Radiation pneumonitis (RP) is defined as an exudative inflammation which occurs as a result of ionizing radiation and it is considered as an alveolitis resulting from irradiation of the lungs with a single dose of 7 Gy or more. It involves changes in the irradiated areas only.

Apoptosis (programmed cell death) is important for cellular homeostasis in a variety of tissues. The mechanisms regulating apoptosis are complex and involve the interaction of nuclear and cytoplasmic proteins. Recently, the sensitivity of tissues to apoptosis induced by a variety of external stimuli has been closely linked to the intracellular concentrations of a family of cell-death regulators, which include Bcl-2. High levels of Bcl-2 protect against apoptosis through inhibition of caspases activation, the key enzymes of apoptosis.

Pentoxifylline (PTX) ameliorates radiation-induced histological changes in irradiated lungs. Funk et al. observed the in vitro influence of cytokine production on PTX-treated cells, which were stimulated by phytohemaglutinin. These results suggest that PTX inhibits the production of TNF-α, IL-2, and IFN-γ, proinflammatory cytokines.

Summary: We measured the number of Bcl-2, apoptotic, neutrophil, and surfactant apoprotein D (SP-D) positive cells in irradiated rat lungs during different time points after the sublethal whole-thorax irradiation of rats. We also investigated the influence of pentoxifylline (PTX) therapy on these markers. Wistar rats were given 15 Gy thoracic irradiation and PTX (35 mg/kg) twice a week. Animals were examined histologically and immunohistochemically at intervals from 1 to 12 weeks. In non-treated rats compared with treated rats, Bcl-2 expression was significantly inhibited from 4 weeks after irradiation. A higher apoptosis presence in non-treated rats from 4 weeks was found and apoptosis development in PTX-treated animals was delayed and started 8 weeks after irradiation. Similar differences were measured during neutrophil granulocytes examination. Neutrophil penetration in non-treated rats was found 5 weeks after irradiation in contrast to the RP onset of PTX-treated animals 8 weeks after irradiation. The number of SP-D positive cells in non-treated rats observed until 5 weeks after irradiation was higher than in the control group. PTX-treated animals expressed higher number of SP-D positive cells during the whole experiment than the control group. We suggest that apoptosis is linked to neutrophil granulocyte actions during the RP onset and that PTX therapy causes diminished inflammation development.

Key words: Irradiation, Lung, Pentoxifylline, Apoptosis, Bcl-2

Abbreviations: RP - radiation pneumonitis; ARDS - adult respiratory distress syndrome; DAD - diffuse alveolar damage; TNF - tumor necrosis factor; IL - interleukin; S.E.M - standard error of mean. PTX - pentoxifylline; PBS - phosphate-buffered saline; cAMP - cyclic adenosine monophosphate; ATP - adenosine triphosphate; DXM - dexamethasone; IFN - interferon; SP-D - surfactant apoprotein D.
with 10% neutral buffered formalin, embedded into paraffin and 4 mm thick tissue sections were cut, stained with chloroacetate esterase to detect neutrophil granulocytes and Gramm’s staining for bacterial infection.

Immunohistochemical examinations for bcl-2, direct detection of apoptotic cells, and surfactant apoprotein D positive cells were performed with a standard immunoperoxidase technique. After blocking endogenous peroxidase activity for 20 min, tissue sections were incubated with individual polyclonal antibodies (anti-bcl-2 rabbit antibody, Santa Cruz, CA, USA) diluted 1:300 in phosphate-buffered saline (PBS, pH 7.2) for 24 h at 4°C, and then washed three times in PBS. The slides for bcl-2 detection were then incubated with a horseradish peroxidase-coupled anti-rabbit antibody (Santa Cruz, CA, USA) for 45 minutes at 37°C. Excess antibodies were washed off with PBS. Finally, a 0.05% 3,3-diaminobenzidine tetrahydrochloride chromogen solution (Sigma, Prague, Czech Republic) in PBS containing 0.02% hydrogen peroxide was added for 10 min to visualise the antigen-antibody complex in situ.

A M30 Cytodeath kit (Roche Diagnostics, Mannheim, Germany) was employed for the detection of apoptotic cells. The staining procedure was as for bcl-2, whereas incubation with the primary antibody was for 1 hour at room temperature and incubation with the mouse secondary horseradish peroxidase-coupled antibody for 30 minutes at room temperature.

