

Towards disease-oriented dosing of rapamycin for longevity

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Relations between aging and age-related diseases (ARDs). Credit: *Aging* (2023). DOI: 10.18632/aging.204920

A new research perspective titled "Towards disease-oriented dosing of rapamycin for longevity: does aging exist or only age-related diseases?" has been published in *Aging*.



In his new research perspective, Dr. Mikhail V. Blagosklonny from Roswell Park Comprehensive Cancer Center discusses aging and rapamycin (Sirolimus)—the only drug that consistently extends <u>life span</u> in countless animal studies in all species tested. He writes that individuals taking rapamycin and those not taking it will ultimately succumb to age-related diseases. However, if administered in diseaseoriented dosages for an extended period of time, individuals taking rapamycin may experience a delayed onset of such diseases, and live longer.

"The goal is to delay a particular disease that is expected to be lifelimiting in a particular person," says Dr. Blagosklonny.

Age-related diseases, quasi-programmed during development, progress at varying rates in different individuals. Rapamycin is a prophylactic antiaging drug that decelerates early development of <u>age-related diseases</u>. Dr. Blagosklonny further discusses the hyperfunction theory of quasiprogrammed diseases, which challenges the need for the traditional concept of aging itself.

"I emphasize that aging is not programmed, but in contrast, quasiprogrammed. 'Quasi' means pseudo; seemingly; apparently but not really. Some scientists deliberately represent hyperfunction theory as theory of programmed aging. It's the opposite. Quasi-program is a continuation of a real program. Quasi-program has no intent, no purpose and it is always harmful," concludes Dr. Blagosklonny.

More information: Mikhail V. Blagosklonny, Towards diseaseoriented dosing of rapamycin for longevity: does aging exist or only agerelated diseases?, *Aging* (2023). <u>DOI: 10.18632/aging.204920</u>



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