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Abstract

Maintaining leanness and a physically active lifestyle during adulthood reduces systemic inflammation, an underlying factor in multiple chronic diseases. The anti-inflammatory influence of near-daily physical activity in lowering C-reactive protein, total blood leukocytes, interleukin-6, and other inflammatory cytokines may play a key role in lowering risk of cardiovascular disease, certain types of cancer, type 2 diabetes, sarcopenia, and dementia. Moderate exercise training causes favorable perturbations in immunity and a reduction in incidence of upper respiratory tract infection (URTI). During each bout of moderate exercise, an enhanced recirculation of immunoglobulins, neutrophils, and natural killer cells occurs that persists for up to 3-h post-exercise. This exercise-induced surge in immune cells from the innate immune system is transient but improves overall surveillance against pathogens. As moderate exercise continues on a near-daily basis for 12–15 weeks, the number of symptoms days with URTI is decreased 25%–50% compared to randomized sedentary controls. Epidemiologic and animal studies support this inverse relationship between URTI risk and increased physical activity.

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1. Introduction

Exercise immunology, a relatively new area of scientific endeavor, is the study of acute and chronic effects of various exercise workloads on the immune system and immunosurveillance against pathogens. Two areas of investigation from exercise immunology have clinical and public health implications: (1) the chronic anti-inflammatory influence of exercise training; (2) the reduction in risk of upper respiratory tract infections (URTI) from regular moderate exercise training.

2. Anti-inflammatory influence of exercise training

Acute inflammation is a normal response of the immune system to infection and trauma. Intense and prolonged exercise similar to marathon race competition causes large but transient increases in total white blood cells (WBC) and a variety of cytokines including interleukin-6 (IL-6), IL-8, IL-10, IL-1 receptor antagonist (IL-1ra), granulocyte colony stimulating factor (GCSF), monocyte chemoattractant protein 1 (MCP-1), macrophage inflammatory protein 1β (MIP-1β), tumor necrosis factor-α (TNF-α), and macrophage migration inhibitory factor (MIF). C-reactive protein (CRP) is also elevated following heavy exertion, but the increase is delayed in comparison to most cytokines.

Despite regular increases in these inflammation biomarkers during each intense exercise bout, endurance athletes have lower resting levels in contrast to overweight and unfit adults. For example, mean CRP levels in long distance runners (rested state) typically fall below 0.5 mg/L in comparison to 4.0 mg/L and higher in obese, postmenopausal women.
The persistent increase in inflammation biomarkers is defined as chronic or systemic inflammation, and is linked with multiple disorders and diseases including atherosclerosis and cardiovascular disease (CVD), the metabolic syndrome, diabetes mellitus, sarcopenia, arthritis, osteoporosis, chronic obstructive pulmonary disease, dementia, depression, and various types of cancers.⁵–⁷ CRP is the most frequently measured inflammatory biomarker, and individuals with CRP values in the upper tertile of the adult population (>3.0 mg/L) have a 2-fold increase in CVD risk compared to those with a CRP concentration below 1.0 mg/L.⁷ An elevated fasting IL-6 concentration is a significant component of the chronic low-grade inflammation that underlies the metabolic syndrome, CVD, diabetes, and various cancers.⁶ Athletes typically have plasma IL-6 concentrations that fall below 1.0 pg/mL in contrast to values above 2.0 pg/mL in older and obese individuals.⁵,⁸

2.1. Physical activity, fitness, and chronic inflammation

Large population observational studies consistently show reduced WBC, CRP, IL-6, TNF-α, and other inflammatory biomarkers in adults with higher levels of physical activity and fitness, even after adjustment for potential confounders.⁹–¹⁴ The inverse association between physical activity/fitness and inflammation is related in part to the effect of activity on fat mass.¹¹ In most studies, however, adjustment for body mass index (BMI) and adiposity attenuates but does not negate the strength of the relationship between inflammatory biomarkers and physical activity/fitness.¹¹,¹⁵ For example, in a study of 1002 community-dwelling adults (18–85 years), a general linear model (GLM) analysis adjusted CRP means for frequency of physical activity, BMI, and several other lifestyle and demographic factors.¹⁵ BMI had the strongest effect on CRP followed by gender (higher in females), exercise frequency, age, and smoking status (see Fig. 1).

