The effect of influenza vaccine on cytokine production by human mononuclear cells

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Abstract

Background: Seasonal influenza outbursts present a considerable challenge for health providers. Therefore, great efforts are made to vaccinate as many individuals as possible to avoid loss of labor days, unnecessary admissions and complications. The vaccine used in our country during the 2014-2015 seasons contains the following influenza strains, i.e., A/California (H1N1), A/Texas (H3N2) and B/Massachusetts. Vaccinated people react by mobilizing all components of the immune system. The aim of the study was to examine the vaccine effect on human peripheral blood mononuclear cells (PBMC) activation and their capacity for inflammatory cytokine production.

Methods: PBMC were incubated for 24 hrs without or with lipopolysaccharide or for 48 hrs with phorbol meristate acetate and ionomycin to determine the secretion of TNF-α, IL-1β, IL-6, IL-1ra and IL-10, IL-2 and IFN-γ by non-stimulated or stimulated PBMC. Influenza vaccine was added at the onset of cultures at the following concentrations: 0.08 µl/ml, 0.2 µl/ml and 0.5 µl/ml.

Results: Incubation of non-stimulated PBMC with influenza vaccine caused a marked increase in IL-2 and IFN-γ production. The secretion of the other pro- and anti-inflammatory cytokines tested was not affected by incubation with the vaccine.

Conclusions: Considering the increased production of IL-2 and IFN-γ following incubation of PBMC with influenza vaccine viruses and the central role of these cytokines as stimulators of cellular and humoral activity, efforts should be made to vaccinate as many individuals as possible to prevent influenza spreading.

Condensed abstract

The vaccine against influenza used in our country during the 2014-2015 seasons contains the following strains: A/California (H1N1), A/Texas (H3N2) and B/Massachusetts. The study was designed to examine the vaccine effect on capacity for cytokine production by human peripheral blood mononuclear cells (PBMC). PBMC were incubated with influenza vaccine and the secretion of TNF-α, IL-1β, IL-6, IL-1ra, IL-10, IL-2 and IFN-γ was examined. Influenza vaccine caused a marked increase in IL-2 and IFN-γ production. Since these cytokines act as stimulators of cellular and humoral activity, efforts should be made to vaccinate as many individuals as possible to prevent influenza spreading.

Introduction

Seasonal outbursts of influenza are associated with considerable health risks terminating with comorbidity and even mortality. According to Laidlaw, [1] influenza virus epidemics are the cause of about 36,000 annual deaths in the United States. Needless to say, that these influenza epidemics present a serious therapeutic and financial burden to health providers. Since older individuals, immune-compromised and immunosuppressed patients are easy targets to viral invasions, health insurance establishments make great efforts to protect as many individuals as possible by vaccination with inactivated influenza viruses. It is notable that following influenza virus invasion the organism mobilizes various arms of its immune defenses, such as production of specific antibodies, macrophages stimulated for cytokine production, as well as activation of influenza virus specific T cells in order to protect it. It has been shown that a combination between CD8+ T cells with influenza virus-specific non-neutralizing antibodies provides solid protective immunity [1]. Similarly, virus-specific CD4+ T cells, not only exert a direct cytotoxic effect, but also provide help to CD8+ T cells [2,3]. The protective role of the peripheral blood mononuclear cells (PBMC) against influenza viral infections merits attention. The interactions between these cells and influenza viruses triggers several functions such as chemotaxis, phagocytosis, enzyme synthesis, activation of toll-like receptors, inflammatory cytokine generation, to cite a few of them [4-7]. Due to repeated exposure to influenza virus infections, the organism develops strain-specific antibodies that react against circulating viruses which consequently endure antigenic mutations [3]. While the inflammatory response is not a prerogative to immune cells only, it has been reported that these cells, challenged with influenza A virus infections, react by an intensive production of a lengthy list of chemokines and cytokines including TNF-α, IL-1β, IL-6, IL-18 and IFN-α/β, a capacity lesser pronounced by respiratory epithelial cells [8, 9]. Exposure of human macrophages to influenza virus resulted in a marked production of...
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Materials and methods

Cell preparation and culture conditions

Peripheral blood mononuclear cells (PBMC) were separated from venous blood of adult donors (n=20, aged 49.3 ± 6.7 years) by gradient centrifugation using Lymphoprep-1077 (Axis-Shield PoC AS, Oslo, Norway). The cells were washed twice in phosphate buffered saline (PBS) and suspended in RPMI-1640 medium (Biological Industries, Beith Haemek, Israel) containing 1% penicillin, streptomycin and nystatin and supplemented with 10% fetal calf serum (designated as complete medium CM).

