

Hippo/YAP-mediated rigidity-dependent motor neuron differentiation of human pluripotent stem cells

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Our understanding of the intrinsic mechanosensitive properties of human pluripotent stem cells (hPSCs), in particular the effects that the physical microenvironment has on their differentiation, remains elusive¹. Here, we show that neural induction and caudalization of hPSCs can be accelerated by using a synthetic microengineered substrate system consisting of poly(dimethylsiloxane) micropost arrays (PMAs) with tunable mechanical rigidities. The purity and yield of functional motor neurons derived from hPSCs within 23 days of culture using soft PMAs were improved more than fourfold and tenfold, respectively, compared with coverslips or rigid PMAs. Mechanistic studies revealed a multi-targeted mechano-transductive process involving Smad phosphorylation and nucleocytoplasmic shuttling, regulated by rigidity-dependent Hippo/YAP activities and actomyosin cytoskeleton integrity and contractility. Our findings suggest that substrate rigidity is an important biophysical cue influencing neural induction and subtype specification, and that microengineered substrates can thus serve as a promising platform for large-scale culture of hPSCs.

Human pluripotent stem cells including human embryonic stem cells (hESCs) and induced pluripotent stem cells (hiPSCs) are a promising resource for regenerating complex neural tissues, including motor neurons (MNs; refs 2,3). However, poorly defined culture conditions and inefficient protocols for derivation of MNs from hPSCs have hindered their use. Current hPSC-based MN differentiation relies on soluble morphogens including sonic hedgehog (SHH) and retinoic acid (RA) to direct MN specification. Because morphogenesis during embryonic development occurs through dynamic modulation of extracellular physical signals, matrix rigidity is likely to be important for the differentiation and functional maturation of neural subtypes from hPSCs. Here we investigated intrinsic mechanosensitive properties of hPSCs, and determined how they could be leveraged to improve production of functional MNs using a synthetic, micromoulded poly(dimethylsiloxane) (PDMS) micropost array (PMA) system (Supplementary Fig. 1 and Methods), which has a uniform surface geometry and different post heights to modulate substrate rigidity independent of effects on adhesion and other material surface properties⁴.

The influence of substrate rigidity on neural induction of hPSCs was assessed using vitronectin-coated coverslips (with bulk modulus $E = 10^4$ kPa) and PMAs with a broad range of rigidities ($E = 1.0$ – $1,200$ kPa). hESCs were seeded at $20,000$ cells cm^{-2} on coverslips and PMAs in growth medium. After 24 h, hESCs were switched to neural induction medium containing the dual Smad inhibitors, SB 431542 (SB, TGF- β inhibitor) and LDN 193189 (LDN, BMP4 inhibitor), to promote neural induction (Fig. 1a; ref. 3). No significant difference in cell attachment was observed between coverslips and PMAs of different rigidities (not shown). On coverslips and PMAs with $E = 1,200$ kPa (rigid PMAs), hESCs spread to form monolayers, whereas on PMAs with $E \leq 5.0$ kPa (soft PMAs), hESCs spontaneously migrated towards each other to form compact clusters. Cell spreading and nucleus size of hESCs were significantly reduced on soft PMAs relative to on coverslips or rigid PMAs (referred to henceforth as controls; Supplementary Fig. 2a–c). Notably, within 24 h, $22.3 \pm 6.2\%$ of hESCs on soft PMAs had differentiated, indicated by loss of Oct4 (pluripotency-associated transcription factor) expression, whereas only $4.9 \pm 1.0\%$ (coverslip) and $3.9 \pm 0.4\%$ (rigid PMA) of hESCs on controls started differentiating (Supplementary Fig. 2d).

Expression of the early neuroectodermal differentiation marker, Pax6, was used to monitor neural induction. On soft PMAs, Pax6⁺ neuroepithelial cells (NEs) were detected as early as day 4, and reached $95.1 \pm 2.1\%$ by day 8. In contrast, on controls, Pax6⁺ NEs appeared at day 6 and constituted only $28.2 \pm 10.8\%$ (coverslip) and $33.4 \pm 7.2\%$ (rigid PMA) of total cells at day 8, respectively (Fig. 1b,c; ref. 5). Immunoblots revealed higher Pax6 and Sox1 (neuroectodermal transcription factor) protein expression by hESCs on soft PMAs relative to controls (Fig. 1d). Paralleling definitive neural stem cells in mouse embryos⁶, Pax6⁺ NEs derived from soft PMAs were responsive to bFGF and readily formed polarized neural tube-like rosettes, whereas the controls did not (Supplementary Fig. 3). A screening assay revealed a threshold of PMA rigidity E for neural induction: $E \leq 5$ kPa had a potent effect, whereas $E \geq 14$ kPa did not (Supplementary Fig. 4).