Randomized, controlled exercise-intervention studies provide equivocal support for the inverse relationship between increased physical activity and reduced systemic inflammation.¹¹,¹⁶–²² One explanation is that in comparison to the large variance evaluated in observational studies, the change in aerobic fitness and activity levels is typically of low magnitude in randomized exercise trials, the duration of training seldom extends beyond 6 months, and the number of subjects is relatively low.¹⁷,¹⁸,²⁰,²³ Nonetheless, data from both study formats support that in order for reductions in chronic inflammation to be experienced, a large change in a combination of lifestyle factors is needed including weight loss, near-daily moderate-to-vigorous physical activity of 30–60 min duration, avoidance of cigarette smoking, and increased intake of fruits and vegetables.²²,²³ For example, if an obese, older individual adds three weekly 30-min walking sessions to the lifestyle, reductions in chronic inflammation are unlikely to be experienced unless the exercise workload is increased in combination with significant weight loss and improved diet quality.

2.2. Potential mechanisms

When successful, exercise training may exert anti-inflammatory influences through a reduction in visceral fat mass and the induction of an acute anti-inflammatory environment with each bout of exercise that over time becomes chronic.²⁴,²⁵ These effects may be mediated in part through muscle-derived peptides or myokines, but this proposed mechanism needs further testing.²⁵ Contracting skeletal muscles release myokines (e.g., IL-6, IL-8, IL-15) that may exert both direct and chronic anti-inflammatory effects.

The first identified and most studied myokine is IL-6. During prolonged and intense exercise, IL-6 is produced by muscle fibers and stimulates the appearance in the circulation of other anti-inflammatory cytokines such as IL-1ra and IL-10.²⁶ IL-6 also inhibits the production of the proinflammatory cytokine TNF-α and stimulates lipolysis and fat oxidation.²⁶ With weight loss from energy restriction and exercise, plasma levels of IL-6 fall, skeletal muscle TNF-α decreases, and insulin sensitivity improves.²⁷,²⁸ Thus IL-6 release from the exercising muscle may help mediate some of the health benefits of exercise including metabolic control of type 2 diabetes.²⁷,²⁸

Fig. 1. C-reactive protein (CRP) adjusted means for frequency of physical activity, body mass index (BMI), and several other lifestyle and demographic factors.¹⁵
Muscle IL-6 release, however, is very low during moderate physical activity. For example, during a 30-min brisk walk on a treadmill, plasma IL-6 concentrations increased from 1.3 pg/mL to 2.0 pg/mL in female subjects. The increase in IL-6 during brisk walking is probably insufficient to mediate anti-inflammatory and other beneficial health effects, and additional research is needed to determine the relative contribution of myokines compared to other exercise-induced factors. The acute exercise-induced increase in IL-6 after heavy exertion (e.g., typically above 5 pg/mL, 10 pg/mL, and 50 pg/mL following 1-h, 2-h, and marathon race running bouts, respectively) may indeed orchestrate anti-inflammatory influences, lipolysis, and improved insulin sensitivity, but this amount of physical activity is beyond levels achievable by most overweight/obese individuals.

A moderate exercise program of near daily 30-min walking bouts, without diet control, has small influences on visceral fat, even in long-term studies. This is further evidence that the myokine hypothesis does not apply at the activity level attainable by most middle-aged and elderly individuals. Thus moderate physical activity training must be increased to the highest levels acceptable to an individual (e.g., 60 min a day) and combined with weight loss through tight control of energy intake and improved diet quality to achieve reductions in systemic inflammation.

3. URTI risk reduction from regular moderate exercise training

URTI is the most frequently occurring infectious disease in humans worldwide. More than 200 different viruses cause the common cold, and rhinoviruses and coronaviruses are the culprits 25%—60% of the time. The National Institute of Allergy and Infectious Diseases reports that people in the USA suffer one billion colds each year with an incidence of 2.22. The increase in IL-6 during brisk walking is probably insufficient to mediate anti-inflammatory and other beneficial health effects, and additional research is needed to determine the relative contribution of myokines compared to other exercise-induced factors. The acute exercise-induced increase in IL-6 after heavy exertion (e.g., typically above 5 pg/mL, 10 pg/mL, and 50 pg/mL following 1-h, 2-h, and marathon race running bouts, respectively) may indeed orchestrate anti-inflammatory influences, lipolysis, and improved insulin sensitivity, but this amount of physical activity is beyond levels achievable by most overweight/obese individuals.

