Accepted Manuscript

Targeting Chromatin <u>Aging</u> - The Epigenetic Impact of <u>Longevity-Associated</u> Interventions

Adam Field, Peter D. Adams

 PII:
 S0531-5565(16)30583-6

 DOI:
 doi:10.1016/j.exger.2016.12.010

 Reference:
 EXG 9954

To appear in: Experimental Gerontology

Received date:28 OctRevised date:5 DeceAccepted date:10 Dec

28 October 2016 5 December 2016 10 December 2016 Experimental Gerontology



Please cite this article as: Field, Adam, Adams, Peter D., Targeting Chromatin <u>Aging</u> - The Epigenetic Impact of Longevity-Associated Interventions, *Experimental Gerontology* (2016), doi:10.1016/j.exger.2016.12.010

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

ACCEPTED MANUSCRIPT

"Targeting Chromatin Aging - The Epigenetic Impact of Longevity-Associated

Interventions"

Adam Field, Peter D Adams

Glasgow G61 1BD United Kingdom

E-mail: a.field.1@research.gla.ac.uk

0.0 Abstract

A rapidly growing body of evidence has shown that chromatin undergoes radical alterations as an organism ages, but how these changes relate to <u>aging</u> itself is an open question. It is likely that these processes contribute to genomic instability and loss of transcriptional fidelity, which in turn drives deleterious <u>age-related</u> phenotypes. Interventions associated with increased healthspan and longevity such as reduced insulin / IGF signalling (IIS), inhibition of mTOR and energy depletion resulting in SIRT1 / AMPK activation, all have beneficial effects which ameliorate multiple facets of <u>age-associated</u> decline. The impact of these interventions to act directly upon the epigenome and promote a youthful chromatin landscape, maintaining genetic and transcriptional memory throughout the lifecourse. We propose that this is a fundamental mechanism through which these interventions are able to curtail the incidence of <u>age-related</u> disease. By revisiting these well characterised interventions, we may be able to identify targetable effectors of chromatin function and use this knowledge to enhance healthspan and longevity in human populations through the measured application of dietary and small molecule interventions.

Keywords

Epigenetics; <u>Aging</u>; Histone; Methylation; Longevity; Chromatin; Rapamycin; Insulin; Calorie Restriction; Healthspan; mTOR; AMPK; SIRT1; IGF; Download English Version:

https://daneshyari.com/en/article/5501317

Download Persian Version:

https://daneshyari.com/article/5501317

Daneshyari.com