Two sets of PMAs with different post diameters but matching effective moduli E were compared, with results indicating that neural induction by soft PMAs was not sensitive to micropost geometries (Supplementary Fig. 5). By modulating the PDMS

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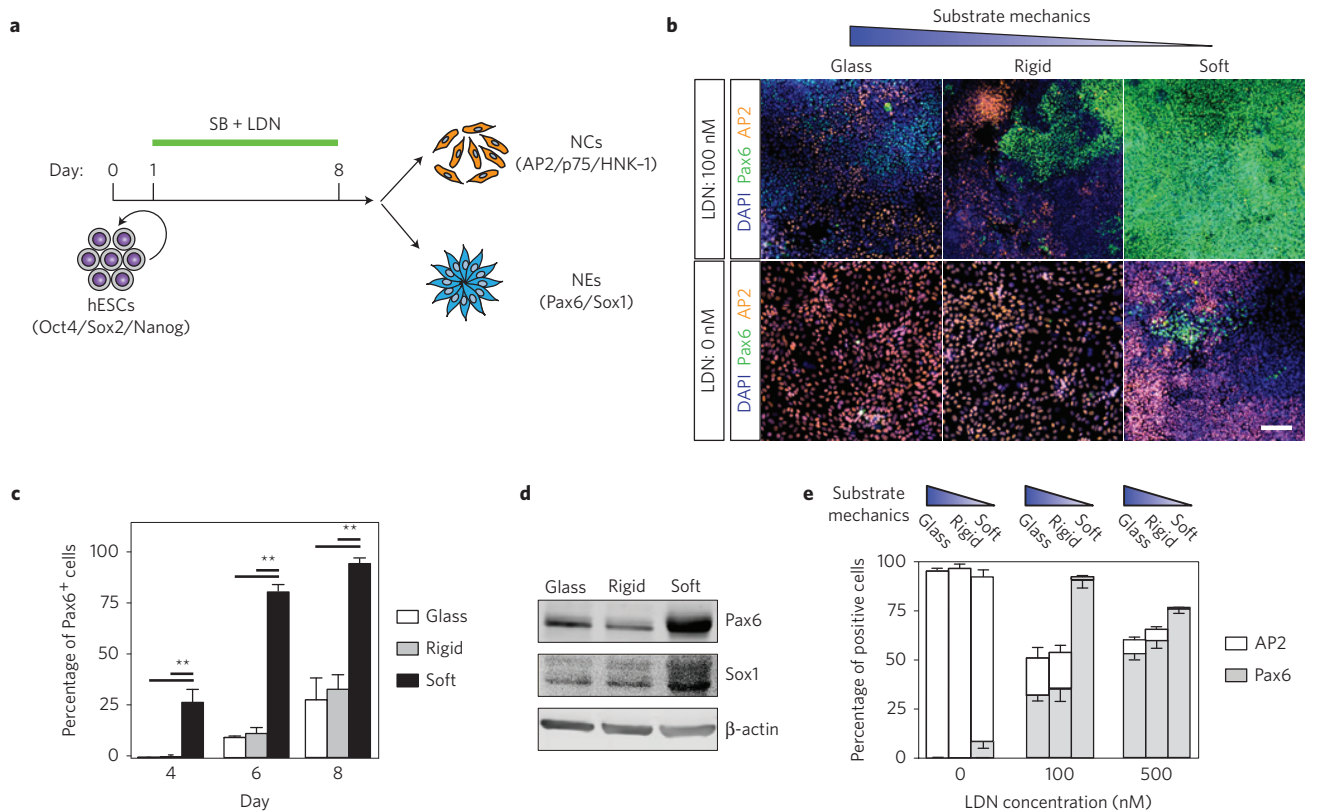


Figure 1 | Soft substrates promote neuroepithelial conversion while inhibiting neural crest differentiation of hESCs in a BMP4-dependent manner.

a, Schematic diagram showing experimental design of hESC neural induction. hESCs were cultured for 8 days in neural induction medium containing the dual Smad inhibitors, SB 431542 (SB, 10 μ M) and LDN 193189 (LDN, 100 nM). **b**, Representative immunofluorescence images showing Pax6⁺ NEs and AP2⁺ NCs after 8 days of culture with (top; 100 nM) or without (bottom) LDN on vitronectin-coated coverslips and rigid ($E=1,200$ kPa) and soft ($E=5$ kPa) PMAs. Scale bar, 100 μ m. **c**, Bar graph showing percentage of Pax6⁺ NEs on coverslips and rigid and soft PMAs at day 4, 6 and 8. **d**, Western blotting showing Pax6 and Sox1 expression levels in hESCs cultured for 8 d on coverslips and rigid and soft PMAs. **e**, Bar plot showing percentages of Pax6⁺ NEs and AP2⁺ NCs at day 8 as a function of substrate rigidity and LDN concentration. Data represents the mean \pm s.e.m with $n=3$. P values were calculated using one-way ANOVA, followed by Tukey post hoc analysis. **, $P < 0.01$.

