co-culture system can be used as a novel platform for 1) drug discovery in the treatment of neuromuscular disorders, 2) deciphering vital features of NMJ formation, regulation, maintenance, and repair, and 3) exploring neuromuscular diseases, age-associated degeneration of the NMJ, muscle aging, and diabetic neuropathy and myopathy.

Keywords: motor unit, neuromuscular junctions, human embryonic stem cells, neuronal progenitor cells, human myoblasts

Introduction

Neuromuscular junctions (NMJs) serve as the interface between nerves and skeletal muscles. Maintenance, structure, and formation of NMJs depend on the bidirectional molecular interaction between the muscle and motor neuron.¹ The NMJ consists of a presynaptic motor neuron terminal, a postsynaptic motor end plate and a synaptic cleft. If chemical or molecular communication is disrupted, NMJ deterioration can follow. This involves axon degeneration, synapse disruption, impaired NMJ transmission, and muscle fiber degradation² which are the features of neuromuscular diseases, myopathies, and age-associated neuromuscular impairments.³

Despite decades of intensive research to characterize the structure and function of NMJs by utilizing animals and ex vivo models,⁴ effective treatment of neuromuscular

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and neurodegenerative diseases remains a significant unmet clinical need. This is mainly due to the failure of experimental animal models to reflect complex processes of human aging and disease progression.⁵ In order to advance this field, novel, alternative, experimental models are needed.

There has been recent progress toward the development of in vitro co-culture models using human induced pluripotent stem cells (iPSCs);^{6,7} mouse,⁸ rat,⁹ and human primary myoblasts;^{10,11} and human embryonic stem cells (hESCs)¹²⁻¹⁴ and cross-species models.^{15,16} However, existing in vitro motor neuron and skeletal muscle co-culture systems typically require a complex neural growth medium that contains serum and cocktails of around 15 neural growth factors (some of which are derived from animals).^{11,12,17} This further complicates drug discovery and toxicology studies due to possible cross-communication of the novel compound with factors contained within the added media, possibly explaining why many promising therapies do not translate to clinics. Another issue with existing models is that muscle contraction is induced by applied electrical or chemical stimulation, which does not replicate the native physiological stimulation required for muscle contractions.^{8,17-19} Recent innovation in the use of iPSCs offers the potential to derive myoblasts and motor neurons for use with in vitro NMJ models. However, cells derived from iPSCs may exhibit genetic inconsistency and genetic modification, which limit their use.²⁰ Recent human iPSC-based studies have failed to recapitulate the severe neuronal loss observed in human neurodegenerative diseases.²¹⁻²³ Human skeletal myoblasts which were used in some of the abovementioned models^{10,11} were obtained from primary cells (eg, muscle biopsy or surgical samples), but their life span is limited to just a few passages which restricts experimentation and necessitates repeated supply of the primary cells.^{24,25} Furthermore, primary cells have varied cell purity²⁶ and experience phenotypic changes when expanded, rendering primary myoblasts a problematic choice for a consistently reproducible co-culture system.^{24,25} Therefore, there is a clear need for a more relevant human experimental model to study motor units and NMJs to overcome the limitations of existing models.

Methods

Human immortalized myoblast cultures

The human immortalized myoblasts cell line ("C25") was obtained from the Institute of Myology.²⁷ This cell line was established using a biopsy of semi-tendinosis from a 25-year-old male (obtained anonymously from Myobank, a tissue bank affiliated to EuroBioBank which is authorized by the

