Pharmacological regulation of neutrophil activity and apoptosis.

Contribution to new strategy for modulation of inflammatory processes

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ABSTRACT

Novel strategies of antiinflammatory therapy are based upon pharmacological agents capable to enhance the resolution – i.e. the termination of the beneficial inflammation before it may turn into an adverse chronic stage. In contrast to the current therapy, which antagonises the formation of proinflammatory mediators, the "proresolving" therapy promotes natural antiinflammatory processes. It is likely that several drugs and phytochemicals would act in this way, but this point has not been investigated and thus might be totally overlooked. In this paper, effects of curcumin (diferuloylmethane) were analysed, considering the ability of this natural compound to affect resolution of inflammation through modulation of its important inputs – activity and apoptosis of neutrophils. The presented data indicate that, besides its well-known ability to suppress mechanisms engaged at the onset and progression of inflammation, curcumin could support resolution of inflammation through decreased activity and enhanced apoptosis of neutrophils. This substance decreased the formation of oxidants in neutrophils, both under in vitro conditions and after oral administration to arthritic rats. Moreover, curcumin accelerated spontaneous apoptosis of neutrophils, as indicated by increased externalisation of phosphatidylserine, by intercalation of propidium iodide and by enhanced activity of the executioner caspase-3.

KEY WORDS: curcumin; neutrophils; apoptosis; caspase-3 activity; reactive oxygen species

Introduction

Inflammation is a universal defense response of the organism to infection or to tissue injury. During the inflammatory process, different cytokines, growth, transforming and chemotactic factors are synthesised and several types of cells become activated. Neutrophils (polymorphonuclear leukocytes) are considered to be central cells of acute inflammation. These cells most rapidly reach the site of injury or infection and liberate antimicrobial proteins, proteases and produce reactive oxygen species. Prolonged or excessive liberation of these very effective and toxic substances could intensify the inflammatory process and enhance tissue damage. Therefore activity of neutrophils declines consecutively by the effect of antiinflammatory mediators (IL-10, lipoxins, resolvins, protectins) and programmed death of these cells (apoptosis) is initiated. These processes are essential for the termination (resolution) of the beneficial inflammation (Kohli & Levy, 2009; Serhan et al., 2007). An abnormal, ineffective or absent resolution contributes to tissue damage and to persisting inflammation. Moreover, it is thought to participate in the pathogenesis of chronic inflammatory disorders, such as asthma, rheumatoid arthritis or chronic obstructive pulmonary disease. Therefore novel strategies of the antiinflammatory therapy are based upon pharmacological agents capable to enhance the resolution of inflammation. In contrast to the current therapy, which antagonises the formation of proinflammatory mediators, the "proresolving" therapy promotes natural antiinflammatory processes (Filep & El Kebir, 2009; Hallett et al., 2008; Rossi et al., 2007). It is likely that several drugs and phytochemicals would act in this way, but this point has not been investigated and thus might be totally overlooked. In this paper effects of curcumin (diferuloylmethane) were analysed, considering the ability of this natural compound to affect resolution of inflammation through modulation of its important inputs – activity and apoptosis of neutrophils.
Materials and methods

Chemicals

The highly purified curcumin (diferuloylmethane), lumi-
nol, isoluminol and PMA (phorbol-myristate-acetate) were obtained from Sigma (Steinheim, Germany) and methot-
rexate was from Pliva-Lachema (Brno, Czech Republic).

Human Annexin V/FITC Kit was purchased from Bender
MedSystems (Vienna, Austria), Caspase-Glo 3/7 Assay from
Promega (Madison, Wis, USA) and human purified cas-
pase-3 was from Enzo Life Sciences (Lausen, Switzerland).

