PEAK PERFORMANCE & PREVENTION

# **Clinical Applications**

- Supports Healthy Immune Function\*
- · Promotes Healthy Skin and Mucous Membranes\*
- Provides Vitamin A Repletion in Cases of Deficiency\*

**P3:** A provides an excellent source of supplemental vitamin A. Vitamin A is a fat-soluble vitamin necessary for growth, bone development, vision, hormone receptor function, cellular health, immune function, and the integrity of epithelial and mucosal surfaces.\*

### All Peak Performance & Prevention Formulas Meet or Exceed cGMP Quality Standards

### Discussion

Vitamin A is a fat-soluble vitamin that has long been recognized as an essential micronutrient for growth and bone development, eye health/ night vision, cellular proliferation and differentiation, immune function, reproduction, and the integrity of cells that line the epithelial and mucosal surfaces. The generic term "vitamin A" encompasses a number of related metabolic compounds: retinyl esters and retinol (which is converted to retinoic acid) are considered to be preformed vitamin A and are herein referred to as vitamin A.\*

Vitamin A supplementation plays a role in conditions that challenge the immune system and impair cellular health.<sup>[1]</sup> P3: A is formulated to provide support for healthy immune function and to promote healthy skin and mucous membranes.\*

#### **Immune Support**

Research supporting the foundational role of vitamin A in immune system function dates back to the 1920s.<sup>[2]</sup> Modern molecular and animal studies attribute the value of vitamin A to its involvement in maintaining mucosal barriers that line the airways, digestive tract, and urinary tract. The neutrophils, macrophages, and natural killer cells in these tissues provide the body's first line of immune defense. Vitamin A also plays a role in cellular differentiation and proliferation and in the generation of antibody responses to T-cell dependent and independent antigens.<sup>\*[1,3-5]</sup>

Vitamin A metabolites are involved in promoting differentiation of regulatory T cells (Tregs), a subpopulation of T cells involved in maintaining cellular tolerance to self-antigens and regulating autoimmune response. In vitro studies suggest that vitamin A contributes to enhancing the stability and functionality of Tregs under inflammatory conditions.<sup>[6,7]</sup> The changes in mucosal epithelial regeneration and immune function attributed to vitamin A activity may account for the increased mortality seen in vitamin A-deficient populations throughout the world. <sup>[6,8]</sup> However, although the beneficial effects of vitamin A on immune function are well-established at the molecular level, more research is needed to elucidate ongoing therapeutic effects.\*

#### **Epithelial and Mucous Membranes**

The function of vitamin A in the maintenance of epithelial membranes is crucial. With essential roles in cell morphology and as a cell differentiation enhancer, vitamin A is an integral part of nonspecific, innate immunity function when intestinal and respiratory tissues require defense against invaders. It has been suggested that vitamin A also improves the mechanistic defense of oral mucosa, increases the integrity of intestinal mucus barrier, and maintains the morphology and amount of urothelial cells.<sup>[6,9]</sup> Under conditions of vitamin A deficiency, epithelial cells shrink, leading to a wide variety of symptoms such as a decreased resistance to pathogens that leaves cells vulnerable due to impaired function.<sup>\*[6,10,11]</sup>

Vitamin A retinoids also regulate the growth and differentiation of many cell types within skin. In distressed tissue, vitamin A appears to stimulate epidermal turnover, increase the rate of re-epithelialization, and restore epithelial structure. Additionally, retinoids appear to have the ability to counteract the inhibitory effects of steroids in impaired tissues. In a double-blind, placebo-controlled study in animals, it was suggested that retinoids stimulate the expression of transforming growth factor-beta (TGF-beta) and insulin-like growth factor (IGF) and increase collagen production.<sup>[12]</sup> Mechanisms continue to be elucidated in order to increase the understanding of the beneficial effects of vitamin A and its metabolites on skin. Vitamin A continues to be used both orally and topically for a wide variety of skin issues.<sup>\*[13]</sup>

#### Vitamin A Repletion in Cases of Deficiency

Vitamin A deficiency is common in many developing countries, and health consequences are most commonly seen during high-nutritionaldemand life stages, such as infancy, childhood, pregnancy, and lactation. The most common symptom of deficiency is an inability to see in low light, and this can progress to further visual complications if not corrected. Other symptoms can include frequent bouts of impaired immune health, delayed growth, skin issues, and infertility. Although most people in the United States get enough vitamin A from their diet, certain groups are more susceptible to deficiency and have impaired storage or transport of vitamin A due to a variety of health factors. Individuals who suffer from poor absorption of lipids due to bowel inflammation or submaximal pancreatic or biliary secretions are at particular risk. Oral ingestion of vitamin A is effective for replenishing and improving deficiency symptoms that may develop from a diet lacking in vitamin A or in individuals who cannot fully metabolize what they consume.<sup>\*(15,8)</sup> P3: A

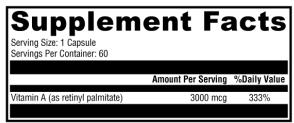
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\*These statements have not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure, or prevent any disease.



Other Ingredients: Microcrystalline cellulose, capsule (hypromellose and water), ascorbyl palmitate, and silica.

# Directions

Take one capsule daily, or as directed by your healthcare practitioner.

Consult your healthcare practitioner prior to use. Individuals taking medication should discuss potential interactions with their healthcare practitioner. Not recommended for long-term use or for use during pregnancy, during lactation, or when trying to conceive. Do not use if tamper seal is damaged.