French Ministry of Research [authorization AC-2013-1868]). After attaining 80% confluence, cells were seeded in six-well plates recoated with gelatin (0.5%) at a concentration of 1.5×10^5 cells/mL in growth media. The growth media was supplemented with DMEM from Lonza (Basel, Switzerland), 60% (v/v) Medium 199 with Earle's Balanced Salt Solution from Lonza, 20% (v/v) heat-inactivated FBS from Thermo Fisher Scientific (Waltham, MA, USA), 20% (v/v) L-glutamine from Lonza, 1% (v/v) fetuin from FBS from Sigma-Aldrich (St Louis, MO, USA) 25 μ g/mL, recombinant human basic fibroblast growth factor from Thermo Fisher Scientific 0.5 ng/mL, recombinant human EGF from Thermo Fisher Scientific 5 ng/mL, recombinant human hepatocyte growth factor from Sino Biological Inc. (Beijing, China) 2.5 ng/mL, recombinant human insulin from Sigma-Aldrich 5 μ g/mL, dexamethasone from Sigma-Aldrich 0.2 μ g/mL and gentamicin from Thermo Fisher Scientific.

Neural differentiation of hESCs

Induction of neuroepithelial clusters (NECs)

Mouse embryonic fibroblasts (MEFs; Cell Biolabs, San Diego, CA, USA) were cultured within MEF growth media (summarized in Table 1) and were passaged at a 1:4 ratio. At passage 4 (p4), MEFs were inactivated mitotically using 0.1 μ g/mL mitomycin C (Sigma-Aldrich). The Shef3 hESC line was obtained from the UK StemCell Bank under the project SCSC10-48 and maintained on mitotically inactivated MEFs

Table 1 MEF growth media

MEF growth media components	Volume
DMEM from Lonza (Basel, Switzerland)	500 mL
L-glutamine from Lonza	1% (v/v)
Penicillin/streptomycin (Sigma-Aldrich, St Louis, MO, USA)	5 mL (2 mM)
Heat-inactivated FBS from Thermo Fisher Scientific (Waltham, MA, USA)	10% (v/v)

Abbreviation: MEF, mouse embryonic fibroblast.

Table 2 hESC media for cells on MEFs

hESC/MEF cell medium components	Volume
DMEM-F12 (1:1) from Lonza (Basel, Switzerland)	38.5 mL
MEM nonessential amino acids (Thermo Fisher Scientific, Waltham, MA, USA)	0.5 mL (1X)
Penicillin/streptomycin (Sigma-Aldrich, St Louis, MO, USA)	0.5 mL (1X)
bFGF (R&D Systems, Minneapolis, MN, USA)	5 μ L (10 ng/mL)
Knockout serum replacement (Thermo Fisher Scientific)	10 mL (20%)

Abbreviations: bFGF, basic fibroblast growth factor; hESC, human embryonic stem cell; MEF, mouse embryonic fibroblast; MEM, Minimum Essential Medium.

in hESC medium (summarized in Table 2) in 96-well plates. hESCs were mechanically passaged every 5–7 days and then conditioned to feeder-free culture by TrypLE Express (Thermo Fisher Scientific) enzyme dissociation and plated at a high density (1:1) onto hESC-qualified Matrigel® (Corning)-coated flasks in mTESR (STEMCELL Technologies). For neural induction, feeder-free hESCs were dissociated with TrypLE and replated in neural induction medium (NIM; summarized in Table 3) in uncoated, V-shaped 96-well plates at a density of 1×10^4 cells/well. Within 24 hours, NECs formed.

Generation of neural rosette-forming progenitor cells (NRPCs)

Medium was replaced daily for 5 days. On day 6, aggregates were replated in 96-well plates in NIM onto 20 $\mu\text{g}/\text{mL}$ laminin (Millipore)-coated dishes to allow neural rosette formation (2–3 days).

Expansion of neural progenitor cells

Neural rosette clusters were mechanically isolated and replated in 96-well plates onto laminin-coated dishes in neural expansion medium (NIM plus 1X B27 supplement; Thermo Fisher Scientific). Early- and late-passage neural progenitor cells (NPCs) were cultured in the same conditions and passaged at a 1:3 ratio using TrypLE. To monitor NPCs differentiation into motor neurons, some of the NPCs were transfected with a GFP-reporter lentivirus system.

