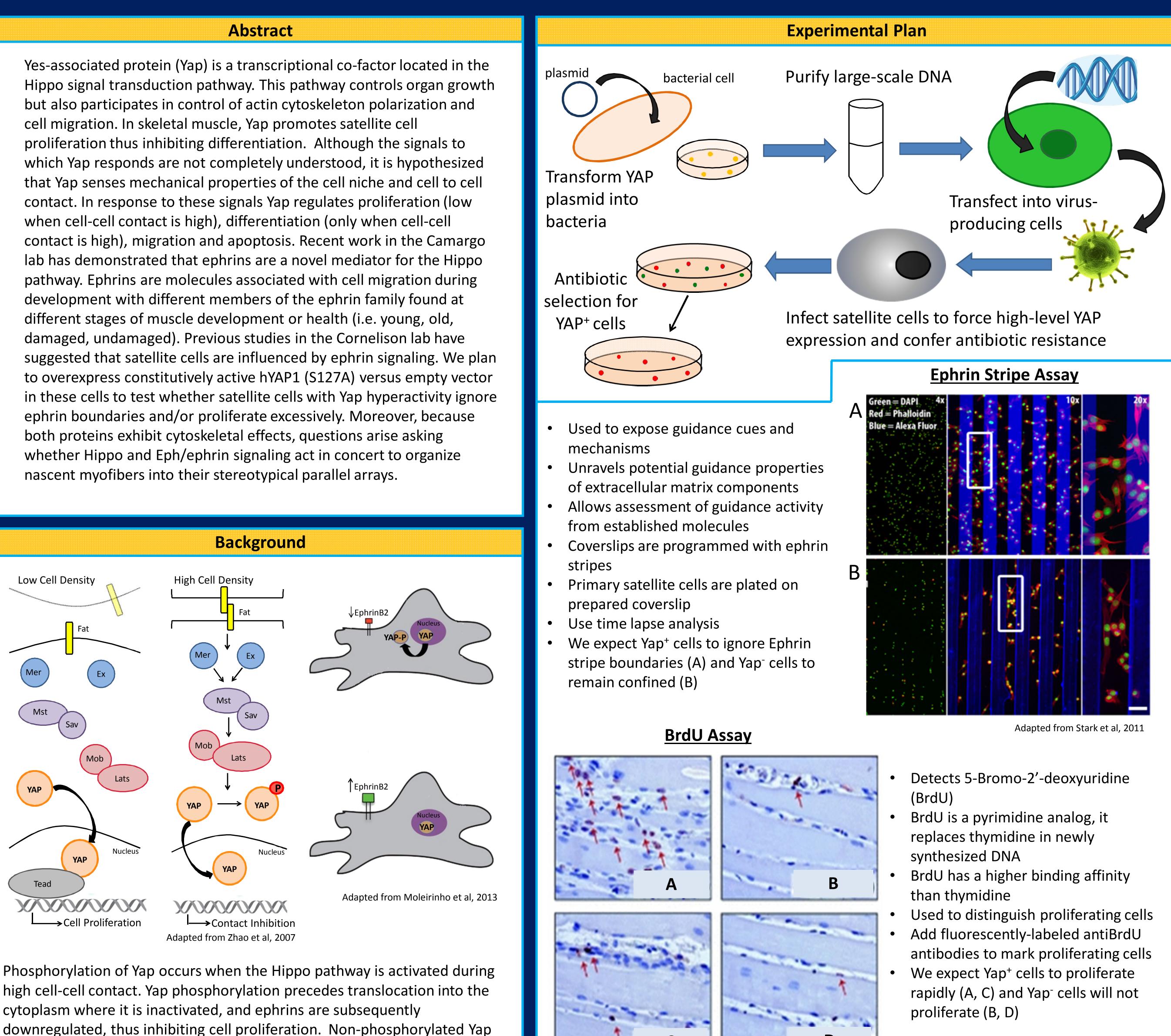
College of Veterinary Medicine<sup>1</sup>, Department of Biological Sciences<sup>2</sup> and Bond Life Sciences Center<sup>3</sup>, University of Missouri, Columbia, MO



cytoplasm where it is inactivated, and ephrins are subsequently downregulated, thus inhibiting cell proliferation. Non-phosphorylated Yap (active) remains within the nucleus where it functions as a transcriptional cofactor, promoting cell proliferation and upregulating ephrin expression.



# Alicia K. Tutino<sup>1</sup>, DDW Cornelison<sup>2,3</sup>

Adapted from Stratos et al, 2012

### **Potential Results**

# • Constitutively Active Yap

- Increase in ephrin expression
- Satellite cells do not obey ephrin boundaries
- Increase in proliferation of satellite cells
- Increase in activation of satellite cells
- Decrease in differentiation of satellite cells

### **Empty Vector**

- No change in ephrin expression
- Satellite cells obey ephrin boundaries
- No change in satellite cell proliferation, activation, or differentiation

# **Conclusion:**

- Effect on Regeneration:
  - In the constitutively active Yap, although the satellite cells are proliferating continuously, they will not be able to differentiate into mature myofibers. Therefore, regeneration is enhanced with the increase in satellite cells but stunted when they reach the point of differentiation • In the empty vector there is no effect on Yap and it would
  - act as it does normally
- Organization of Myofibers
  - Both the Hippo pathway and the Eph/ephrin signaling pathway work together in the organization of muscle fibers
  - Eph/ephrin repulsion is important in maintaining muscle type patterning
  - Hippo pathway is required to polarize actin cytoskeleton • The state of Yap can increase or decrease ephrin possibly
  - having an effect on this system

# **Future Studies**

The roles of the Eph/ephrin pathway and the Hippo/Yap pathway in muscle regeneration are not yet identified. If it is established that the Eph/ephrin and Hippo/Yap signaling pathways interact in muscle satellite cells, future studies will aim to identify specific Eph/ephrin proteins involved in Yap signaling. Ideally, this work will continue in vivo by testing genetic mouse mutants for specific Ephs and ephrins as well as Hippo and Yap.

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Dawn D. Cornelison cornelisond@missouri.edu Alicia K. Tutino akt4b2@mail.missouri.edu