Influenza vaccine

Influenza vaccine – Fluarix (split virion, inactivated GlaxoSmithKline Biologicals, Branch of SB Pharma GmbH & Co. KG, Dresden, Germany) containing antigens equivalent to the following subtypes: A/California/7/2009 (H1N1)pdm09-like strain, A/Texas/50/2012 (H3N2)-like strain, B/Massachusetts/02/2012-like strain. This vaccine complies with the WHO recommended strains (Northern Hemisphere) for the season 2014/2015. Each 0.5 ml vaccine contains 15 μg hemagglutinin of each of the recommended strains.

Effect of influenza vaccine on cytokine production

2×10^6/ml of PBMC suspended in CM were incubated in 24 well plates ((Nunc, Roskilde, Denmark) for 24 hrs without or with 20 ng/ml lipopolysaccharide (LPS, E. coli, Sigma) to determine the secretion of TNF-α, IL-1β, IL-6, IL-1ra and IL-10 by non-stimulated or stimulated PBMC. Cells were incubated for 48 hrs without or with 1 μg/ml of phorbol meristate acetate (PMA-Sigma, Israel) and 0.5 μg/ml of ionomycin (Sigma, Israel) to evaluate the spontaneous or stimulated production of IL-2 and IFN-γ. Influenza vaccine was added at the onset of cultures at the following concentrations: 0.08 μl/ml, 0.2 μl/ml and 0.5 μl/ml.

Table 1. Effect of influenza vaccine on pro- and anti-inflammatory cytokine secretion

| Influenza vaccine | Pro-inflammatory Cytokines | Anti-inflammatory Cytokines |
|-------------------|----------------------------|-----------------------------|
|                   | TNF-α, nm/gl               | IL-1β, nm/gl                | IL-6, nm/gl             | IL-10, nm/gl  | IL-1ra, nm/gl |
| 0                 | Spon n=20                  | LPS n=12                    | Spon n=12               | LPS n=16      | Spon n=12     | LPS n=12    |
| 0.08 μl/ml        | 0.33 ± 0.03                | 1.0 ± 0.4                   | 1.6 ± 0.1               | 5.5 ± 0.6     | 11.4 ± 1.8    | 38.1 ± 3.6  |
| 0.2 μl/ml         | 0.32 ± 0.04                | 1.0 ± 0.3                   | 1.7 ± 0.1               | 4.9 ± 0.8     | 11.8 ± 1.9    | 36.2 ± 3.0  |
| 0.5 μl/ml         | 0.35 ± 0.04                | 1.0 ± 0.2                   | 1.9 ± 0.2               | 4.6 ± 0.6*    | 11.8 ± 1.3    | 37.3 ± 2.8  |

Effect of influenza vaccine on the secretion of cytokines by non-stimulated (spontaneous), or LPS-stimulated PBMC. 2 ×106 PBMC were incubated for 24 hrs in the absence (0) or presence of influenza vaccine at concentrations as indicated. The level of the cytokines in the supernatant was tested by ELISA.

Statistical analysis

Data was analyzed using one way ANOVA with repeated measures to analyze the effect of influenza vaccine various concentrations on each cytokine secretion and two tailed paired Student’s t-test to compare between cytokine secretion with each dose of the vaccine with that of the control. The results are expressed as mean ± SEM. P value <0.05 was considered as statistically significant.

Results

Effect of influenza vaccine on pro-inflammatory cytokine production

Table 1 shows that incubation of either non-stimulated or LPS stimulated PBMC with the three concentrations of influenza vaccine, had no significant effect on the production of TNF-α (F3,63=0.012, p=0.99 or F3,43=0.668, p=0.58, respectively), IL-1β (F3,47=0.298, p=0.826 or F3,63=0.717, p=0.55, respectively) or IL-6 (F3,47=0.043, p=0.99 or F3,47=0.701, p=0.56, respectively). However, 5µl/ml influenza vaccine caused 17% reduction in the secretion of IL-1β induced by LPS being statistically significant (p=0.0006) (Table 1).