Immunohistochemistry of NECs, NRPCs, and NPCs

For immunostaining, at each stage of neural differentiation (NECs and NRPCs), cells were sectioned onto glass slides. Sectioned NECs and NRPCs and cultured NPCs in six-well plates were fixed with 4% paraformaldehyde. Fixed sections and NPCs were permeabilized with 0.3% Triton X in PBS and then subsequently blocked with 2% BSA in PBS/Tween

20. The slides were then incubated overnight at 4°C with a primary antibody (MAB2736; R&D Systems). The following day, sections and NPCs were washed with PBS and incubated for 1 hour at room temperature (RT) with goat anti-mouse secondary antibody (Alexa Flour 568-red for NECs, NRPCs and NPCs; Thermo Fisher Scientific) and imaged under a fluorescent microscope (Leica CTR 6000; Leica Microsystems, Wetzlar, Germany).

Co-culture of NPCs and human myoblasts

Human myoblasts were incubated over 24 hours at 37°C within a 5% CO₂ environment in six-well plates. Then, the growth media was replaced with co-culture media supplemented with DMEM (500 mL), 10 $\mu\text{g}/\text{mL}$ recombinant human insulin, and 10 $\mu\text{g}/\text{mL}$ gentamicin. NPCs were present at a concentration of 25×10^3 cells/mL and incubated at 37°C with 5% CO₂ for up to 7 days. At day 7, myotube contractions were observed.

Immunohistochemistry

At day 7, cells were stained with α -Bungarotoxin (α -BTX) (T0195, 1/200; Sigma-Aldrich), and then fixed with 4% paraformaldehyde for 10 minutes at RT. Co-cultures were washed twice with PBS (Thermo Fisher Scientific). After α -BTX staining, cells were washed twice with PBS and permeabilized with 1X perm/wash buffer (BD, Franklin Lakes, NJ, USA) for 30 minutes at RT. Cells were washed twice with PBS and then blocked with PBS containing 1% BSA and 10% goat serum (Thermo Fisher Scientific) for 1 hour at RT. The following antibodies were used: mouse anti- β III-tubulin (MAB1195, clone #TuJ1, 1/400; R&D Systems), rabbit anti-ryanodine receptor (anti-RyR; AB9078, 1/200; Millipore), goat anti-choline acetyltransferase (anti-ChAT; ABN100, 1/200; Millipore), goat anti-ChAT (AB144, 1/200; Millipore), and mouse anti-dihydropyridine receptor (anti-DHPR; Ab2864, 1/400; Abcam). These antibodies were incubated overnight at 4°C in 1X perm/wash buffer. Co-cultures were washed with PBS and stained with the corresponding secondary antibodies supplemented with DAPI (1/10,000; Sigma-Aldrich) for 1 hour at RT. Stained co-culture cells were visualized using a Leica SP5 confocal microscope sourced by Leica Microsystems for fluorescent microscopy.

Results

A functional human motor unit platform

The present work describes a functional human motor unit platform established using immortalized skeletal myoblasts

Table 3 Neural induction medium

NIM components	Volume
DMEM-F12 (1:1) from Lonza (Basel, Switzerland)	48.5 mL
MEM nonessential amino acids (Thermo Fisher Scientific, Waltham, MA, USA)	0.5 mL (1X)
Penicillin/streptomycin (Sigma-Aldrich, St Louis, MO, USA)	0.5 mL (1X)
bFGF (R&D Systems, Minneapolis, MN, USA) (100 $\mu\text{g}/\text{mL}$)	10 μL (20 ng/mL)
N2 supplement (Thermo Fisher Scientific)	0.5 mL (1X)
Heparin (Sigma-Aldrich) (2 mg/mL)	50 μL (2 $\mu\text{g}/\text{mL}$)
Poly (vinyl alcohol) (Sigma-Aldrich)	4 mg/mL

Abbreviations: bFGF, basic fibroblast growth factor; MEM, Minimum Essential Medium; NIM, neural induction medium.

and hESCs-derived NPCs that develop into motor neurons in muscle differentiation media (co-culture media) devoid of any complex neural growth factors or serum.