curing agent to base monomer ratio (1:10–1:100), flat featureless PDMS surfaces with different bulk moduli E (0.5 kPa–2.5 MPa) were generated and assayed for neural induction. At day 6, Pax6⁺ NEs constituted $32.5 \pm 3.3\%$ (1:70 PDMS, $E = 5$ kPa) and $27.7 \pm 4.9\%$ (1:100 PDMS, $E = 0.5$ kPa) of cells on soft, flat PDMS surfaces, respectively, in contrast to $14.3 \pm 2.5\%$ (coverslip) and $14.3 \pm 1.8\%$ (1:10 PDMS, $E = 2.5$ MPa) on controls (Supplementary Fig. 6). Noticeably, neural induction by soft, flat PDMS surfaces was not as drastic as soft PMAs, possibly owing to their viscoelastic nature, different from the elastic PMAs.

Temporal mRNA expression of pluripotency markers (*NANOG* and *SOX2*) and neuroectodermal markers (*PAX6*, *SOX1*, and *TUJ-1*) was quantified using qRT-PCR. Soft PMAs accelerated disruption of the transcriptional circuitry maintaining pluripotency, while promoting neuroectodermal gene expression (Supplementary Fig. 7). Experiments with another hESC line (CHB-10) and a hiPSC line derived from human foreskin fibroblasts showed that at day 6, Pax6⁺ NEs constituted $60.8 \pm 1.1\%$ (CHB-10) and $78.8 \pm 5.8\%$ (hiPSCs) of cells on soft PMAs, respectively, in contrast to $23.2 \pm 3.0\%$ (CHB-10) and $24.5 \pm 1.8\%$ (hiPSCs) on rigid PMAs (Supplementary Fig. 8).

Treatment of hESCs with both Smad inhibitors led to Pax6⁺ NEs, as well as Pax6⁻ cells co-expressing neural crest (NC) markers AP2, p75, and HNK-1 (Fig. 1b and Supplementary Fig. 9; ref. 3). At day 8, $18.7 \pm 5.3\%$ and $18.1 \pm 3.6\%$ of cells were AP2⁺ on coverslips and rigid PMAs, respectively. Strikingly, only $1.1 \pm 0.5\%$ of cells were AP2⁺ on soft PMAs (Fig. 1b). Similar reductions in p75⁺

and HNK-1⁺ NCs were observed on soft PMAs (Supplementary Fig. 9). Paralleling the effects of substrate rigidity, manipulation of the concentration of the BMP4 inhibitor, LDN, altered the ratio of Pax6⁺ NEs to Pax6⁻/AP2⁺ NCs. Increasing LDN levels promoted differentiation towards Pax6⁺ NEs, whereas LDN's absence favoured NC differentiation (Fig. 1e).

To induce functional neurons, NEs require anterior/posterior and dorsal/ventral patterning cues^{2,7}. When the caudalization factor, RA, was added to neural induction medium from day 4 for rigid PMAs, the posterior genes, *HOXB1*, *HOXB4*, and *HOXC8* were significantly upregulated, whereas *OTX2*, an anterior identity gene, was suppressed (Supplementary Fig. 10a,b). Surprisingly, soft PMAs promoted *HOXB1*, *HOXB4*, and *HOXC8* expression in the absence of RA (Supplementary Fig. 10b). Although RA's effects exceeded those of soft PMAs on *OTX2*, *HOXB1*, or *HOXB4* expression, expression of *HOXB4* and *HOXC8* remained responsive to varying substrate rigidity in the presence of RA, suggesting an additive effect of substrate rigidity on neural caudalization. *Otx2* and *HoxB4* protein expression levels confirmed the effect of soft PMAs on promoting neural patterning and specification of posterior identity (Supplementary Fig. 10c,d).