Neutrophil activity

Activity of neutrophils was evaluated on the basis of ox-
idant formation measured by the chemiluminescence
method. Under in vitro conditions, isolated human neu-
 trophils were stimulated with 0.05 μM PMA and chemilu-
minescence was enhanced either with isoluminol (extra-
cellular) or with luminol in the presence of extracellular
scavengers – superoxide dismutase and catalase (Nosáľ
et al., 2009). The effect of orally administered curcumin
on neutrophil activity was assessed in arthritic rats, in
compliance with Principles of Laboratory Animal Care
(Bauerová et al., 2009). The study was approved by the
State Veterinary and Food Administration of the Slovak
Republic and by the Ethical Committee at the Institute of
Experimental Pharmacology and Toxicology. Curcumin
(50 mg/kg, daily) or methotrexate (0.5 mg/kg, twice a
week) was administered orally over a period of 28 days
after arthritis induction. Then the number of neutrophils
and the luminol enhanced whole blood chemilumin-
escence (spontaneous or stimulated with 0.01 μM PMA)
were assessed (Drábiková et al., 2009). The presented
data, expressed in RLU (relative light units), are based on
integral values of chemiluminescence over 1800 s
(isolated neutrophils) or 3600 s (whole blood).

Neutrophil apoptosis

Viability of human neutrophils was recorded by flow
cytometry after double staining with Annexin-V conju-
gated with FITC (fluorescein isothiocyanate) and prop-
idium iodide. Only Annexin positive cells were considered
pre-apoptotic cells and double positive cells (Annexin+/ propidium+) were considered late-apoptotic or dead cells.

Caspase-3 activity

Effect of curcumin on caspase-3 activity was evaluated in
cell-free system by the luminescence method, using
Caspase-Glo 3/7 Assay kit and human purified caspase-3
(Perečko et al., 2010). The method is based on caspase
cleavage of Z-DEVD-amino-luciferin substrate and on
the light production resulting from the reaction of amino-
luciferin with luciferase.

Statistical analysis

All values are given as means ± SEM. The statistical sig-
nificance of differences between means was established by
Student’s t-test and p-values below 0.05 were considered
statistically significant.

Results

Under in vitro conditions, curcumin decreased the
concentration of oxidants, produced by activated neu-
traphils both intra- and extracellularly. However its
concentration needed for a significant modulation of the
radicals formed inside neutrophils was 100-times higher
than that effective extracellularly (Table 1). The effect of
curcumin on neutrophil activity was further tested under
in vivo conditions – in arthritic rats. Adjuvant arthritis
was accompanied by an increased number of neutrophils
in blood and by a more pronounced spontaneous and
stimulated chemiluminescence (Table 2). Whereas the
arthritis related alteration in neutrophil count and in
spontaneous chemiluminescence were not modified by
curcumin, the increased reactivity of neutrophils to PMA
was less evident in curcumin treated animals. The effects
of curcumin were similar to those of methotrexate – a
drug widely used in the therapy of arthritic patients.

Curcumin (10 and 100 μM) elevated the percentage of
preapoptotic and apoptotic neutrophils (Figure 1A) and
increased the activity of human recombinant caspase-3
(Figure 1B).

Discussion

Curcumin is a natural antioxidant, displaying remark-
able antiinflammatory and antiarthritic activities (Funk
et al., 2006; Jagetia & Aggarwal, 2007). These beneficial
effects have been attributed to the capacity of curcumin
to prevent activation of nuclear factor-kappa B and the
subsequent overexpression of proinflammatory media-
tors – cytokines, adhesion molecules, cyclooxygenase-2,
phospholipase A2, myeloperoxidase, collagenase, as well
as to its ability to modulate activities of T lymphocytes
and macrophages (Huang et al., 2004; Sharma et al.,
2005). The presented results indicate that, besides sup-
pression of proinflammatory processes, curcumin could
promote resolution of inflammation owing to its capacity

Table 1. Effect of curcumin on neutrophil activity in vitro.

| Curcumin (μM) | Intracellular chemiluminescence | Extracellular chemiluminescence |
|--------------|-------------------------------|-------------------------------|
| 0.01         | 8.18 ± 4.56                   | 9.92 ± 3.82                   |
| 0.1          | 6.93 ± 5.51                   | 15.66 ± 3.84**                |
| 1            | 8.09 ± 6.07                   | 31.03 ± 1.85**                |
| 10           | 81.40 ± 2.43**                | 86.09 ± 2.30**                |
| 100          | 93.88 ± 1.38**                | 99.70 ± 0.04**                |

Formation of reactive oxygen species was measured on the basis of lum-
nol/isoluminol enhanced chemiluminescence in human neutrophils treated
with curcumin and stimulated with PMA. Absolute control values of chemi-
luminescence, expressed in relative light units, were 465.200 ± 75.789 RLU
(intracellular) and 9487833 ± 1125 207 RLU (extracellular). Mean ± SEM,
n = 6, **p<0.01
to accelerate apoptosis of neutrophils and to inhibit oxidative burst of these cells.