Surfactant apoprotein D was detected as with M30 Cyto-death, whereas samples were put in citrate saline (pH 6.0) and twice heated in a microwave (750W) for a period of 5 minutes. Samples were then incubated with the mouse monoclonal anti-surfactant apoprotein D antibody (clone VI F11, from Prof. Michael Kasper, Technische Universität, Dresden, Germany) diluted 1:10 in PBS for 1 h at room temperature.

As negative controls were used samples stained without primary antibodies.

**Measurement of bcl-2, M 30, and SP-D positive cell numbers**

Immunohistochemical samples were evaluated using an IMT-2 light microscope (Olympus Company, Prague) and computer image analysis (Image Pro, Media Cybernetics, MD, USA). Ten randomly selected viewing fields with a size of 10744,32 μm² without bronchi and large vessels from each sample were evaluated at a 600 fold original magnification.

**Data Processing**

The Mann-Whitney test was used for statistical analysis, giving a mean ± 2 x SEM.

**Results**

The following bcl-2 positive cells were found: type-II pneumocytes, fibroblasts, fibrocytes, endothelial cells and macrophages. Epithelial cells of the bronchial tree were also positive but the bronchi were not evaluated. Apoptosis was detected in the epithelium, in type-II pneumocytes and occasionally macrophages (after phagocytosis) and endothelial cells. SP-D positive staining in type-II pneumocytes, macrophages, and unexamined bronchial epithelial cells was observed. No infectious agents were found.

In Group S, 3 animals died, 2 at 4 weeks and 1 at 12 weeks after irradiation. One rat from the PTX-administered group died at 12 weeks after irradiation.

**Table 1: Average number of bcl-2 positive cells ± 2 x SEM in the lungs.**

| Group          | 1 week | 2 weeks | 3 weeks | 4 weeks | 5 weeks | 8 weeks | 12 weeks |
|----------------|--------|---------|---------|---------|---------|---------|----------|
| Group S        | 13.57±1.12 | 18.64±2.36 | 14.7±1.10 | 10.78±2.64 | 15.3±2.72 | 10.8±1.52 | 17.4±2.88 |
| Group PTX 35   | 11.92±2.20 | 17.06±3.32 | 19.5±4.32 | 29.25±2.50 | 34.06±5.52 | 29.5±4.01 | 31.8±2.49 |
| Group C        | 10.3±2.50 | 15.8±3.50 | 13.5±2.50 | 22.4±4.50 | 27.8±4.50 | 22.5±3.50 | 27.8±4.50 |

**Probability of value differences to Group C: 12 weeks after irradiation:** p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.

**Table 2: Average number of apoptotic cells ± 2 x SEM in the lungs.**

| Group          | 1 week | 2 weeks | 3 weeks | 4 weeks | 5 weeks | 8 weeks | 12 weeks |
|----------------|--------|---------|---------|---------|---------|---------|----------|
| Group S        | 0.18±0.12 | 1.6±0.38 | 0.12±0.16 | 4.25±2.04 | 17.05±2.68 | 10.25±1.08 | 8.4±1.92 |
| Group PTX 35   | 3.38±1.08 | 1.67±0.03 | 0.2±0.16 | 0.08±0.12 | 0.02±0.04 | 2.67±1.29 | 3.8±1.28 |
| Group C        | 0.35±0.26 | 0.3±0.26 | 0.3±0.26 | 0.3±0.26 | 0.3±0.26 | 0.3±0.26 | 0.3±0.26 |

**Probability of value differences to Group C: 12 weeks after irradiation:** p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.

**Table 3: Average number of neutrophils in the lungs ± 2 x SEM.**

| Group          | 1 week | 2 weeks | 3 weeks | 4 weeks | 5 weeks | 8 weeks | 12 weeks |
|----------------|--------|---------|---------|---------|---------|---------|----------|
| Group S        | 0.09±0.05 | 0.10±0.10 | 0.11±0.15 | 0.12±0.15 | 1.24±0.34 | 4.80±0.60 | 5.9±0.56  |
| Group PTX 35   | 0.11±0.06 | 0.21±0.06 | 0.1±0.06 | 0.08±0.04 | 0.16±0.06 | 1.96±0.77 | 3.4±0.45  |
| Group C        | 0.46±0.41 | 0.45±0.41 | 0.45±0.41 | 0.45±0.41 | 0.45±0.41 | 0.45±0.41 | 0.45±0.41 |

**Probability of value differences to Group C: p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.**

**Probability of value differences between Group S and Group PTX 35 in the same time interval:** p<0.05 - a; p<0.01 – b; p<0.001 - c.