To validate the platform, two monoculture controls were included, one of NPCs and the other of immortalized human myoblasts²⁷ with co-culture media for 7 days. Cultured GFP-transfected NPCs did not show any morphological changes or motor neuron differentiation, but instead they deteriorated and died (Figure 1A). The human myoblasts stained with phalloidin-DAPI showed normal morphology and differentiation features with centrally and peripherally located nuclei, but the myotubes did not spontaneously contract (Figure 1B).

Derivation of NPCs from hESCs and establishment of the NMJs model

The NPCs were derived from hESCs, as described previously.²⁸ Neural differentiation of hESCs progressed through three stages of differentiation to NECs (Figure 2A) and NRPCs (Figure 2B) which were stained with Nestin red color. To confirm that differentiated NPCs were homogeneous committed neural lineage, cells were stained with DAPI, anti-GFAP (a specific marker for glial cells) and anti-Nestin. Double-positive staining (DAPI blue and Nestin red) confirmed the formation of NPCs while single DAPI staining was not observed, which confirmed that the differentiated cells were NPCs (Figure 2C). Staining of GFAP was not present, confirming again that all of the cells were NPCs (Figure 2C).

The NPCs were co-cultured with human immortalized myoblasts for 7 days in co-culture media. After myogenic differentiation was initiated, the characteristics of functional motor units began to develop. This included myotube formation and axonal sprouting from the NPCs which subsequently formed NMJs along the myotubes (Figure 2D). The myotubes

showed spontaneous muscle contractions from approximately day 7 onwards in the absence of any exogenous electrical and chemical stimuli (Video S1). Due to the force of contractions, some myotubes were detached from the culture plate, causing large spaces within the co-culture. Since the myotubes cultured without NPCs did not contract, they did not detach from the culture plate and did not open up large spaces (Figure 1B).

Characterization of co-cultures

On day 7, the motor neuron formation was assessed using the specific marker for motor neuron differentiation, β III-tubulin. Figure 3A shows a typical shape of mature motor neurons with axons terminating on myotubes. To further confirm the formation of cholinergic motor neurons, acetyltransferase antibodies (ChAT, a key enzyme for acetylcholine biosynthesis) were used and are shown in green in Figure 3B.

Neurally and aneurally cultured myotubes were characterized by antibody staining against DHPRs and RyRs voltage-gated channels which are located at muscle fiber T-tubules and the sarcoplasmic reticulum, respectively. The DHPR- and RYR-stained images showed transversal triad structures (Figure 3C). These images illustrated mature differentiated myotubes within the co-culture platform with peripherally located nuclei (Figure 3A and B). The aneurally cultured myotubes (control image) differentiated as expected, but did not exhibit appropriate transversal triad structures (Figure 3D).

NMJs display acetylcholine receptor (AChR) clusters along myotubes, and these were assessed by staining for α -BTX, as shown in Figure 3B and E where AChRs are marked by red clusters. As shown in Figure 3A and E, motor neuron axons extend to innervate myotubes and AChR clusters at this same location, marked with α -BTX (red),

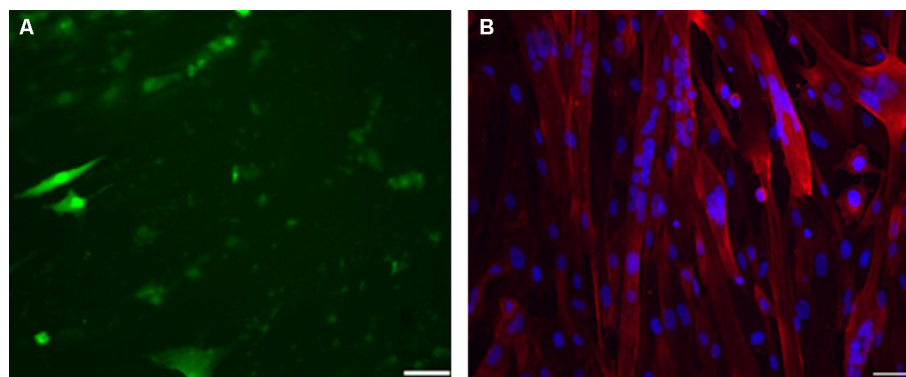


Figure 1 Monoculture of NPCs (A) and myoblasts (B).