Effect of influenza vaccine on anti-inflammatory cytokine production

The production of the anti-inflammatory cytokines IL-10 and IL-1ra by non-stimulated or LPS stimulated PBMC was not affected by the addition of influenza vaccine at the three concentrations used in the study (F3,43=0.774, p=0.515 or F3,43=0.205, p=0.89, respectively) for IL-10 and (F3,43=0.638, p=0.594 or F3,43=0.219 p=0.882 respectively) for IL-1ra, (Table 1).

Effect of influenza vaccine on IL-2 and IFN-γ production

A dose-dependent stimulation of IL-2 or IFN-γ generation was found when non-stimulated PBMC were incubated with increased concentrations of influenza vaccine between 0.08 - 0.5μl/ml (F3,55=3.8, p=0.015 or F3,55=3.35, p=0.025, respectively) (Table 2, Figure 1). Incubation of PMA/ionomycine stimulated PBMC with the same

The plates were incubated at 37°C in a humidified atmosphere supplemented with 5% CO2. At the end of the incubation period, the cells were removed by centrifugation and the supernatants were kept at -70°C until assayed for cytokine content.

Cytokine content in the supernatants

The concentration of cytokines in the supernatants was tested using ELISA kits specific for human cytokines (Biosource International, Camarillo, CA) as detailed in the manufacturer’s guide-line. Each kit is specific for one individual cytokine. The detection level of cytokines in each one of the kits was 30 pg/ml.
concentrations of influenza vaccine had no significant effect on either IL-2 production (F(3,47)=2.05, p=0.104) or IFN-γ secretion (F(3,47)=0.78, p=0.512, Table 2). At vaccine concentrations of 0.08, 0.2 and 0.5 µl/ml the spontaneous secretion of IL-2 was enhanced by 20%, 45% and by 75% (p=0.05, p<0.01, p<0.001 respectively) and that of spontaneous production of IFN-γ by 160%, 240% and 340% (p<0.01, p<0.05, p<0.01 respectively).

**Discussion**

Human peripheral blood monocytes unrelatedly of their phenotype react vividly following contact with influenza or other viruses [12]. These cells were found to be increased in numbers in 90% of all patients with probable influenza, with or without concomitant lymphopenia [13]. It has been reported that there is a rapid differentiation of monocytes to interferon producing myeloid dendritic cells following influenza virus infection [14]. Similar results have been obtained by Hou et al., [15] who have achieved a differentiation of human monocytes to dendritic cells after 18 hours of their contact with influenza and other viruses. The results of the present study indicate that interaction between peripheral blood mononuclear cells and the influenza vaccine did not activate both non-stimulated and LPS-stimulated cells for TNF-α, IL-1β and IL-6 production. Similar results were observed when the production of the anti-inflammatory cytokines IL-10 and IL-1ra by PBMC was evaluated. On the other hand, the results depicted a clear stimulatory effect of the influenza viruses on IL-2 and IFN-γ levels produced by the immune cells. IL-2 is a cytokine that plays a central role in immune response. It stimulates cellular as well as humoral reactivity. IL-2 stimulates T cell and NK cell proliferation, survival and activation in vitro [16]. Natural killer cells (NK) play an essential role in the immune response against influenza infection. NK cells secrete a variety of cytokines, IFN-γ among them being particularly important since it initiates macrophages to kill phagocytosed pathogens [17]. At the early stages following influenza infection NK cells contain viral replication obtained by killing of infected cells [18]. IL-2, affecting predominantly T cells, acts a pivotal factor in the immune defense against viral and other invaders. It has been shown that IL-2 regulates T-cell motility, adhesion properties and cell proliferation by enhancing the synthesis of the endogenous thrombospondin-1 [19]. According to Hua et al., [20] the concurrent activity of IL-2/IL-2 receptor alpha pathway induces expression of CD4+ T cells with cytotoxic potential against influenza viruses. In addition IL-2 has been found to be essential in the optimal activation and survival of CD4+ and CD8+ T cells that contribute to the clearance of influenza virus during primary and secondary infection. This effect can be achieved either in cases that the cytokine acts alone or concomitantly with IL-21 and IFN-γ [21, 22]. During the H1N1 influenza outburst in 2009, Sridhar et al., [23] have observed that individuals with higher incidence of CD8+ T cells producing IFN-γ and IL-2 showed better outcome after the infection. The occurrence of CD8+ T cells' expansion after influenza infection has been supported by another study [24]. Following 2009 H1N1 influenza pandemic, Weaver et al., [25] have found that humans responded to the infection by a distinct expression of TNF-α, IFN-γ and IL-2 producing CD4+ T cells observed one month post-infection. Considering the fact that T cells and particularly CD4+ cells are the primary target of viruses' attack in general, the observed in vitro increased production of IL-2 and IFN-γ by PBMC in the present study underlines the importance of these cells in the resistance to influenza virus infection. However, it should be emphasized that immune cells of other types, such as CD14+ and CD16+ monocytes producing TNF-α and IL-6, have been found to increase in number following influenza vaccination [11]. In our hands, there was no alteration in the production of the pro-inflammatory cytokine IL-6, although according to Lauder et al., [26] it has been shown to exert anti-viral responses by modulation of T-cells function. Working with IL-6 deprived mice the authors have observed severe lung pathological changes in the animals concomitant with increased production of the pro-inflammatory cytokines IFN-α and TNF-α. Giamarellos-Bourboulis et al., [27] have examined TNF-α, IL-1β and IL-6 concentrations of healthy volunteers at the beginning of signs of influenza A (H1N1) infection, cases with flu-like syndrome and patients with H1N1 fully developed illness. While TNF-α and IL-1β serum concentrations did not differ between the three groups, IL-6 was found to be elevated in the individuals with flu-like syndrome. Conversely, human macrophages exposed to influenza H1N1 virus did not show increase of IL-1β, TNF-α or IL-6 [6], as has been observed in the present work. One has to keep in mind that the capacity of the immune cells to produce pro-inflammatory cytokines following influenza virus infections depends conspicuously on the pathogen's virulence degree and on the viral load [28]. The increased production of IFN-γ found in the present work is in line with other reports that have delineated that influenza A virus stimulated human macrophages to produce not only IFN-α/β, but also a substantial number of inflammatory cytokines including IL-18, a cytokine capable to further induce IFNγ expression [9,29].