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3.1. Moderate physical activity and URTI risk

Several lines of evidence support the linkage between moderate physical activity and improved immunity and lowered infection rates: survey, animal, epidemiologic, and randomized training data. Survey data consistently support the common belief among fitness enthusiasts that regular exercise confers resistance against infection. In surveys, 80%—90% of regular exercisers perceive themselves as less vulnerable to viral illnesses compared to sedentary peers.

Animal studies are difficult to apply to the human condition, but in general, support the finding that moderate exercise lowers morbidity and mortality following pathogen inoculation, especially when compared to prolonged and intense exertion or physical inactivity. Mice infected with the herpes simplex virus, for example, and then exposed to 30-min of moderate exercise experience a lower mortality during a 21-day period compared to higher mortality rates after 2.5 h of exhaustive exercise or rest. Another study with mice showed that 3.5 months of moderate exercise training compared to no exercise prior to induced influenza infection decreased symptom severity and lung viral loads and inflammation.

Retrospective and prospective epidemiologic studies have measured URTI incidence in large groups of moderately active and sedentary individuals. Collectively, the epidemiologic studies consistently show reduced URTI rates in physically active or fit individuals. A one-year epidemiologic study of 547 adults showed a 23% reduction in URTI risk in those engaging in regular versus irregular moderate-to-vigorous physical activity. In a group of 145 elderly subjects, URTI symptomatology during a one-year period was reduced among those engaging in higher compared to lower amounts of moderate physical activity. During a one-year study of 142 males aged 33—90 years, the odds of having at least 15 days with URTI was 64% lower among those with higher physical activity patterns. A cohort of 1509 Swedish men and women aged 20—60 years were followed for 15 weeks during the winter/spring. Subjects in the upper tertile for physical activity experienced an 18% reduction in URTI risk, but this proportion improved to 42% among those with high perceived mental stress.

A group of 1002 adults (18—85 years, 60% female, 40% male) were followed for 12 weeks (half during the winter, half during the fall) while monitoring URTI symptoms and severity using the Wisconsin Upper Respiratory Symptom Survey. Subjects reported frequency of moderate-to-vigorous aerobic activity, and rated their physical fitness level using a 10-point Likert scale. The number of days with URTI was 43% lower in subjects reporting an average of five or more days of aerobic exercise (20-min bouts or longer) compared to those who were largely sedentary (<1 day per week) (see Fig. 2). This relationship occurred after adjustment for important confounders including age, education level, marital status, gender, BMI, and perceived mental stress. The number of days with URTI was 46% lower when comparing subjects in the highest versus lowest tertile for perceived physical fitness, even after adjustment for confounders.

Regular physical activity may lower rates of infection for other types of diseases, but data are limited due to low disease prevalence. For example, women with a high frequency of walking experienced an 18% lower risk of pneumonia compared with women who walked the least. In the same cohort, women who reported running or jogging more than 2 h per week had a reduced pneumonia risk compared with women who spent no time running or jogging.
Randomized experimental trials provide important data in support of the viewpoint that moderate physical activity reduces URTI symptomatology. In a randomized, controlled study of 36 women (mean age, 35 years), subjects walked briskly for 45-min, five days a week, and experienced one-half the days with URTI symptoms (5.1 vs. 10.8) during the 15-week period compared to that of the sedentary control group.46

The effect of exercise training (five 45-min walking sessions/week at 60%–75% maximum heart rate) and/or moderate energy restriction (1200–1300 kcal per day) on URTI was studied in obese women (n = 91, BMI 33.1 ± 0.6 kg/m²) randomized to one of four groups: control, exercise, diet, exercise and diet.47 Energy restriction had no significant effect on URTI incidence, and subjects from the two exercise groups were contrasted with subjects from the two nonexercise groups. The number of days with URTI for subjects in the exercise groups was reduced 40% relative to the nonexercise groups (5.6 vs. 9.4), similar to the level of nonobese, physically active controls (n = 30, 4.8 days with URTI).

