



吕东媛

中国科学院力学研究所副研究员。2003年本科毕业于内蒙古农业大学获理学学士，2008年在内蒙古农业大学获发育生物学博士。2008-2011年在中国科学院力学研究所国家微重力实验室从事博士后流动工作(合作导师：龙勉研究员)，主要从事成体干细胞及胚胎干细胞力学-生物学耦合规律及分子机制研究。主持3项国家自然科学基金项目。已发表文章20余篇，其中在 *Biomaterials*、*ACS Appl Mater Inter*、*FASEB J* 等杂志发表的SCI文章，多次被 *Nat Rev Mol Cell Bio*、*Prog Mater Sci*、*Mat Sci Eng R*、*Adv Mater*、*Adv Funct Mater* 等杂志引用。授权发明专利1项。

代表性论文：

1. Li Z.[#], Gong Y.W., Sun S.J., Du Y., **Lü D.Y.**, Liu X.F., and Long M*. 2013. Differential Regulation of Stiffness, Topography, and Dimension of Substrate on Rat Mesenchymal Stem Cells. *Biomaterials*, **34**: 7616-7625.
2. **Lü D.Y.**[#], Liu X.F., Gao Y.X., Huo B., Kang Y.Y., Chen J., Sun S.J., Luo X.D., and Long M*. 2013. Asymmetric migration of human keratinocytes under mechanical stretch and cocultured fibroblasts in a wound repairing model. *PLoS One*, **8(9)**: e74563.
3. **Lü D.Y.**[#], Luo C.H., Zhang C., Li Z., and Long M*. 2014. Differential regulation of substrate stiffness and topography on morphology and stemness of mouse embryonic stem cells. *Biomaterials*, **35**: 3945-3955.
4. Wang J.W.[#], **Lü D.Y.**, Mao D.B., and Long M*. 2014. Mechanomics: An emerging field between biology and biomechanics. *Protein & Cell*, **5(7)**: 518-531.
5. Liu X.F.[#], Vargas D.A.[#], **Lü D.Y.**[#], Zhang Y., Zaman M.H.*[#], and Long M*. 2014. Computational modeling of stem cell migration: A mini review. *Cell. Mol. Bioengi.*, **7(2)**: 196-204.
6. Wang S.M.[#], Zhang Z.Y.*[#], **Lü D.Y.**[#], and Xu Q.X. 2015. Effects of mechanical stretching on the morphology and cytoskeleton of vaginal fibroblasts from women with pelvic organ prolapsed. *Int. J. Mol. Sci.*, **16(5)**: 9406-9419.

Regulation of morphology and differentiation of embryonic stem cells by substrate stiffness and topography 报告摘要：

Results and Discussion: Hepatic differentiation of hESCs on topographical PA hydrogel with two typical stiffnesses was systematically tested using a four-stage induction protocol, we proposed a working model to illustrate how the two mechanical factors cooperatively manipulate the hepatic differentiation of hESCs. Substrate stiffness and topography are critical in regulating cell functions, coupled impacts of topography and stiffness were found differentially at distinct differentiation stages. Substrate stiffness is dominant in stemness maintenance and defined endoderm differentiation. Substrate topography plays a decisive role in manipulating clone or cell morphology and spreading and favors the differentiation to hepatocyte-like cells formation and maturation. Both substrate stiffness and topography cooperatively support clone formation or cell shape maintenance, and the differentiated cells present liver-specific functions, such as albumin excretion, glycogen synthesis, and ICG engulfment.

Conclusions: These results indicate that stem cell morphology and differentiation are affected by substrate stiffness and topography that mimics physiological conditions, combined with those biochemical factors. Biomechanical cues are critical for enhancing the efficiency and maturity of hepatic differentiation of ESCs.