In short, the present work indicates that three viral strains integrated in a seasonal influenza vaccine recommended by the Health

**Table 2. Effect of Influenza vaccine on IL-2 and IFN-γ secretion.**

| Cytokine | IL-2, ng/ml | IFN-γ, ng/ml |
|----------|-------------|--------------|
| Influenza | Spont. | n=14 | PMA/iono | n=12 | Spont. | n=14 | PMA/iono | n=12 |
| 0 | 0.39 ± 0.02 | 23.7 ± 1.9 | 0.37 ± 0.05 | 33.6 ± 2.2 |
| 0.08 µl/ml | 0.35 ± 0.04 | 20.9 ± 1.9 | 0.97 ± 0.25* | 32.1 ± 2.9 |
| 0.2 µl/ml | 0.42 ± 0.05** | 18.0 ± 2.4 | 1.26 ± 0.31* | 31.8 ± 2.0 |
| 0.5 µl/ml | 0.51 ± 0.06*** | 20.6 ± 2.1 | 1.63 ± 0.41** | 34.0 ± 2.2 |

Table 2. Effect of Influenza vaccine on IL-2 and IFN-γ secretion.

Effect of influenza virus on the production of IL-2 and IFN-γ by non-stimulated or PMA/ ionomycin –stimulated PBMC. 2 x 106 PBMC were incubated for 48 hrs in the absence (0) or presence of influenza vaccine at concentrations as indicated. The level of the cytokines in the supernatant was tested by ELISA. Asterisks represent statistically significant difference from cells incubated without influenza vaccine (*p<0.05; **p<0.01; ***p<0.001).

Spont. - spontaneous cytokine production by non-stimulated cells.

PMA/iono (phorbol meristate acetate) stimulated cells. *p<0.05, **p<0.01, ***p<0.001.

![Figure 1. Bessler et al](image)

**Figure 1.** Effect of influenza vaccine on the secretion of IL-2 and IFN-γ by non-stimulated PBMC. 2 x 106 PBMC were incubated for 48 hrs in the presence of influenza vaccine at volumes as indicated. The level of the cytokines in the supernatant was tested by ELISA. Each column represents the mean of 14 experiments. Bars represent SEM. Asterisks represent statistically significant difference from PBMC incubated without influenza vaccine (p=0.05;**p<0.01;***p<0.001).
Authorities in this country promoted in vitro production of both IL-2 and IFN-γ, two cytokines pivotal for the antiviral defense of the organism.

Author disclosure statement

No competing financial interests. The authors declare that they have no competing interests. There were no funding sources for the study.

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