Notes: NPCs and myoblasts were separately cultured for 7 days. The same growth media was used for monocultures and co-cultures. (A) A representative image of GFP-transfected NPCs. The NPCs did not exhibit any morphological changes or motor neuron differentiation; instead, they deteriorated and died. (B) A representative image of phalloidin-DAPI-stained human myoblasts showed normal morphology and differentiation features with centrally and peripherally located nuclei. Scale bar: 50 μ M.

Abbreviation: NPC, neural progenitor cell.

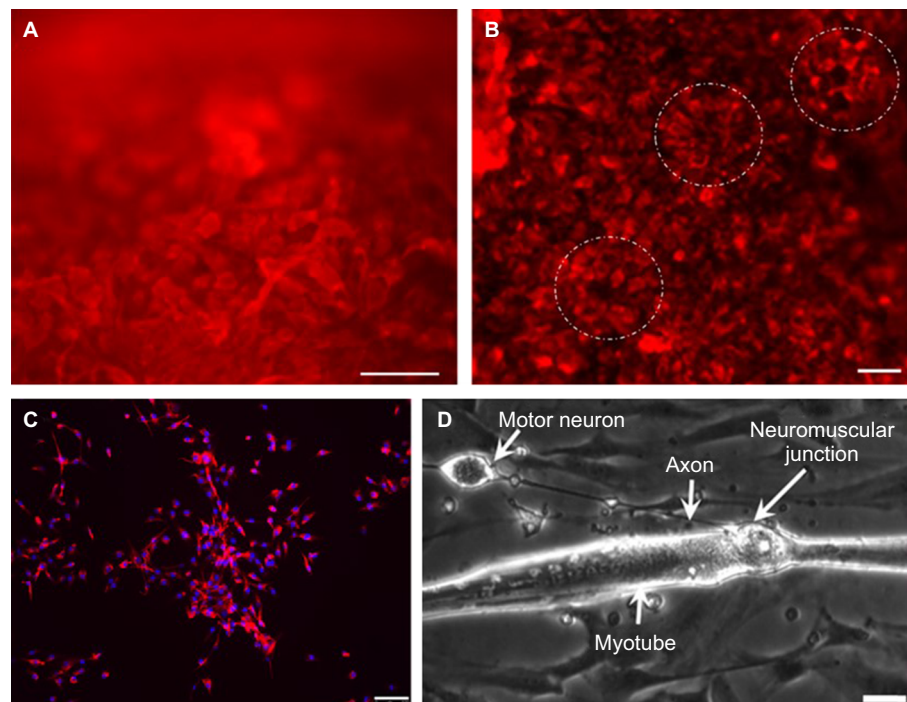


Figure 2 Neural differentiation of hESCs.

Notes: Identification of NPCs from hESCs using Nestin red color as a specific neural differentiation marker. (A) NECs. (B) NRPCs. (C) Nestin-red color for NPCs. (D) A representative phase-contrast image of NMJ formation showing an axon extended to contact a myotube. White dotted circle indicates a rosette shape. Scale bars are 100 μ M for images (A)–(C) and 25 μ M for image (D).

Abbreviations: hESC, human embryonic stem cell; NEC, neuroepithelial cluster; NMJ, neuromuscular junction; NPC, neural progenitor cell; NRPC, neural rosette-forming progenitor cell.

indicating NMJ formation. Spontaneous myotube contractile activity was determined on day 7. The myotubes co-cultured with motor neurons showed high levels of spontaneous contractile activity (Video S1). Muscle contractions were absent from aneural myoblast cultures.