As shown by flow cytometric analysis, curcumin enhanced spontaneous apoptosis of neutrophils. This was indicated by an increased number of Annexin positive cells – i.e. cells displaying a more pronounced externalisation of phosphatidylserine. The expression of phosphatidylserine on the external side of plasma membrane facilitates the recognition of apoptotic neutrophils by macrophages and their safe removal from the site of inflammation (Luo & Loison, 2007). At high concentration (100 μM), curcumin increased the number of late apoptotic or dead neutrophils, disclosed on the basis of nucleus staining by propidium iodide. Integrity of these cells was not destroyed by curcumin treatment and thus the discharge of toxic neutrophil contents into the extracellular space was prevented.

The proapoptotic effect was not confirmed under in vivo conditions – by a decreased number of neutrophils in rats treated with curcumin. This might be due to the lower oral bioavailability of curcumin (Aggarwal & Sung, 2008) and to the dosage/period of administration – not sufficient to achieve effective concentration in neutrophils.

The ability of curcumin to increase apoptosis has been extensively studied in cancer cells (Nair et al., 2010). In neutrophils, it was deduced from the altered morphology and from the activation of the executioner caspase-3 in these cells, resulting from the activation of p38 mitogen-activated protein kinase (Hu et al., 2005). Our experiments, performed in cell-free system, indicated that direct interaction with caspase-3, resulting in increased activity of this enzyme, could not be excluded.

Reduced formation of reactive oxygen species in neutrophils may represent another mechanism involved in
the proresolving activity of curcumin. Oxidants released outside neutrophils (potentially dangerous for host tissues) were reduced more effectively than oxidants formed within neutrophils (involved in elimination of phagocytosed pathogens and fulfilling a regulatory role). Compared to previous studies (Limasset et al., 1993; Jackson et al., 2006), the inhibition was observed at a lower concentration of curcumin. Reduced chemiluminescence could result from the well-known scavenging activity of curcumin, which originates from the presence of phenolic and central methylene hydrogens in its molecule (Weber et al., 2005; Pari & Murugan, 2006). Moreover, curcumin was found to decrease phosphorylation of the signalling enzyme protein kinase C (PKC), namely of its two isoforms PKCcα and PKCcβII (Jančinová et al., 2009). Since these isoforms directly participate in the activation of the neutrophil NADPH oxidase (Fontayne et al., 2002), their inhibition could result in reduced oxidant formation and may explain the decreased chemiluminescence observed in the presence of curcumin. The decreasing effect of curcumin on neutrophil activity was further observed under in vivo conditions – in arthritic rats. Neutrophils modified by inflammation produced several times more oxidants than did healthy controls and this hyperreactivity was suppressed by orally administered curcumin. Since the excessive formation of radicals was ascribed to the enhanced activity of NADPH oxidase (Fairhurst et al., 2007; Miesel et al., 1996), the capacity of curcumin to prevent activation of nuclear factor-kappa B and the subsequent expression of proinflammatory mediators (Funk et al., 2006; Huang et al., 2004; Jagietta & Aggarwal, 2007; Sharma et al., 2005) could be involved in this effect.

Curcumin-related inhibition of neutrophil oxidative burst was comparable with the activity of methotrexate – a drug widely used in the therapy of artritic patients, which triggers synthesis of the endogenous anti-inflammatory mediator adenosine (Cronstein, 2005) and is thus considered a “resolution friendly” drug (Serhan et al., 2007).

The presented data indicate that, besides its ability to suppress mechanisms engaged at the onset and progression of inflammation, curcumin could support resolution of inflammation through decreased activity and enhanced apoptosis of neutrophils.

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