We next examined whether the ventralization factor, Purmorphamine (Pur), could induce MNs from caudalized NEs derived from soft PMAs, using oligodendrocyte transcription factor 2 (*Olig2*) as a MN marker. When NEs were cultured continuously for an additional 8 d in the presence of Pur and RA, $65 \pm 6.3\%$ of cells on soft PMAs became *Olig2*⁺, whereas only

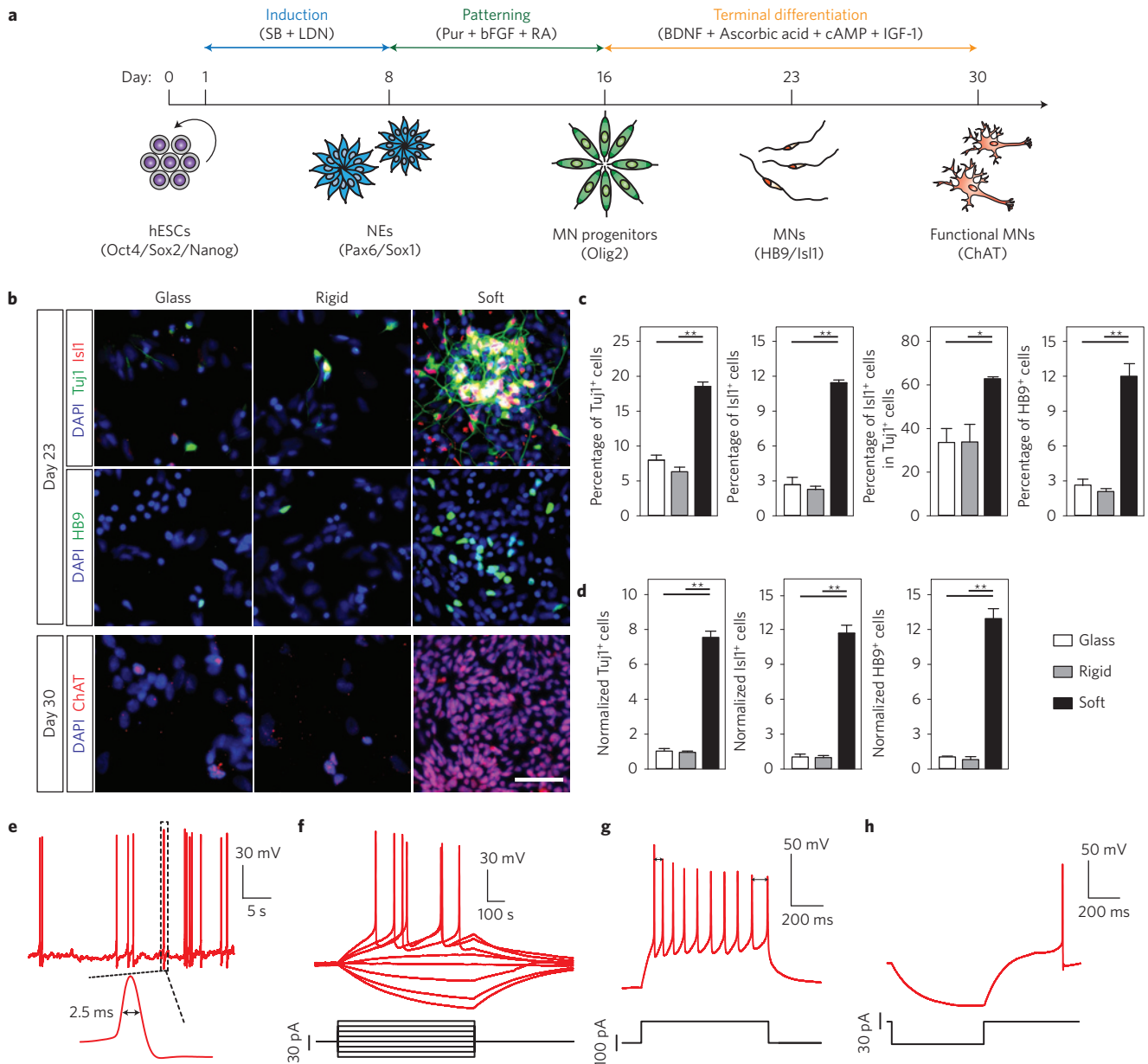


Figure 2 | Purity and yield of functional motor neurons (MNs) derived from hESCs are improved on soft substrates. **a**, Schematic diagram showing experimental design for sequential neural induction, patterning and functional maturation of MNs from hESCs. hESCs were cultured on vitronectin-coated coverslips and rigid ($E=1,200$ kPa) and soft ($E=5$ kPa) PMAs in neural induction medium containing the dual Smad inhibitors SB and LDN for 8 days and then in MN differentiation medium containing purmorphamine (Pur), basic fibroblast growth factor (bFGF) and retinoic acid (RA) for an additional 8 days. Putative MN progenitor cells collected at day 16 were transferred onto coverslips and cultured in MN maturation medium containing brain-derived neurotrophic factor (BDNF), ascorbic acid, cyclic adenosine monophosphate (cAMP) and insulin-like growth factor 1 (IGF-1) for another 14 days. **b**, Representative immunofluorescence images showing Tuj1⁺, Isl1⁺ and HB9⁺ cells at day 23 (top), and ChAT⁺ cells at day 30 (bottom). Scale bar, 100 μ m. **c,d**, Bar plots showing percentages (**c**) and relative numbers (**d**) of Tuj1⁺, Isl1⁺ and HB9⁺ cells at day 23 as a function of substrate rigidity. Data in **d** was normalized to values from coverslips. Data represents the mean \pm s.e.m with $n \geq 3$. P values were calculated using two-side unpaired student t -tests. *, $P < 0.05$; **, $P < 0.01$. **e-h**, Electrophysiological characterization of functional MNs derived from soft PMAs at day 30 using whole-cell patch clamp. **e**, Spontaneous action potential (AP) with resting membrane potential of -64.5 mV. **f**, Voltage response to current step injection. **g**, Instantaneous frequency change or spike frequency adaptation (SFA) evoked with positive current injection. **h**, Post-inhibitory rebound (PIR) after hyperpolarizing current injection.