**Table 4: Average number of SP-D positive cells in the lungs ± 2 x SEM.**

| Group          | 1 week | 2 weeks | 3 weeks | 4 weeks | 5 weeks | 8 weeks | 12 weeks |
|----------------|--------|---------|---------|---------|---------|---------|----------|
| Group S        | 9.0±1.00 | 10.4±0.90 | 10.5±1.93 | 9.8±0.84 | 11.2±0.76 | 9.3±0.83 | 9.7±1.20  |
| Group PTX 35   | 12.8±0.92 | 5.5±0.83 | 14.3±1.40 | 13.3±1.65 | 9.8±0.96 | 9.5±0.83 | 9.7±1.20  |
| Group C        | 6.3±0.89 | 6.3±0.89 | 6.3±0.89 | 6.3±0.89 | 6.3±0.89 | 6.3±0.89 | 6.3±0.89  |

**Probability of value differences to Group C: p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.**

**Probability of value differences between Group S and Group PTX 35 in the same time interval:** p<0.05 - a; p<0.01 – b; p<0.001 - c.

**The bcl-2 expression measurement**

Significantly higher values of Group C compared with Group S were measured 12 weeks after irradiation. The effect of PTX therapy was expressed by a higher number of positive staining cells beyond 4 weeks than in Group S.

**Apopotic cell measurement**

Significantly higher values of this marker of Group C compared with Group S were measured 12 weeks after irradiation. In Group PTX 35, the effect of therapy expressed...
ly confirmed inhibition of TNF-α and other pro-inflammatory cytokines during PTX-therapy. If the hypothesis of RP development via the overexpressed pro-inflammatory cytokine cascade (TNF-α, IL-1) and others is correct (26), then the therapeutic period influencing the expression of cytokines starts immediately after irradiation because an increased production of pro-inflammatory cytokines was found immediately after irradiation.

In this study, we sought to test the effect of PTX administered from day 1 to day 84 (12 weeks) after irradiation and to study the link with bcl-2 expression. We investigated apoptosis by the caspase cleavage product of cytokeratin 18 as well as the number of neutrophils and surfactant apo-protein D positive cells in the irradiated lungs by computer image analysis.

**Material and Methods**

Male Wistar rats (Konaroviec, Czech Republic) aged 8-12 weeks and weighing 150-200 g were given local thoracic irradiation using a 90Co unit (Chisotom Chirana) at a dose rate of 1.0 Gy/min, target distance 1 m. The animals were slightly anaesthetised before irradiation using a mixture of one volume of Rometal (Spofa Company, Prague), 3 volumes of Narkamon (Leciva Company, Prague) and 12 volumes of physiological saline. This solution was injected intratracheally at 10 ml/kg. Local thoracic irradiation was performed in a jig in which a 10 cm thick layer of lead reduced the dose to other parts of the body to around 2-3% of the lung dose (Table 1).

Three groups of rats were used: Group S was given physiological saline only. Group PTX 35 was given pentoxifylline (Pentilin, Krka, 100 mg/5 ml ampoules) diluted with physiological saline and administered at a dose of 35 mg/kg; while Group C involved sham-irradiated controls.

PTX and saline were injected subcutaneously twice a week.

Total number of examined rats was 90. Group C contained 6 non-irradiated rats. An each irradiated group (Group S and PTX 35) included 7 irradiated subgroups by 6 animals in each subgroup. Total number of 84 rats (except group C) was irradiated with 15 Gy and treated 1.2, 3.3, 4.5, 6 and 8 weeks after irradiation, respectively. This radiation dose resulted in significant pneumonitis and fibrosis but no mortality up to 30 weeks, whereas 16 Gy resulted in over 50% mortality 6 weeks after irradiation (30). The first drug administration was given 1 h after irradiation and the last at 2 days before the animals were sacrificed.