In another study, 30 sedentary elderly women (mean age, 73 years) were assigned to walking or sedentary groups.48,49 The exercise group walked 30–40 min, 5 days per week, for 12 weeks at 60% heart rate reserve. Incidence of URTI in the walking groups was 21% compared to 50% in the calisthenic control group during the study (September–November).

A one-year randomized study of 115 overweight, postmenopausal women showed that regular moderate exercise (166 min per week, ~4 days per week) lowered URTI risk compared to controls (who engaged in a stretching program).50 In the final three months of the study, the risk of colds in the control group was more than threefold that of the exercisers.

### 3.2. Moderate physical activity and enhanced immunosurveillance

During moderate exercise several transient changes occur in the immune system.29,51–53 Moderate exercise increases the recirculation of immunoglobulins, and neutrophils and natural killer cells, two cells that play a critical role in innate immune defenses. Animal data indicate that lung macrophages play an important role in mediating the beneficial effects of moderate exercise on lowered susceptibility to infection.54 Stress hormones, which can suppress immunity, and pro- and anti-inflammatory cytokines, indicative of intense metabolic activity, are not elevated during moderate exercise.29

Although the immune system returns to pre-exercise levels within a few hours after the exercise session is over, each session may represent an improvement in immune surveillance that reduces the risk of infection over the long term. Other exercise-immune related benefits include enhanced antibody-specific responses to vaccinations. For example, several studies indicate that both acute and chronic moderate exercise training improves the body’s antibody response to the influenza vaccine.55–58 In one study, a 45-min moderate exercise bout just before influenza vaccination improved the antibody response.55

These data provide additional evidence that moderate exercise favorably influences overall immune surveillance against pathogens. Taken together, the data on the relationship between moderate exercise, enhanced immunity, and lowered URTI risk are consistent with guidelines urging the general public to engage in near-daily brisk walking.

### 4. Conclusion

Although methodology varies widely and evidence is still emerging59 epidemiologic and randomized exercise training studies consistently report a reduction in URTI incidence or risk of 18%–67%. This is the most important finding that has emerged from exercise immunology studies during the past two decades. Animal and human data indicate that during each exercise bout, transient immune changes take place that over time may improve immunosurveillance against pathogens, thereby reducing URTI risk. The magnitude of reduction in URTI risk with near-daily moderate physical activity exceeds...
levels reported for most medications and supplements, and bolsters public health guidelines urging individuals to be physically active on a regular basis.

Regular physical activity should be combined with other lifestyle strategies to more effectively reduce URTI risk. These strategies include stress management, regular sleep, avoidance of malnutrition, and proper hygiene.\(^{33,60}\) URTI is caused by multiple and diverse pathogens, making it unlikely that a unifying vaccine will be developed.\(^{33}\) Thus lifestyle strategies include stress management, regular sleep, avoidance physically active on a regular basis.

The anti-inflammatory effect of near-daily physical activity may play a key role in many health benefits, including reduced cardiovascular disease, type 2 diabetes, various types of cancer, sarcopenia, and dementia.\(^{9–18}\) This is an exciting area of scientific endeavor, and additional research is needed to determine how immune perturbations during each exercise bout accumulate over time to produce an anti-inflammatory influence. As with URTI, multiple lifestyle approaches to reducing chronic inflammation should be employed with a focus on weight loss, high volume of physical activity, avoidance of smoking, and improved diet quality.