$7.9 \pm 1.3\%$ of cells were Olig2⁺ on rigid PMAs (Supplementary Fig. 11). In the absence of RA, $19.3 \pm 1.7\%$ of cells on soft PMAs became Olig2⁺, whereas on rigid PMAs only $3.5 \pm 1.6\%$ of cells were Olig2⁺.

Because soft PMAs accelerate neural induction and caudalization, we investigated whether functional MN production could be expedited using soft PMAs. hESCs were cultured

on vitronectin-coated coverslips and PMAs for 16 d to allow neural induction and caudalization before passage onto poly-l-ornithine/laminin/fibronectin-coated coverslips and treatment with MN maturation medium containing neurotrophic factors BDNF and IGF-1 for an additional 14 d (Fig. 2a). At day 23, $18.5 \pm 0.6\%$, $11.4 \pm 0.3\%$, and $12.0 \pm 1.1\%$ of cells derived from soft PMAs became Tuj1⁺ (neuronal form of tubulin), Isl1⁺

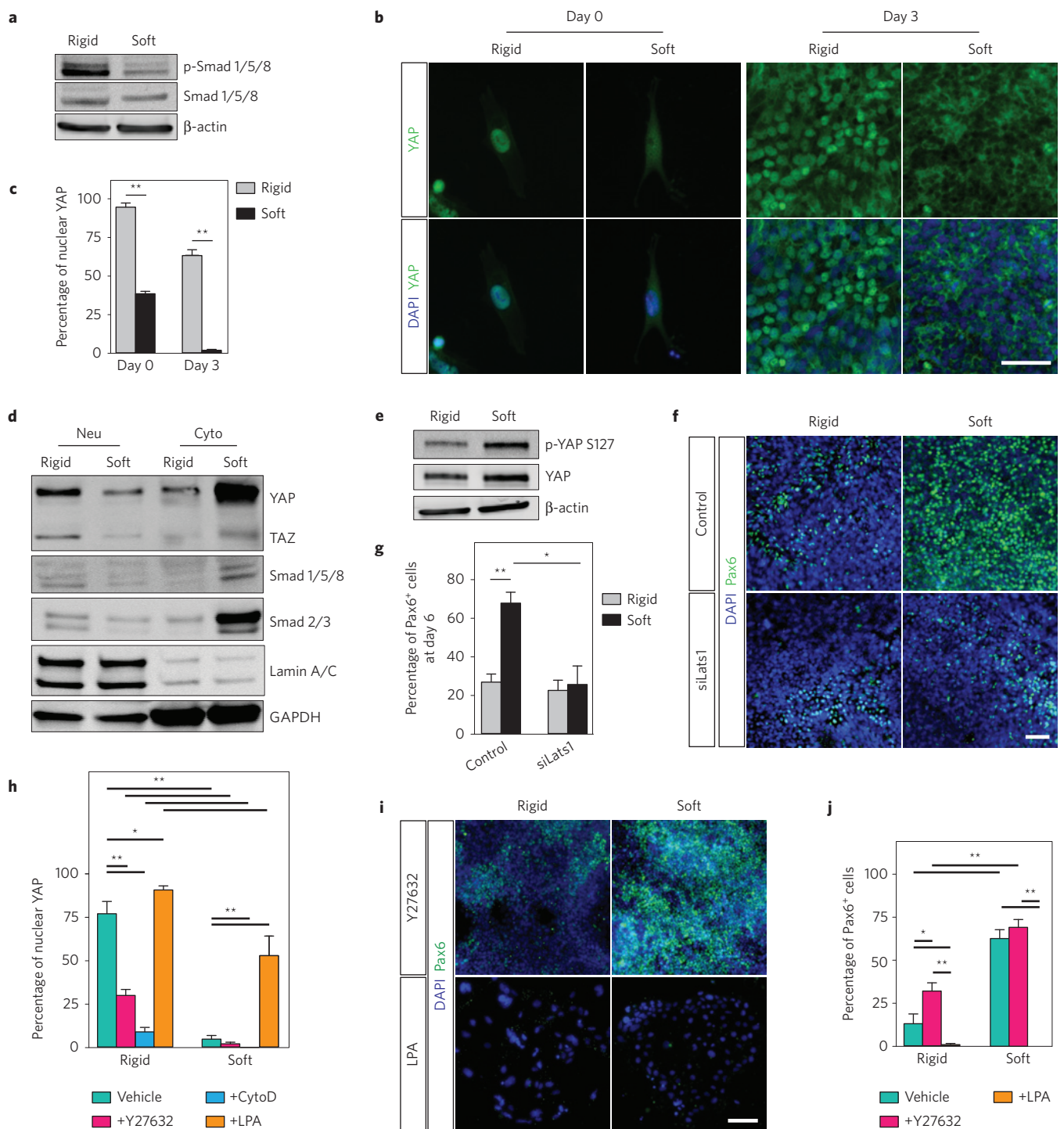


Figure 3 | Soft substrates promote hESC neuroepithelial conversion through a multi-targeted mechanotransductive process involving mechanosensitive Smad phosphorylation and nucleocytoplasmic shuttling regulated by rigidity-dependent Hippo/YAP activities and the actomyosin cytoskeleton.

a, Western blotting of total and phosphorylated Smad 1/5/8 (p-Smad 1/5/8) in hESCs cultured in neural induction medium for 3 days on rigid ($E = 1,200$ kPa) and soft ($E = 5$ kPa) PMAs. **b,c**, Representative immunofluorescence images (**b**) and bar plot (**c**) showing rigidity-dependent subcellular localization of YAP in hESCs at day 0 and day 3. Scale bar in **b**, 50 μm . **d**, Western blotting for YAP, TAZ, Smad 1/5/8 and Smad 2/3 in nuclear and cytoplasmic protein fractions from hESCs cultured for 3 days on rigid and soft PMAs. **e**, Western blotting for phosphorylated YAP S127 (p-YAP S127) and YAP in whole cell lysates of hESCs after 3 days of culture on rigid and soft PMAs. **f,g**, Representative immunofluorescence images (**f**) and bar plot (**g**) showing the percentage of Pax6⁺ NEs derived from scramble control and *siLats1* knockdown hESCs after 