

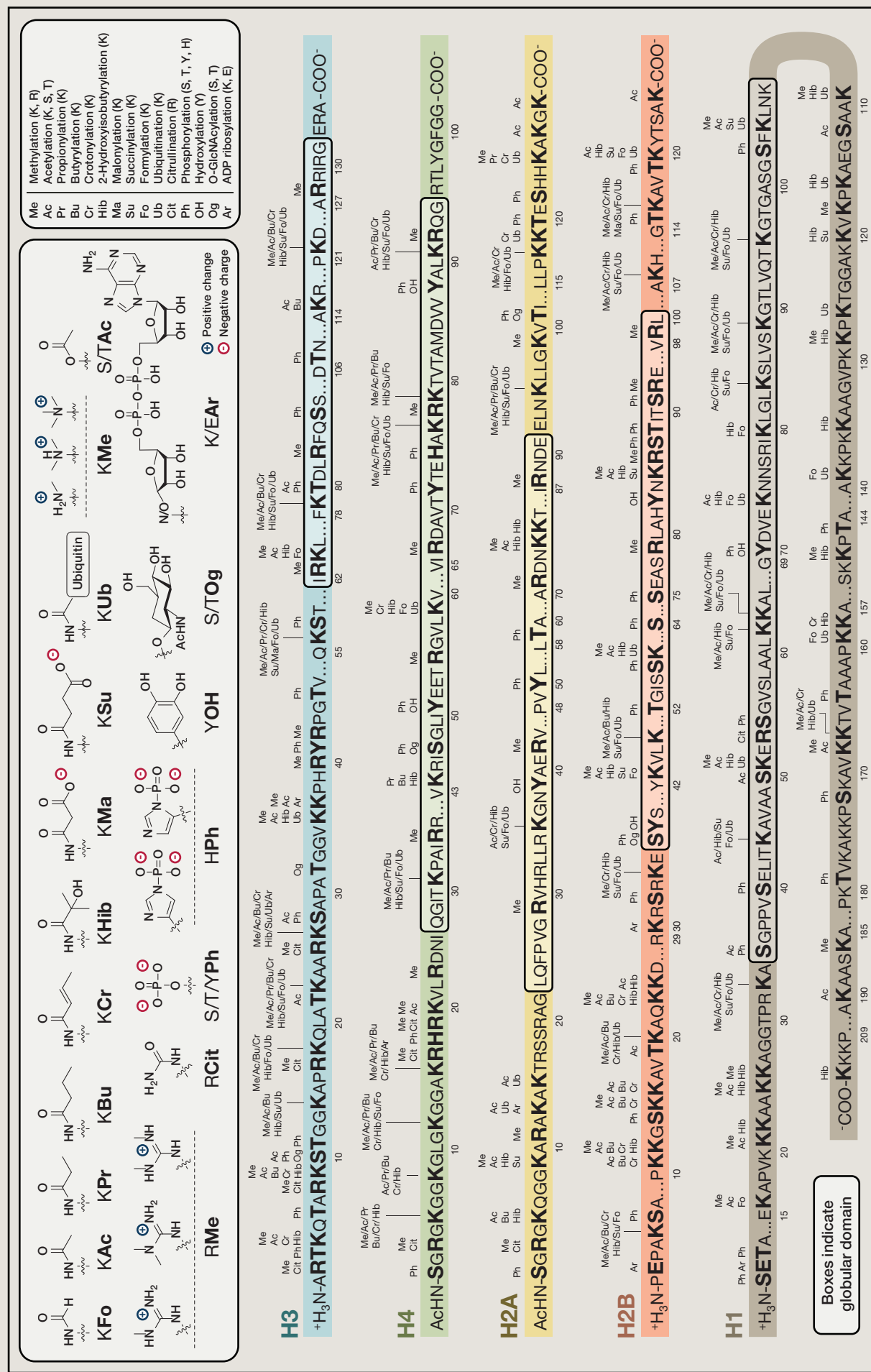
# SnapShot: Histone Modifications

He Huang,<sup>1</sup> Benjamin R. Sabari,<sup>2</sup> Benjamin A. Garcia,<sup>3</sup> C. David Allis,<sup>2</sup> and Yingming Zhao<sup>1</sup>