**Histologic Examination**

Rats were killed by cervical dislocation and lung histology examined 1, 2, 3, 4, 5, 8, and 12 weeks after irradiation; controls were killed at 12 weeks. Measured marker levels in 6 control rats 12 weeks after irradiation were used as reference values, except in the M30 and bcl-2 examinations. During dissection, the lungs were carefully fixed per trachea

| Tab. 1: Average number of bcl-2 positive cells ± 2 x SEM in the lungs. |
|----------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
|          | 1 week          | 2 weeks         | 3 weeks         | 4 weeks         | 5 weeks         | 8 weeks         |
| Group S  | 13.57±5.12      | 18.65±2.36      | 14.75±1.00      | 10.78±2.64      | 15.30±2.72      | 10.80±1.52      |
| Group PTX 35 | 11.92±2.0b    | 19.70±3.32      | 19.36±3.42      | 29.25±5.20      | 34.08±5.52      | 29.55±4.01      |
| Group C  | 5.38±2.56       | 7.28±1.04       | 3.12±0.54       | 2.67±1.20       | 2.33±1.28       | 0.55±0.26       |

**Probability of value differences to Group C 12 weeks after irradiation:** p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.

**Probability of value differences between Group S and Group PTX 35 in the same time interval:** p<0.05 - a; p<0.01 – b; p<0.001 - c.

**Probability of value differences between Group S and Group C in the same time interval:** p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.

**Probability of value differences between Group PTX 35 and Group C in the same time interval:** p<0.05 - b; p<0.01 - c; p<0.001 - f.

**Probability of value differences between Group S and Group PTX 35 in the same time interval:** p<0.05 - a; p<0.01 – b; p<0.001 - c.

**Probability of value differences between Group PTX 35 and Group C in the same time interval:** p<0.05 - b; p<0.01 - c; p<0.001 - f.

**Probability of value differences between Group S and Group C 12 weeks after irradiation:** p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.

**Probability of value differences between Group S and Group PTX 35 in the same time interval:** p<0.05 - a; p<0.01 – b; p<0.001 - c.

**Probability of value differences between Group PTX 35 and Group C in the same time interval:** p<0.05 - b; p<0.01 - c; p<0.001 - f.

**Probability of value differences between Group S and Group C in the same time interval:** p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.

**Probability of value differences between Group PTX 35 and Group C in the same time interval:** p<0.05 - b; p<0.01 - c; p<0.001 - f.

**Probability of value differences between Group S and Group PTX 35 in the same time interval:** p<0.05 - a; p<0.01 – b; p<0.001 - c.

**Probability of value differences between Group PTX 35 and Group C in the same time interval:** p<0.05 - b; p<0.01 - c; p<0.001 - f.

**Probability of value differences between Group S and Group C 12 weeks after irradiation:** p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.

**Probability of value differences between Group S and Group PTX 35 in the same time interval:** p<0.05 - a; p<0.01 – b; p<0.001 - c.

**Probability of value differences between Group PTX 35 and Group C in the same time interval:** p<0.05 - b; p<0.01 - c; p<0.001 - f.

**Probability of value differences between Group S and Group C in the same time interval:** p<0.05 - 1; p<0.01 - 2; p<0.001 - 3.
Until 5 weeks after irradiation, a significantly higher number of SP-D positive cells were measured in Group S in the lungs of PTX-treated rats during the whole experiment, a higher number of SP-D immuno-reactive cells were detected than in Group C. The effect of PTX therapy was observed in all time intervals, except 2 and 5 weeks after sublethal irradiation.

Discussion

Many of the known etiologies of diffuse alveolar damage, including radiation, may cause DNA damage and thereby induce apoptosis (13,17). In animal models, DAD is associated with the generation of free radicals, which include the superoxide anion (O₂⁻), hydrogen peroxide (H₂O₂), the hydroxyl radical (OH•), and singlet oxygen (O²⁺) (6,23). These free radicals have been shown to induce apoptosis, presumably through damage to DNA (32). Korsmeyer et al. (20) postulated that DNA damage and p53-dependent apoptosis may contribute to the pathogenesis and/or evolution of this disorder.

In our study, the number of apoptotic cells was significantly reduced and delayed by the number of apoptotic cells was found at 1 week and 4 weeks after irradiation to the end of the experiment in comparison with Group S.

The number of neutrophil granulocytes

Fig. 2: Average number of bcl-2 positive cells in the field in the lungs after 15 Gy irradiation. The error bars are in the form of 95% confidence intervals. (Δ Group S - O Group PTX 35, α Group C).

From 5 weeks after irradiation to the end of the experiment a significantly higher number of neutrophils was found in Group S than in Group C. The penetration of neutrophils in irradiated lungs was delayed. A higher number of neutrophils were observed 8 weeks after irradiation, significantly lower than in Group S.