6 days of culture on rigid and soft PMAs. Scale bar in **f**, 50 μm . **h**, Bar graph showing the percentage of hESCs with nuclear localization of YAP after 3 days of culture with different drug supplementations, as indicated. **i,j**, Representative immunofluorescence images (**i**) and bar plot (**j**) showing the percentage of Pax6⁺ NEs derived from hESCs on rigid and soft PMAs after 6 days of culture with different drug supplementations, as indicated. Scale bar in **i**, 50 μm . In Western blots β -actin and GAPDH were used as protein loading controls and lamin A/C for nuclear fraction control. Data represents the mean \pm s.e.m with $n \geq 3$. *P* values in **c,g** and **h,j** were calculated using two-side unpaired student *t*-tests and one way ANOVA followed by Tukey post hoc analysis, respectively. *, $P < 0.05$; **, $P < 0.01$.

(MN-associated transcription factor), and HB9⁺ (MN-specific transcription factor), respectively. In contrast, Tuj1⁺, Isl1⁺, and HB9⁺ cells derived from coverslips constituted $8.0 \pm 0.7\%$, $2.6 \pm 0.6\%$, and $2.6 \pm 0.5\%$ of total cells, respectively (Fig. 2b,c). Similarly, only $6.3 \pm 0.7\%$, $2.2 \pm 0.3\%$, and $2.0 \pm 0.2\%$ of cells derived from rigid PMAs were Tuj1⁺, Isl1⁺, and HB9⁺, respectively (Fig. 2b,c). $62.6 \pm 1.1\%$ of Tuj1⁺ cells derived from soft PMAs co-expressed Isl1, whereas on controls only $33.3 \pm 6.2\%$ (coverslip) and $33.5 \pm 8.1\%$ (rigid PMA) of Tuj1⁺ cells were Isl1⁺ (Fig. 2c). Notably, significantly more soft PMA-derived cells expressed choline acetyltransferase (ChAT), an enzyme catalysing the formation of neurotransmitter acetylcholine, at day 30, compared with controls (Fig. 2b). The purity of MNs, defined as the percentage of HB9⁺ cells, from soft PMAs was thus improved about 4.6-fold, compared with controls (Fig. 2c). Furthermore, subculture of MN progenitors derived from soft PMAs led to 7.5-, 11.7-, and 12.9-fold increases in the numbers (and thus yields) of Tuj1⁺, Isl1⁺, and HB9⁺ cells, respectively (Fig. 2d).