# **Does Not Contain**

Wheat, gluten, yeast, soy, animal or dairy products, fish, shellfish, peanuts, tree nuts, egg, ingredients derived from genetically modified organisms (GMOs), artificial colors, artificial sweeteners, or artificial preservatives.

### <u>SAFETY</u>

Vitamin A is an essential nutrient required for many physiological functions; however, like other fat-soluble vitamins, it is capable of building to toxic levels when consumed in excess. Vitamin A is stored in the body, primarily in liver tissue, as retinyl ester. Because of the potential for toxicity, the Institute of Medicine's Food and Nutrition Board set an upper level (UL) of 3,000 mcg retinol activity equivalents (RAE), or 10,000 IU, in 2001. This value was based on research looking at teratogenicity for women of childbearing age and at potential for liver toxicity in all populations.<sup>\*[1]</sup>

Although healthy fetal development requires sufficient vitamin A intake, levels in excess of 3,000 mcg RAE/day during early pregnancy have been linked to birth defects. Because vitamin A excess and deficiency have the potential for increasing the risk of birth defects, particular caution should be taken with high-potency vitamin A products during pregnancy or when trying to conceive.\*

Research at the time the UL was set suggested a link between changes in bone density and high vitamin A intake, but the findings were confounding and therefore not considered. Since that time, researchers have published limited experimental data suggesting that vitamin A, when taken at levels above the UL, may affect the development of bone remodeling cells and may stimulate bone matrix degradation. <sup>[14]</sup> Prospective research has suggested that vitamin A in excess of the UL may be linked to an increased risk of hip fracture.<sup>[15]</sup> As with prenatal development, too little or too much can impact bone health. Sensitive populations can include but are not limited to postmenopausal women and patients with a history of bone density issues or frequent fractures. Further research is needed to update monitoring protocols for practitioners prescribing vitamin A supplementation to ensure optimal serum retinol levels.\*

Dose levels beyond the UL can be necessary in some populations to prevent or reverse health consequences from vitamin A deficiency. Because the safety of vitamin A is dependent on both the daily level of intake and duration of consumption, P3: A is offered in two potency options to meet individual needs. However, due to potential safety concerns in certain populations, P3: A is labeled with a precautionary statement and is not endorsed for long-term use.\*

## References

1. Vitamin A. In: Food and Nutrition Board, Institute of Medicine. Dietary Reference Intakes for Vitamin A, Vitamin K, Arsenic, Boron, Chromium, Copper, Iodine, Iron, Manganese, Molybdenum, Nickel, Silicon, Vanadium, and Zinc. Washington, DC: National Academy Press; 2001:82-161. https://www.nap.edu/read/10026/chapter/6 Accessed August 29, 2019.

2. Green HN, Mellanby E. Vitamin A as an anti-infective agent. Br Med J. 1928 Oct 20;2(3537):691-6. [PMID: 20774205]

3. Blomhoff HK, Smeland EB, Erikstein B, et al. Vitamin A is a key regulator for cell growth, cytokine production, and differentiation in normal B cells. J Biol Chem. 1992 Nov 25;267(33):23988-92. [PMID: 1429735]

4. Brown CC, Noelle RJ. Seeing through the dark: new insights into the immune regulatory functions of vitamin A. Eur J Immunol. 2015 May;45(5):1287-95. [PMID: 25808452]

5. Semba RD. Vitamin A, immunity, and infection. Clin Infect Dis. 1994 Sep;19(3):489-99. [PMID: 7811869]

6. Huang Z, Liu Y, Qi G, et al. Role of vitamin A in the immune system. J Clin Med. 2018 Sep;7(9): 258. [PMID: 30200565]

7. Lu L, Lan Q, Li Z, et al. Critical role of all-trans retinoic acid in stabilizing human natural regulatory T cells under inflammatory conditions. Proc Natl Acad Sci USA. 2014 Aug19;111(33):E3432-40. [PMID: 25099355]

8. Stephensen CB. Vitamin A, infection, and immune function. Annu Rev Nutr. 2001;21:167-92. [PMID: 11375434]

9. McCullough FS, Northrop-Clewes CA, Thurnham DI. The effect of vitamin A on epithelial integrity. Proc Nutr Soc. 1999 May;58(2):289-93. [PMID: 10466169]

10. Oliveira LM, Teixeira FME, Sato MN. Impact of retinoic acid on immune cells and inflammatory diseases. *Mediators Inflamm.* 2018 Aug 9;2018:3067126. IPMID: 301588321

11. de Medeiros PHQS, Pinto DV, de Almeida JZ, et al. Modulation of intestinal immune and barrier functions by vitamin A: implications for current understanding of malnutrition and enteric infections in children. Nutrients. 2018 Aug 21;10(9):1128. [PMID: 30134532]

12. Wicke C, Halliday B, Allen D, et al. Effects of steroids and retinoids on wound healing. Arch Surg. 2000 Nov;135(11):1265-70. [PMID: 11074878]

13. Polcz ME, Barbul A. The role of vitamin A in wound healing. Nutr Clin Pract. 2019 Aug 7. [PMID: 31389093]

14. Conaway HH, Henning P, Lerner UH. Vitamin a metabolism, action, and role in skeletal homeostasis. Endocr Rev. 2013 Dec;34(6):766-97. [PMID: 23720297]

15. Wu AM, Huang CQ, Lin ZK, et al. The relationship between vitamin A and risk of fracture: meta-analysis of prospective studies. J Bone Miner Res. 2014 Sep;29(9):2032-9. [PMID: 24700407]

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