The beneficial effects of repeated applications of PTX, DXM and their combination to sublethally irradiated mice were noted (25). PTX diminishes and/or delays the neutrophil granulocyte penetration through the vascular wall in the lungs and the interstitial edema intensity during the RP onst (24,25). In addition, it has been found in the Paragard-tissue culture model formed by isolated pulmonary cells (28) that PTX reduces the production of oxygen radicals and scavenge free radicals.

The observed apoptosis inhibition during the RP onst phase in PTX-treated rats may have been caused via PTX-induced TNF-α inhibition and subsequently a lower inflammatory response in irradiated lungs. Moreover, PTX-mediated overproduction of CAMP (3) and subsequently ATP (10) after phosphodiesterase inhibition may play an additional role in the establishment of the rapid neotrophil granulocyte chemotaxis. In our model this results of surfactant apoprotein D positive cells show that PTX facilitates surfactant production in our model.

We suggested that single or repeated applications of PTX-exposure to the lungs is linked to inflammation onset after thoracic irradiation. Moreover, PTX therapy would cause a delayed and diminished radiation-induced inflammatory response in the lungs.

Further investigation of the early molecular postradiation pulmonary changes with cell identifications will be the objective of our work.

Conclusion

The reduced and delayed expression of apoptosis in irradiated lungs caused by PTX might be used as an effective tool for inhibition of radiation-induced changes of lungs. The inflammatory process and apoptosis in irradiated lungs are closely related phenomena with a possible bilaterally influencing relationship.

Acknowledgements

We would like to thank Pro. Dr. Michael Kasper for his rat reactive anti-SP-D antibody and we thank Mrs. Sarka Prichová and Hana Burzová for their skillful assistance. This study has been supported by MO 60020398127, MO 0231010006, and FJ MSM 111100005 1 grants.

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The number of SPD positive cells

Until 5 weeks after irradiation, a significantly higher number of SPD-positive cells were measured in Group S in the lungs of PTX-treated rats during the whole experiment, a higher number of SPD immunoreactive cells were detected than in Group C. The effect of PTX therapy was observed in all time intervals, except 2 and 5 weeks after sublethal irradiation.

Discussion

Many of the known etiologies of diffuse alveolar damage, including radiation, may cause DNA damage and thereby induce apoptosis (13,17). In animal models, DAD is associated with the generation of free radicals, which include the superoxide anion (O$_2^-$), hydrogen peroxide (H$_2$O$_2$), the hydroxyl radical (OH•), and singlet oxygen (O$_2$. ($\Delta$)). These free radicals have been shown to induce apoptosis, presumably through damage to DNA (32). Korsmeyer et al. (20) postulated that free radical damage and its potential to induce apoptosis are regulated by the intracellular levels particularly of bcl-2. In this regard, bcl-2 has been localized to mitochondria, nuclear membranes, and the endoplasmic reticulum, which are sites where free radicals are generated (20). In DAD, the absence of bcl-2 expression in type II pneumocytes suggest that there is a role for the effects of free radical-induced apoptosis (14). Guinee et al. (15) have shown that apoptosis in DAD is associated with the expression of p53 and WAF1, suggesting that DNA damage and p53-dependent apoptosis may contribute to the pathogenesis and/or evolution of this disorder.

In our study, we investigated the physiological saline–administered animals, an inhibition of bcl-2 followed by the presence of the apoptotic cells in the next time interval were seen. We assume that the presence of the apoptotic cells early after irradiation has a relationship to the action of the radiation-induced free radicals. Apoptotic cells increased significantly beyond 4 weeks after irradiation, one week before production of the neutrophils. On the other hand, the relationship between inflammation and apoptotic phenomena might be bilateral. Apoptotic cells and bodies are primarily removed by nonprofessional phagocytes, such as type II pneumocytes in the lungs. However, if they are not readily removed by phagocytosis, they undergo “secondary necrosis”. During the secondary necrosis, the lysosomes in type II pneumocytes fuse and form hydrolytic enzymes into the cytoplasm causing further destruction of the internal components including the plasma membrane. Once the plasma membrane loses integrity, the cell lysates, releasing hydrolytic enzymes into the extracellular space and guarantee an inflammatory response with possible tissue damage (12).