We next evaluated action potentials (APs) for functional MNs derived at day 30 from soft PMAs using whole-cell recording (Supplementary Methods). Among 27 cells recorded, 12 generated APs with an average amplitude >65 mV and were considered functionally mature⁸. Another five cells fired APs with the average amplitude <65 mV. Nine exhibited spontaneous APs with an average amplitude of 85.3 mV and a half-height width of 2.5 ms (Fig. 2e). When injected with depolarizing currents, 15 cells exhibited firing activities with an average first spike amplitude of 71.7 mV (Fig. 2f). MNs can decrease firing rates responding to constant stimuli, a phenomenon called spike frequency adaption (SFA; ref. 9). Here, 6 of 12 'mature' MNs exhibited SFA with a depolarizing current (Fig. 2g). Moreover, post-inhibitory rebound (PIR, depolarization and/or spiking following a release from hyperpolarization) was observed in six MNs (Fig. 2h). These data suggest that MNs derived from soft PMAs exhibit electrophysiological activities comparable to those from primary neurons *in vivo*⁸. In comparison, hPSCs require up to 50 days under conventional differentiation conditions for complete functional maturation into a MN (refs 10,11).

To examine whether rigidity-mediated MN production might depend on specific ventralizing and neurotrophic factors, additional 23-day and 35-day MN differentiation assays were conducted using SHH for patterning and ventralization, and neurotrophic factors BDNF, GDNF, CNTF, and IGF-1 for terminal differentiation (Supplementary Figs 12 and 13). For the 23-day assay, $32.8 \pm 5.7\%$ and $17.7 \pm 1.7\%$ of cells derived from soft PMAs became Tuj1⁺ and HB9⁺, respectively, at day 23 (Supplementary Fig. 12b,c). In contrast, for controls, $14.6 \pm 1.3\%$ (coverslip) and $10.3 \pm 2.5\%$ (rigid PMA) of cells were Tuj1⁺, and $3.8 \pm 0.4\%$ (coverslip) and $3.0 \pm 0.8\%$ (rigid PMA) cells were HB9⁺ (Supplementary Fig. 12b,c). The purity and total number (and thus yield) of HB9⁺ MNs derived from soft PMAs were improved 4.6- and 12.9-fold, respectively, compared with controls (Supplementary Fig. 12c,d). Similarly, in the 35-day assay, the percentage of HB9⁺ cells at day 35 improved from $7.3 \pm 3.1\%$ (coverslip) and $7.8 \pm 1.2\%$ (rigid PMA) to $47.7 \pm 2.3\%$ (soft PMA; Supplementary Fig. 13b,c). The purity and yield of HB9⁺ MNs derived from soft PMAs were increased 6.1- and 10-fold, respectively, compared with controls (Supplementary Fig. 13c,d).

Our data presented in Fig. 1 suggested a connection between mechanotransduction and inhibition of BMP signalling in hPSCs. Indeed, immunoblots substantiated an inhibitory role of soft PMAs on Smad 1/5/8 phosphorylation (Fig. 3a). Phosphorylated Smads (phosphoSmads) require translocation to the nucleus for transcriptional activation¹². YAP/TAZ, nuclear transducers in the canonical Hippo pathway, bind phosphoSmads and control their nucleocytoplasmic shuttling in hESCs (ref. 13). In the Hippo pathway, Lats1/2 kinase phosphorylates YAP/TAZ,

which in turn bind the scaffolding protein 14-3-3 to remain cytoplasmic¹⁴. YAP/TAZ-mediated nuclear accumulation of phosphoSmads is required for hESC pluripotency and its loss results in neuroectoderm differentiation¹³. A recent study also shows that YAP/TAZ are critical in mechanotransduction^{15,16}.

Supporting our hypothesis that YAP/TAZ-mediated nuclear accumulation of Smads regulates rigidity-dependent neural induction of hESCs, nuclear YAP/TAZ were observed in $95 \pm 2.7\%$ and $38.2 \pm 1.6\%$ of hESCs on rigid and soft PMAs, respectively (Fig. 3b,c). Rigidity-dependent subcellular localization of YAP/TAZ was maintained after 3 d of neural induction and coincided with nucleocytoplasmic shuttling of Smad 2/3 and Smad 1/5/8 (Fig. 3d). Co-localization of YAP/TAZ and phosphoSmad 1/5/8 was also observed before and after neural induction (Supplementary Fig. 14).