References

1. Shephard RJ. Development of the discipline of exercise immunology. Exerc Immunol Rev 2010;16:194–222.
2. Nieman DC, Henson DA, Smith LL, Utter AC, Vinci DM, Davis JM, et al. Cytokine changes after a marathon race. J Appl Physiol 2001;91:109–14.
3. Nieman DC, Dunke CL, Henson DA, MacAnulty SR, Gross SJ, Lind RH. Muscle damage is linked to cytokine changes following a 160-km race. Brain Behav Immun 2005;19:398–403.
4. Arsenault BJ, Earnest CP, Després JP, Blair SN, Church TS. Obesity, coffee consumption and CRP levels in postmenopausal overweight/obese women: importance of hormone replacement therapy use. Eur J Clin Nutr 2009;63:1419–24.
5. Khansari N, Shakiba Y, Mahmoudi M. Chronic inflammation and oxidative stress as a major cause of age-related diseases and cancer. Recent Pat Inflamm Allergy Drug Discov 2009;3:73–80.
6. Devaraj S, Valleggi S, Siegel D, Jialal I. Role of C-reactive protein in contributing to increased cardiovascular risk in metabolic syndrome. Curr Atheroscler Rep 2010;12:110–8.
7. Pearson TA, Mensah GA, Alexander RW, Anderson JL, Cannon III RO, Criqui M, et al. Markers of inflammation and cardiovascular disease: application to clinical and public health practice: a statement for health-care professionals from the Centers for Disease Control and Prevention and the American Heart Association. Circulation 2003;107:499–511.
8. Dekker MJ, Lee S, Hudson R, Kilpatrick K, Graham TE, Ross R, et al. A exercise intervention without weight loss decreases circulating interleukin-6 in lean and obese men with and without type 2 diabetes mellitus. Metabolism 2007;56:332–8.
9. Hsu FC, Kritchevsky SB, Liu Y, Kanaya A, Newman AB, Perry SE, et al. Association between inflammatory components and physical function in the aging, health, and body composition study: a principal component analysis approach. J Gerontol A Biol Sci Med Sci 2009;64:581–9.
10. Lavoie ME, Rabasa-Lhoret R, Doucet E, Mignaud D, Messier L, Bastard JP, et al. Association between physical activity energy expenditure and inflammatory markers in sedentary overweight and obese women. Int J Obes (Lond) 2010;34:1387–95.
11. Beavers KM, Brinkley TE, Nicklas BJ. Effect of exercise training on chronic inflammation. Clin Chim Acta 2010;411:785–93.
12. Ford ES. Does exercise reduce inflammation? Physical activity and C-reactive protein among U.S. adults. Epidemiology 2002;13:561–8.
13. Borodulin K, Laatikainen T, Salomaa V, Jousilahti P. Associations of leisure time physical activity, self-rated physical fitness, and estimated aerobic fitness with serum C-reactive protein among 3,803 adults. Atherosclerosis 2006;185:381–7.
14. Brooks GC, Blaha MJ, Blumenthal RS. Relation of C-reactive protein to abdominal adiposity. Am J Cardiol 2010;106:56–61.
15. Shanela RA, Nieman DC, Henson DA, Jin F, Knab A, Sha W. Inflammation and oxidative stress are lower in physically fit and active adults. Scand J Med Sci Sports 2011 Aug 18. doi:10.1111/j.1600-0838.2011.01373.x [Epub ahead of print].
16. Thompson DS, Earnest CP, Thompson AM, Priest EL, Rodarte RQ, Saunders T, et al. Exercise without weight loss does not reduce C-reactive protein: the INFLAME study. Med Sci Sports Exerc 2010;42:708–16.
17. Arsenault BJ, Cote M, Cartier A, Lemieux I, Després JP, Ross R. Effect of exercise training on cardiometabolic risk markers among sedentary, but metabolically healthy overweight or obese post-menopausal women with elevated blood pressure. Atherosclerosis 2009;207:530–3.
18. Kelley GA, Kelley KS. Effects of aerobic exercise on C-reactive protein, body composition, and maximum oxygen consumption in adults: a meta-analysis of randomized controlled trials. Metabolism 2006;55:1500–7.
19. Stewart LK, Earnest CP, Blair SN, Church TS. Effects of different doses of physical activity on C-reactive protein among women. Med Sci Sports Exerc 2010;42:701–7.