<sup>1</sup>Ben May Department of Cancer Research, The University of Chicago, Chicago, IL 60637, USA

<sup>2</sup>Laboratory of Chromatin Biology and Epigenetics, The Rockefeller University, New York, NY 10021, USA

<sup>3</sup>Department of Biochemistry and Biophysics, University of Pennsylvania, Philadelphia, PA 19104, USA



# SnapShot: Histone Modifications



He Huang,<sup>1</sup> Benjamin R. Sabari,<sup>2</sup> Benjamin A. Garcia,<sup>3</sup> C. David Allis,<sup>2</sup> and Yingming Zhao<sup>1</sup>

<sup>1</sup>Ben May Department of Cancer Research, The University of Chicago, Chicago, IL 60637, USA

<sup>2</sup>Laboratory of Chromatin Biology and Epigenetics, The Rockefeller University, New York, NY 10021, USA

<sup>3</sup>Department of Biochemistry and Biophysics, University of Pennsylvania, Philadelphia, PA 19104, USA

The histone proteins are decorated by a variety of protein posttranslational modifications (also called histone marks). Histone marks are critical to dynamic modulation of chromatin structure and function, contributing to the cellular gene expression program. In addition to the well-studied acetylation and methylation modifications, recent studies have revealed several new types of histone marks, including lysine propionylation, lysine butyrylation, lysine crotonylation, lysine 2-hydroxyisobutyrylation, lysine malonylation, and lysine succinylation. Preliminary studies on some of the new histone marks (e.g., crotonylation and 2-hydroxyisobutyrylation) suggest that their effects on chromatin function are distinct from those of lysine acetylation. Given that the newly discovered lysine acylation reactions likely use the corresponding acyl-CoA molecules as cofactors, it is proposed that histone acylations provide a link between cellular metabolism and epigenetic mechanisms.

This SnapShot summarizes the reported human, mouse, and rat histone marks, including recently identified lysine acylation marks.

## ACKNOWLEDGMENTS

This work was funded by institutional support from The Rockefeller University to C.D.A.; by NIH grants GM105933, CA160036, and RR020839 to Y.Z.; and by DP2OD007447 to B.A.G.

## REFERENCES

- Arnaudo, A.M., and Garcia, B.A. (2013). *Epigenet. Chromatin* 6, 24.
- Beck, H.C., Nielsen, E.C., Matthiesen, R., Jensen, L.H., Sehested, M., Finn, P., Grauslund, M., Hansen, A.M., and Jensen, O.N. (2006). *Mol. Cell. Proteomics* 5, 1314–1325.
- Chen, Y., Sprung, R., Tang, Y., Ball, H., Sangras, B., Kim, S.C., Falck, J.R., Peng, J., Gu, W., and Zhao, Y. (2007). *Mol. Cell. Proteomics* 6, 812–819.
- Dai, L., Peng, C., Montellier, E., Lu, Z., Chen, Y., Ishii, H., Debernardi, A., Buchou, T., Rousseaux, S., Jin, F., et al. (2014). *Nat. Chem. Biol.* 10, 365–370.
- Fierz, B., and Muir, T.W. (2012). *Nat. Chem. Biol.* 8, 417–427.
- Jenuwein, T., and Allis, C.D. (2001). *Science* 293, 1074–1080.
- Kouzarides, T. (2007). *Cell* 128, 693–705.
- Tan, M., Luo, H., Lee, S., Jin, F., Yang, J.S., Montellier, E., Buchou, T., Cheng, Z., Rousseaux, S., Rajagopal, N., et al. (2011). *Cell* 146, 1016–1028.
- Xie, Z., Dai, J., Dai, L., Tan, M., Cheng, Z., Wu, Y., Boeke, J.D., and Zhao, Y. (2012). *Mol. Cell. Proteomics* 11, 100–107.
- Young, N.L., Dimaggio, P.A., and Garcia, B.A. (2010). *Cell. Mol. Life Sci.* 67, 3983–4000.