We next examined whether substrate rigidity regulates YAP/TAZ in hESCs in a Hippo-dependent manner. Immunoblots demonstrated that soft PMAs promoted YAP phosphorylation on serine 127 (ser¹²⁷), a key target of Lats1/2 kinase downstream of the Hippo pathway (Fig. 3e). Silencing of *Lats1* expression using siRNA suppressed YAP phosphorylation on ser¹²⁷ (Supplementary Fig. 15). Notably, *Lats1* knockdown hESCs exhibited a spread morphology on both rigid and soft PMAs, indicating loss of mechanosensory properties (Supplementary Fig. 15). Soft PMAs did not promote neural induction of *Lats1* knockdown hESCs (Fig. 3f,g), further supporting that Lats-mediated phosphorylation of YAP may relay mechanical signals exerted by substrate rigidity.

Adult mammalian cell studies suggest functional links between two major intracellular mechanotransductive components, Rho GTPase and the actomyosin cytoskeleton (CSK), and the Hippo/YAP pathway¹⁶⁻¹⁹. Indeed, hESCs cultured on rigid PMAs exhibited bundled actin microfilaments at cell peripheries, whereas actin microfilaments were diffusely distributed throughout cells on soft PMAs (Supplementary Fig. 16). hESCs were treated independently with Y27632 (inhibitor of Rho-associated kinase, or ROCK, a downstream effector of RhoA), which decreases actomyosin contractility while maintaining intact CSK structure; cytochalasin D (CytoD), an inhibitor of actin polymerization; or lysophosphatidic acid (LPA), which stimulates RhoA and facilitates formation of actin microfilaments. Compared with controls, Y27632 promoted cytoplasmic localization of YAP/TAZ and significantly increased the percentage of Pax6⁺ NEs on rigid, but not soft PMAs (Supplementary Fig. 17 and Fig. 3h-j). In contrast, LPA facilitated nuclear localization of YAP/TAZ and inhibited neural induction on both rigid and soft PMAs. Notably, CytoD treatment had a significant effect on YAP/TAZ translocation on both rigid and soft PMAs, and neural induction was significantly more efficient on soft PMAs without treatment, compared with that on rigid PMAs treated with Y27632 (Fig. 3h-j). These data are consistent with findings that ROCK inhibition facilitates neural differentiation of mouse ESCs and embryonal carcinoma stem cells^{20,21}, whereas LPA maintains hESC pluripotency^{22,23}.

Together, our studies suggest a putative multi-targeted mechanotransductive process in hPSCs where soft PMAs inhibit Smad phosphorylation while activating Lats, an actin-binding protein²⁴, during disassembly of actin microfilaments. This in turn phosphorylates YAP to prevent nuclear translocation of phosphoSmads and inhibit subsequent activation of Smad target genes (Supplementary Fig. 18).

The Hippo/YAP signalling pathway is highly conserved and under intense study given its importance in development and cancer. The current view implicates Hippo signalling regulated by cell polarity and adhesion proteins. Our data reveal functional crosstalk between Hippo/YAP signalling and the classic mechanotransductive components, Rho GTPase and the actomyosin CSK, in hPSCs. Other mechanosensory components, such as adhesion molecules including integrins and cadherins, as

well as cell morphology may also regulate Hippo/YAP signalling in hPSCs. Supporting this view, increased internalization of integrins in mesenchymal stem cells on soft substrates represses Smad 1/5/8 phosphorylation and induces neurogenic differentiation²⁵. Also, NIH/3T3 cell morphology regulates the Hippo pathway¹⁶.

By combining dual Smad inhibition with PMAs, we generated high-yield and high-purity functional MNs from hPSCs. PMAs are fully defined substrates that can be mass produced for large-scale hPSC culture. We demonstrate that like many human adult stem cells^{26–30}, hPSCs are intrinsically mechanosensitive, and that substrate rigidity is an *in vitro* extracellular switch that directs NE versus NC lineage decisions and anterior versus posterior patterning. Mechanical properties of the extracellular matrix *in vivo* may act as a ‘mechanical organizer’, synergizing with morphogens to direct neural plate specification and anterior–posterior axis formation during neurulation.

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Author contributions

Y.S. and J.F. designed experiments; Y.S. performed *in vitro* differentiation experiments and Western blotting; K.M.A.Y., L.G.V.-D., and Y.S. generated and analysed gene expression data; K.M.A.Y. and Y.S. performed siRNA transfection; X.Z. performed electrophysiology measurements; L.G.V.-D. derived hiPSCs; W.C. fabricated silicon masters; S.W. developed image processing program; Y.S., L.G.V.-D., R.P., H.X., P.H.K. and J.F. analysed data and wrote the manuscript; J.F. supervised the project. All authors edited and approved the final manuscript.

Additional information

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Competing financial interests

The authors declare no competing financial interests.