At present, PTX is considered to be a preparation with various immunomodulatory effects which were found in vitro in prismatic concentrations as low as 10 µg/ml (9) and 20 mg/kg/d per os in the rat model (27). The main effect of PTX in immune reactions is a decrease in the production of TNF-α (9,22) at the protein and as well as the mRNA level (3). Because of this, we used pentoxifyline for the treatment of post-irradiation apoptotic changes. Ward et al. (31) reported that PTX had only a small beneficial effect on a radiation-induced lung injury in a rat by some indices such as lung wet weight and protein content, but not by hydroxyproline content, the activity of the lung angiotensin converting enzyme and the plasminogen activator. From their results it follows that the PTX effect is not only through the pulmonary endothelial function, but also via other mechanisms.

The beneficial effects of repeated applications of PTX, DXM and their combination to sublethally irradiated mice were noted (25). PTX diminishes and/or delays the neutrophil granulocyte penetration through the vascular wall in the lungs and the interstitial edema intensity during the RP onset (24,25). In addition, it has been found in the Paragrupa-associated tissue culture model formed by isolated pulmonary cells (28) that PTX reduces the production of oxygen radicals and scavenges free radicals.

The observed apoptosis inhibition during the RP onset phase in PTX-treated rats may have been caused via PTX-induced TNF-α inhibition and subsequently a lower inflammatory response in irradiated lungs. Moreover, TNF-α plays a crucial role in the induction of apoptosis in DAD (13,20). Our results of surfactant apoptotic D-positive cells show that PTX facilitates surfactant production in our model.

We suggested that apoptotic type II pneumocytes in our model is linked to inflammation onset after thoracic irradiation. Moreover, PTX therapy would cause a delayed and decreased radiation-induced inflammatory response in the lungs.

Further investigation of the early molecular postradiation pulmonary changes with cell identifications will be the objective of our work.

Conclusion

The reduced and delayed expression of apoptosis in irradiated lungs caused by PTX might be used as an effective tool for inhibition of radiation-induced changes of lungs. The inflammatory process and apoptosis in irradiated lungs are closely-related phenomena with a possible bilateral influence.

Acknowledgements

We would like to thank Prof. Dr. Michael Kasper for his help and thank Drs. Sárka Příhodová and Hana Burková for their skillful assistance. This study has been supported by MŠMT 60020398127, MŠMT 60301100002, and FJ MSMT 111000003 grants.

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Introduction

Continuing search for etiological factors of dyslexia motivated also a lot of recent studies using various methods (PET, functional MRI, electrophysiology of the CNS, psychophysical examination) to verify the hypothesis of the visual magnocellular pathway involvement (5). There are quite new findings supporting the theories about a delayed signal transmission within the magnocellular pathway and/or a decreased activation of the visual association cortex – particularly of the medio-temporal (V5) area which is reported to be specialized for motion processing (4, 13). However, some other results (7, 14) are not in agreement with the opinion that quite large part of the dyslexic people has a deficit in visual functions. Since knowledge of the basis of dyslexia is crucial for strategy of its rehabilitation, we tried to enlarge our previous study (10) of motion related visual evoked potentials (VEPs) specifically testing the magnocellular pathway (1, 9) in dyslexic children. Simultaneously, we have tested whether the suspected visual deficit in dyslexics is attributable to a delay in magnocellular system maturation (according to some reports – e.g. by Bar- nard et al. (2)) and also if it is possible to influence the function of this system with the use of various light wavelengths (reported e.g. by Williams et al. (16) or Solan (11)).

Methods and subjects

Three groups of subjects were used for electrophysiological testing of visual function. We have examined 10 dyslexics (mean age 40.0 ± 1.1 years) from the group of 20 children who have been tested already 4 years ago (10) – group No. 1 in Tab. 1. The second group consisted of 25 dyslexic children (mean age 10.0 ± 1.9 years) randomly selected in the 2nd and 3rd classes of a specialised school for dyslexics. In the third group 7 control subjects (normal readers of the mean age 13.7 ± 1.1 years) from the previous study (10) were repeatedly examined after 4 years interval. All subjects had visual acuity 6/6 or better (with correction if needed).

The same set of VEPs examinations was done in each subject. Transient pattern-reversal visual evoked potentials (PVEPs) were acquired with high contrast (96%) square-wave black and white checkerboard (element size 40’). Two variants of motion-onset VEPs (MVEPs) were used. The first one - linear motion (random order of fundamental directions, velocity 10 deg/s) of low contrast (10%) isolated checks (40’ check size and 120’ check-to-check distances) had grey, green, blue, yellow and pink modifications (equivalent wavelengths to recommendations by Wilkins et al. (15)). Second motion stimulus consisted of low contrast (10%) grey concentric frames with increasing size and motion.