Discussion

Implications and future uses

This report describes a novel functional human motor unit platform engineered from immortalized skeletal myoblasts and NPCs derived from hESCs. This unique platform was established for investigation of human NMJs and motor unit formation, maintenance, and disease, as well as for drug discovery and toxicity research.

The motor neuron and skeletal muscle co-cultures matured without adding complex cocktails of serum or neural growth factors. NMJ formation was observed and spontaneous myotube contractions were recorded (Video S1) over a period as short as 6–7 days, which is very early compared with the 14 days needed in previous studies¹⁶ or where NMJs formed after 20–25 days.²⁹ These are crucial advances for studying the basic aspects of development and for recognizing pathophysiological systems of NMJ disorders associated with disease

or aging. It is important that this type of work is carried out with relevant human cultures to increase the possibility for translation to clinical practice. Most previous cell culture model systems used cross-species cell types^{15,19} and required cocktails of serum^{30,31} and growth factors.^{10,17,19,29} These were absent from the present model without any adverse effects, which suggests that nerve and muscle cells release all of the necessary factors needed to stimulate nerve axonal sprouting and formation of NMJs with myotubes. Moreover, eliminating serum, which contains unknown factors that may affect assay reproducibility, simplifies the interpretation of pharmacology and toxicity studies.³² Functional NMJ formation is strongly supported by bidirectional communication¹ between nerve and muscle as well as by neural growth factors (such as brain-derived neurotrophic factor, glial cell line-derived neurotrophic factor and neurotrophin-3/4) secreted by muscle to support NMJ formation, maturation and maintenance.^{33,34} Future investigations using this model will identify key factors released from nerve and muscle to orchestrate axonal sprouting, localization, and NMJ maintenance.

Although animal models can represent essential parts of physiological changes in a human disease,³⁵ *in vitro* human cell cultures offer many advantages because they are formed