20. Thompson DS, Markovitch D, Betts J, Mazzotti D, Turner J, Tyrell RM. Time course of changes in inflammatory markers during a 6-mo exercise intervention in sedentary middle-aged men: a randomized-controlled trial. J Appl Physiol 2010;108:769–79.
21. Stewart LK, Flynn MG, Campbell WW, Craig BA, Robinson JP, Timmerman KL, et al. The influence of exercise training on inflammatory cytokines and C-reactive protein. Med Sci Sports Exerc 2007;39:1714–9.
22. Christiansen T, Paulsen SK, Bruun JM, Pedersen SB, Richelsen B. Exercise training versus diet-induced weight-loss on metabolic risk factors and inflammatory markers in obese subjects: a 12-week randomized intervention study. Am J Physiol Endocrinol Metab 2010;298:E824–31.
23. Herder C, Peltonen M, Koenig W, Sutcliffe K, Lindstrom J, Martin S, et al. Anti-inflammatory effect of lifestyle changes in the Finnish Diabetes Prevention Study. J Physiol 2010;581:1433–42.
24. Brandt C, Pedersen BK. The role of exercise-induced myokinase in muscle homeostasis and the defense against chronic diseases. J Biomed Technol 2010;2010:520258 [Epub 2010 Mar 9].
25. Pedersen BK. The diseasome of physical inactivity—and the role of myokinases in muscle—fat cross talk. J Physiol 2009;587( Pt 23):5559–68.
26. Petersen AM, Pedersen BK. The anti-inflammatory effect of exercise. J Appl Physiol 2005;98:1154–62.
27. Ryan AS, Nicklas BJ. Reductions in plasma cytokine levels with weight loss improve insulin sensitivity in overweight and obese postmenopausal women. Diabetes Care 2004;27:1699–705.
28. Ferrier KE, Nestel P, Taylor A, Drew BC, Kingwell BA. Diet but not aerobic exercise training reduces skeletal muscle TNF-alpha in overweight humans. Diabetologia 2004;47:630–7.
29. Nieman DC, Henson DA, Austin MD, Brown VA. The immune response to a 30-minute walk. Med Sci Sports Exerc 2005;37:57–62.
30. Nicklas BJ, Wang X, You T, Lyles MF, Demons J, Easter L, et al. Effect of exercise intensity on abdominal fat loss during calorie restriction in overweight and obese postmenopausal women: a randomized, controlled trial. Am J Clin Nutr 2009;89:1043–52.
31. National Institute of Allergy and Infectious Diseases. The common cold. Available at: http://www.niaid.nih.gov/topics/commoncold [accessed 03.07.2010].
32. Fendrick AM, Monto AS, Nightengale B, Sarnes M. The economic burden of non-influenza-related viral respiratory tract infection in the United States. Arch Intern Med 2003;163:487–94.
33. Monto AS. Epidemiology of viral respiratory infections. Am J Med 2002;112(Suppl 6A):S4–12.
34. Nieman DC. Is infection risk linked to exercise workload? Med Sci Sports Exerc 2000;32(Suppl 7):S406–11.
35. Nieman DC. Immune function responses to ultramarathon race competition. Med Sportiva 2009;13:189–96.
36. Shephard RJ, Kavanagh T, Mertens DJ, Qureshi S, Clark M. Personal health benefits of Masters athletics competition. Br J Sports Med 1995;29:35–40.
37. Davis JM, Kohut ML, Colbert LH, Jackson DA, Ghaffar A, Mayer EP. Exercise, alveolar macrophage function, and susceptibility to respiratory infection. J Appl Physiol 1997;83:1461–6.
38. Sim YJ, Yu S, Yoon KJ, Loiacono CM, Kohut ML. Chronic exercise reduces illness severity, decreases viral load, and results in greater anti-inflammatory effects than acute exercise during influenza infection. J Infect Dis 2009;200:1434–42.
39. Matthews CE, Ockene IS, Freedson PS, Rosal MC, Merriam PA, Hebert JR. Moderate to vigorous physical activity and risk of upper-respiratory tract infection. Med Sci Sports Exerc 2002;34:1242–8.
40. Kostka T, Praczko K. Interrelationship between physical activity, symptomatology of upper respiratory tract infections, and depression in elderly people. Gerontology 2007;53:187–93.
41. Kostka T, Drygas W, Jegier A, Praczko K. Physical activity and upper respiratory tract infections. Int J Sports Med 2008;29:158–62.
