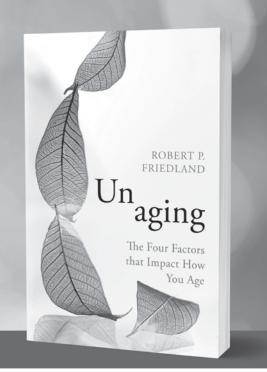
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The robert P. FRIEDLAND aging

The Four Factors that Impact How You Age



9781009087742| Paperback| £14.99 / \$19.95 | October 2022

Aging is a subject of concern to everyone, but is widely misunderstood. If we view it as inevitable, we miss the fact that not everyone is able to grow to an old age. Realization of this reality helps us to understand that aging presents a wonderful opportunity - an opportunity to make choices about how we live which can enhance the aging process and offer a chance to live to our potential. This book clearly presents the four, multiple reserve, factors (cognitive, physical, psychological and social) which impact our ability to have healthy responses to the stresses of aging. By giving the biological basis for the advice given, you will learn the steps to take in your activities, diet and mental outlook to grasp the opportunity that aging offers. Everyone must know that what we do makes a difference.

"Rooted on his vast clinical and research experience, Dr. Friedland takes us on an accessible scientific tour to demystify the inevitability of aging."

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"This book is a 'must read' for anyone who would like to use the latest scientific studies to help them live healthier lives as they age."

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FOR THE PREVENTION OF MIGRAINES, CONSIDER AIMOVIG®

PrAimovig® (erenumab injection) is indicated for prevention of migraine in adults who have at least 4 migraine days per month.

Aimovig® reduced the number of monthly migraine days vs. placebo

In patients with episodic migraines, Aimovig® significantly reduced the mean MMD from baseline to months 4–6 vs. placebo: -3.7 (140 mg), -3.2 (70 mg) vs. -1.8 days, p<0.0011*

Migraine Physical Function Impact Diary (MPFID) data

In patients with episodic migraines, Aimovig®-treated patients reported significantly improved scores from baseline on domains of the MPFID at months 4-6 vs. placebo (secondary endpoint):^{2*†}

- Physical Impairment: -4.8 (140 mg), -4.2 (70 mg) vs. -2.4 points, p<0.001
- Everyday Activities: -5.9 (140 mg), -5.5 (70 mg) vs. -3.3 points, p<0.001

FOR MORE INFORMATION ABOUT AIMOVIG®, VISIT PRO.NOVARTIS.CA



To access the pro.novartis.ca portal for health care professionals (HCP) from the public site, you will need to validate that you are an HCP

Clinical use: Safety and efficacy has not been studied in patients below the age of 18. Safety and efficacy has not been studied in patients aged 65 or older.

Other relevant warnings and precautions:

- · Constipation with serious complications
- · If a serious or severe hypersensitivity reaction occurs (including rash, angioedema and anaphylactoid reactions), discontinue Aimovig® and initiate appropriate therapy.
- · Aimovig® crosses the placenta. Avoid use during pregnancy as a cautionary measure.
- In nursing women, a decision should be made whether to discontinue nursing or discontinue Aimovig®, weighing the potential benefit of Aimovig® to the mother and breastfeeding to the infant.

For more information:

Please consult the Product Monograph at https://health-products.canada.ca/dpd-bdpp/index-eng.jsp for important information relating to adverse reactions, drug interactions, and dosing information which have not been discussed in this piece. The Product Monograph is also available by calling us at 1-800-363-8883.

References: 1. Aimovig® Product Monograph, April 12, 2022. Novartis Pharmaceuticals Canada Inc. 2. Goadsby PJ, et al. A Controlled Trial of Erenumab for Episodic Migraine. N Engl J Med. 2017;377:2123-32. MMD=monthly migraine days

- * Randomized, multi centre, 24 week, placebo controlled, double blind study. Patients had a history of migraine with or without aura for a duration of ≥12 months and 4–14 migraine days per month. Baseline ~8 migraine days/month. AIMOVIG* 70 mg (n=312), 140 mg (n=318), placebo (n=316),
- † The MPFID is a 13-item self-administered instrument measuring physical functioning in the previous 24 hours on migraine and non-migraine days, using an electronic diary. Scores were averaged over 1 month and transformed to a 100-point scale.²



Novartis Pharmaceuticals Canada Inc. 385 Bouchard Blvd. Dorval (Quebec) H9S 1A9 www.novartis.ca Tel: 514-631-6775 Fax: 514-631-1867 Medical Information Tel: 1-800-363-8883 © 2022 Novartis Pharmaceuticals Canada Inc. All rights reserved.

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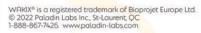
WAKIX® (pitolisant hydrochloride tablets) is indicated for the treatment of excessive daytime sleepiness (EDS) or cataplexy in adult patients with narcolepsy.1

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- · Contraindications in patients with hypersensitivity to pitolisant hydrochloride, severe hepatic impairment and in breastfeeding patients.
- Warnings and precautions regarding QTc prolongation, drug abuse, misuse, dependence and rebound effect, driving or operating machinery, increased exposure with moderate hepatic impairment, seizures or worsening of seizures in patients with a history of epilepsy, suicidal ideation in patients with history of psychiatric disorders, renal impairment, end stage renal disease, reduced effectiveness of hormonal contraceptives, use in pregnancy, avoiding pregnancy, fertility, use in pediatrics and use in geriatrics.
- Adverse reactions, drug interactions, dosing, and conditions of clinical use. The product monograph is also available by calling 1-888-867-7426 or by email at medinfo@paladin-labs.com.
- * Comparative clinical significance is unknown. † Clinical significance is unknown.

1. WAKIX® Product Monograph. Paladin Labs Inc. 2. Data on file. Paladin Labs Inc.













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- provides an integrated, comprehensive, inter-disciplinary approach to stroke prevention and case management for these high-risk patients.
- facilitates planning and implementation of primary and secondary stroke prevention strategies for the respective area of the region.
- works in partnership with primary care, acute care, stroke rehabilitation, community care access centres and other stake-holders in planning regional stroke prevention strategies.
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