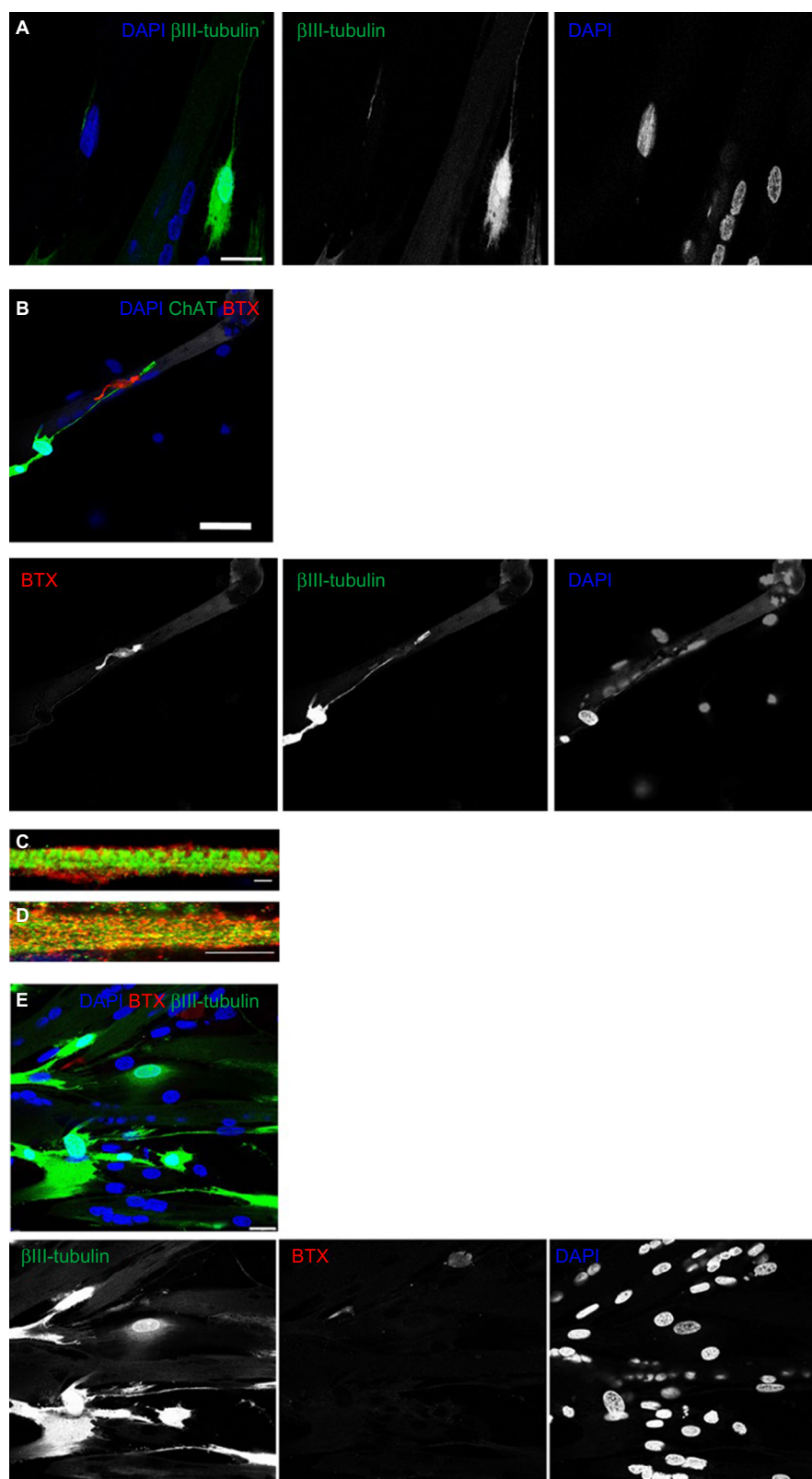


Figure 3 Characterization of muscle–nerve co-cultures.

Notes: (A and B) Characterization of motor neuron formation. (A) A representative image of the co-culture cells stained for βIII-tubulin (green) and DAPI (blue); scale bars: 25 μm. (B) A representative image of the co-culture cells stained for ChAT (green), α-BTX (red), and DAPI (blue); scale bars: 75 μm. (C and D) Characterization of advanced differentiated myotubes. (C) A representative image of the co-culture cells stained for DHPR (red), RyR (green), and DAPI (blue); scale bars: 2.5 μm. (D) A representative image of the co-culture cells stained for DHPR (red), RyR (green), and DAPI (blue); scale bars: 7.5 μm. (E) Characterization of functional NMJs formation. A representative image of the co-culture cells stained for βIII-tubulin (Tuj1; green), α-BTX (red), and DAPI (blue); scale bars: 25 μm.

Abbreviations: α-BTX, α-bungarotoxin; ChAT, choline acetyltransferase; DHPR, dihydropyridine receptor; NMJ, neuromuscular junction; RyR, ryanodine receptor.

of relevant cell types, they can be produced quickly for high-throughput screening, and they are more cost-effective in comparison to animal models.^{10,17}

Conclusion

In summary, human immortalized myoblasts were co-cultured with hESCs-derived NPCs. Over the course of 7 days, myoblasts differentiated into myotubes and NPCs sprouted axons that branched to form multiple NMJ innervation sites along myotubes, and myotubes showed extensive, spontaneous contractile activity. This cell culture platform may be used to study human NMJ growth and disease and may reduce the use of animal models in future related research.

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Author contributions

MAS, JSM, and NAS developed the study concept. All the authors contributed to methodology. MAS, JSM, and NAS carried out the investigation. JSM, TRM, and NAS supervised the study. MAS, JSM, and NAS wrote the original draft of the manuscript. MAS, JSM, VM, and NAS reviewed and edited the draft manuscript. All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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Supplementary material

Video S1 Phase contrast video micrograph of innervated human myotube contractions at Day 7.

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