42. Fondell E, Lagerros YT, Sundberg CJ, Lekander M, Bärlt Ö, Rothman KJ, et al. Physical activity, stress, and self-reported upper respiratory tract infection. Med Sci Sports Exerc 2011;43:272–9.
43. Nieman DC, Henson DA, Austin MD, Shw A. Upper respiratory tract infection is reduced in physically fit and active adults. Br J Sports Med 2011;45:987–92.
44. Barrett B, Brown R, Munt D, Safdar N, Dye L, Maberry R, et al. The Wisconsin upper respiratory symptom survey is responsive, reliable, and valid. J Clin Epidemiol 2005;58:609–17.
45. Neuman MI, Willett WC, Curhan GC. Physical activity and the risk of community-acquired pneumonia in US women. Am J Med 2010;123:281.e7–281.e11.
46. Nieman DC, Neilsen-Cannarella SL, Markoff PA, Balk-Lamberton AJ, Yang H, Chritton DB, et al. The effects of moderate exercise training on natural killer cells and acute upper respiratory tract infections. Int J Sports Med 1990;11:467–73.
47. Nieman DC, Neilsen-Cannarella SL, Henson DA, Koch AJ, Butterworth DE, Fagoaga OR, et al. Immune response to exercise training and/or energy restriction in obese women. Med Sci Sports Exerc 1998;30:679–86.
48. Nieman DC, Henson DA, Gusewitch G, Warren BJ, Dotson RC, Butterworth DE, et al. Physical activity and immune function in elderly women. Med Sci Sports Exerc 1993;25:823–31.
49. Nieman DC. Immune function. In: Gisolfi CV, Lamb DR, Nadel E, editors. Perspectives in exercise science and sports medicine. Exercise in older adults, vol. 8. Carmel, IN: Cooper Publishing Group; 1995. p. 435–61.
50. Chubak J, McTierman A, Sorensen B, Wener MH, Yasui Y, Velasquez M, et al. Moderate-intensity exercise reduces the incidence of colds among postmenopausal women. Am J Med 2006;119:937–42.
51. Neilsen-Cannarella SL, Nieman DC, Jessen J, Chang L, Gusewitch G, Blix GG, et al. The effects of acute moderate exercise on lymphocyte function and serum immunoglobulins. Int J Sports Med 1999;12:591–8.
52. Nieman DC. Exercise effects on systemic immunity. Immunol Cell Biol 2000;78:496–501.
53. Nieman DC, Neilsen-Cannarella SL. The immune response to exercise. Sem Hematol 1994;31:166–79.
54. Murphy EA, Davis JM, Brown AS, Carmichael MD, Van Rooijen N, Ghaffar A, et al. Role of lung macrophages on susceptibility to respiratory infection following short-term moderate exercise training. Am J Physiol Regul Integr Comp Physiol 2004;287:R1354–8.
55. Edwards KM, Burns VE, Reynolds T, Carroll D, Drayson M, Ring C. Acute stress exposure prior to influenza vaccination enhances antibody response in women. Brain Behav Immun 2006;20:159–68.
56. Kohut ML, Arntson BA, Lee W, Rozeboom K, Yoon KJ, Cunnick JE, et al. Moderate exercise improves antibody response to influenza immunization in older adults. Vaccine 2004;22:2298–306.
57. Kohut ML, Lee W, Martin A, Arntson B, Russell DW, Ekkekakis P, et al. The exercise-induced enhancement of influenza immunity is mediated in part by improvements in psychosocial factors in older adults. Brain Behav Immun 2005;19:357–66.
58. Lowder T, Padgett DA, Woods JA. Moderate exercise early after influenza virus infection reduces the Th1 inflammatory response in lungs of mice. Exerc Immunol Rev 2006;12:97–111.
59. Fondell E, Christensen SE, Bärlt O, Bärlt K. Adherence to the Nordic nutrition recommendations as a measure of a healthy diet and upper respiratory tract infection. Public Health Nutr 2011;14:860–9.
60. Cohen S. Keynote presentation at the eight international congress of behavioral medicine: the Pittsburgh common cold studies: psychosocial predictors of susceptibility to respiratory infectious illness. Int J Behav Med 2005;12:123–31.
61. Spiegel K, Sheridan JF, Van Cauter E. Effect of sleep deprivation on response to immunization. JAMA 2002;288:1471–2.
62. Cohen S, Doyle WJ, Alper CM, Janicki-Deverts D, Turner RB. Sleep habits and susceptibility to the common cold. Arch Intern Med 2009;169:62–7.
63. Keusch GT. The history of nutrition: malnutrition, infection and immunity. J Nutr 2003;133:S336–40.