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Data in Brief





Data Article

Data correlations between gender, cytomegalovirus infection and T cells, NK cells, and soluble immune mediators in elderly humans



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NK cell Ageing Cytomegalovirus T cell Sex factors Vitamin D C-reactive protein Adiponectin IL-15 Sphingosine-1-phosphate Retinol binding protein-4

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ABSTRACT

We describe a cohort of 50 elderly subjects, age at least 70 years. We present gender-specific findings in T lymphocyte markers and soluble immune mediators. We show the correlation between cytomegalovirus infection status with CD56^{dim} NK cell responses to a variety of stimuli and with CD56^{bright}/CD56^{dim} NK cell ratio. We also present the correlation of retinol binding protein (RBP)—4 plasma levels with NK cell responses and we explore the relationship between gender and adiponectin, 25(OH)D (vitamin D), and RBP4 in affecting CD56^{dim} NK cell responses. These data are discussed in Al-Attar et al. (2016) [1].

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Specifications Table

Subject area	Immunology
More specific subject area	Sex differences in immune cells and soluble mediators
Type of data	Tables
How data was acquired	Flow cytometry and enzyme-linked immunosorbent assay
Data format	Analyzed
Experimental factors	Immune cells were analyzed immediately ex vivo or were stimulated in vitro
Experimental features	Correlations and mean differences were calculated
Data source location	Lexington, KY – USA
Data accessibility	Data are within this article

Value of the data

- Researchers need to be aware of the gender differences in NK cell responses to various stimuli.
- Exposure to cytomegalovirus (CMV) affects immune responses from T and NK cells and could therefore be an important factor to consider when performing research in elderly human subjects.
- Levels of soluble plasma immune mediators adiponectin and vitamin D (25(OH)D) which affect NK cell development and activity are higher in elderly women compared to men, and are important factors to consider when studying human NK cells.

1. Data

Enclosed are data concerning T cell markers and subsets found in elderly women and men (Table 1). Also shown is the effect of cytomegalovirus (CMV) infection on CD56^{dim} NK cell responses to a variety of stimuli and on the CD56^{bright}/CD56^{dim} NK cell ratio in blood lymphocytes (Table 2). We present the levels of various plasma immune mediators and their levels in elderly women and men (Table 3). We show how plasma RBP4 level correlates with NK cell responses in vitro (Table 4) and we explore if the interaction between gender and plasma adiponectin,RBP4, and 25(OH)D (vitamin D) affects CD56^{dim} NK cell responses in vitro (Table 5). Full details of the data can be found in Al-Attar et al. [1].

2. Experimental design, materials and methods

Male and female subjects > 70 years were recruited from volunteer donor pools at the University of Kentucky Sanders-Brown Center on Aging and by advertisements. Venous blood from 26 males (age mean \pm standard error of the mean (SEM), 77.8 \pm 0.31, range 70–90 years) and 24 females (age mean \pm SEM, 77.0 \pm 0.91, range 70–85 years) were analyzed between October 2012 and April 2014. Prospective donors were screened by telephone interview to exclude those with conditions previously demonstrated to affect NK cells. Exclusion criteria included hospitalization or serious illness in the prior year, history of immunologic illness (rheumatoid arthritis, systemic lupus, scleroderma, polymyositis, Sjögren's syndrome, transplantation, etc), current use of immunomodulatory medications (e.g., corticosteroids), inability to walk one city block, regular consumption of two or more alcoholic beverages per day (28 g ethanol), diabetes, and a history of cancer within the last 10 years, except non-melanoma skin cancers. Two female subjects (but no male subjects) received hormone replacement therapy, one with topical estrogen and one with topical estrogen, progesterone, and testosterone. The gender differences affecting CD56^{dim} NK cell CD38 density and the CD56^{bright} to CD56^{dim} NK cell ratio were no longer significant when the hormone replacement subjects were excluded from analysis. All other gender differences reported below remained significant after exclusion of these two subjects. Blood samples were obtained from people without acute

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