Shortcomings in Treatment of Epilepsy

Jos Utama*, Evi Setiadi*, Wilmar Musram**

INTRODUCTION

As a rule, medical treatment of epileptic patients has been limited to the prescription of common antiepileptic drugs. Only in a small number of cases - mostly in failures - more attention has been paid to secondary conditions, like strenuous and emotional factors, environment and lifestyle; but a real scientific investigation of the nutritional status of the patients has never been done.

Prominent figures in ancient epilepsy, in the beginning of our century, have already stressed the importance of nutritional factors in eliminating epileptic seizures, but the idea of a ketogenic diet (rich in fatty substances), together with a regular lifestyle has never been well formulated.
Recent developments and findings in the field of biotechnology and medical research have stimulated us to take into serious consideration the influences of secondary or external factors in epileptics, next to existing endogenic factors. Environmental factors, strenuous conditions, nutrition, vitamins, neurotransmitters, minerals and other possibilities that could stimulate and enhance the efficacy of the common antiepileptic arsenal became our focus of attention.

Another undeniable fact is that until now only a part of the seizures (approximately 80%) are satisfactorily controlled by antiepileptic drugs. However it is an unstable and dynamic situation, depending upon the optimal condition of the patient.

A relatively small part of the patients seems to be unresponsive to oral antiepileptic treatment, single or in combined doses, and are called refractory or intractable epileptic cases. Only a part of these refractory cases with well defined organic structures could be treated successfully by invasive operative methods.

On the other side, recent experiments have confirmed the key functions of vit. B6 and Ca$^{2+}$ in the process of muscle and cell metabolism, not only clinically but also electrophysiologically in EEG and EMG recordings.

Based on these findings, investigations concerning these accompanying extrinsic factors - like vitamins, Ca$^{2+}$, stress - in epileptics were done, in order to achieve better therapeutic results.

Hypothesis
Shortcomings or failure in treatment of epilepsy are not only related to direct medical treatment but also to secondary extrinsic factors, like patient’s environment, nutritional, educational state and irregular unbalanced lifestyle.

Objectives
To assess the secondary extrinsic factors in epileptic patients, like socio-economic and nutritional status together with the existence of stress factors.

MATERIALS AND METHODS
The material consists of 103 epileptic patients, 46 males and 57 females, ages 11-71 years, selected from the neurological outpatient department of Dr. Cipto Mangunkusumo Hospital, Jakarta. The control group, 101 healthy persons, are recruited from medical students and medical personnel, ages between 18-70 years.

The examination consists of:
Clinical neurology, biochemistry, routine EEG recording and peripheral blood examination while drug monitoring, X-ray or CT scan are only performed in selected cases.

Due to technical factors the procedure is divided into two stages:
Stage I: determination of vit. B1 status in a randomized group of 20 patients.
Stage II: determination of vit. B6, B12, Folic acid and Ca$^{2+}$ status of two groups of patients, consisting of 20 newcomers and 63 chronic epileptics.

The patients are interviewed using a specially prepared check list form, covering personal data together with the patient’s social and educational level and occupation, food consumption and stress factors.

The vit. B1 status is expressed in the Thiamine Pyrophosphate (TPP) effect which is determined by the method of Schouten, a modification of the Dreyfus test. The erythrocyte hemolysate is incubated with and without addition of TPP. The increased transketolase (TK) activity resulting from the addition of TPP is inversely related to thiamine depletion. The TPP effect is the percentage increase of TK activity after addition of TPP.

Vitamin B6 status is determined by the erythrocyte aminotransferase (EAT) activity test and expressed in the Coefficient of EAT activation

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\frac{\text{EAT activity with addition of PLP}}{\text{EAT activity without addition of PLP}}
\]

(PLP, i.e. the ratio PLP (pyridoxal phosphate))

The EAT activity is estimated using the Granutest 25 ASAT Tris Merck Kit no. 12162 and 12165. In vitamin B6 deficiency EAT activity decreases, resulting in a higher CA value. The higher the CA value, the more severe the deficiency.

Plasma vitamin B12 and Folic acid are determined simultaneously using the DPC (Diagnostic Product Corporation) Solid Phase no boil Radioimmunoassay kit.

For the Ca$^{2+}$ determination, venous blood collected in a heparin tube, is aspirated into the ICA2 apparatus (ion selective electrode), providing the Ca$^{2+}$ concentration at the pH of the sample and the converted Ca$^{2+}$ concentration at pH 7.4.

RESULTS
The vitamin and Ca ion status of the epileptic patients are shown in Table 1. Statistical analysis shows a significant difference between the TPP effect of the epileptic patients and the control group (p < 0.001).
The Coefficient of EAT Activation (CA) of the epileptic patients and the control group are compared. Statistical analysis shows a significant difference between both groups (p < 0.001), indicating a vitamin B6 deficiency in the epileptic patients.

Comparing the Ca ion status of the epileptic patients with the references values, a significant Ca ion deficiency is found in the epileptic group.

The vit. B12 status in all the epileptic patients, 20 newcomers and 20 chronic cases is normal, except in one case of chronic epilepsy, treated with luminal 2 x 30 mg daily. The FA status is decreased in 47.4% of the examined patients.

In general we have found no significant differences in the biochemistry values of the chronic epileptic patients and the newcomers, except for a slight decrease in the FA status in the chronic epileptics.

Patients background information obtained by interview is shown in table 2.

DISCUSSION

Although we often do not know what the mechanism of epileptogenesis is, yet continuous research has provided more understanding in experimental epilepsies. Changes in extrinsic factors (environment, ionic concentrations) play an important role on the course of seizure activity next to the intrinsic factors (pace maker cells, synaptic transmission).

Under normal conditions, cell membranes have a resting potential due to the differences in ionic concentration inside and outside the cells. The main factors for sustaining this status of polarization are the selective semipermeability of the cell membrane and the dynamic functions of the ionic pumps. Changes in electrolytes or ionic concentrations of blood plasma, will cause an alteration in electrical activity of the cell membranes, called paroxysmal depolarization shift (PDS). Extracellular Ca\(^{2+}\) and Na\(^{+}\) will enter the cell, while intracellular K\(^{+}\) is shifting outwards. Prolonged depolarization or seizure activity will lead to a dysfunction of the ionic pumps, resulting in cell oedema and irreversible damage.

Apart from control through voltage and repolarizing K\(^{+}\) currents, intracellular Ca\(^{2+}\) accumulation contributes to inactivation of persistent Ca\(^{2+}\) currents. This control is further enhanced by a number of neurotransmitters (NT) such as gamma aminobutyric acid (GABA) and norepinephrine (NE), which depress both persistent and transient Ca\(^{2+}\) currents. GABA is an important inhibitory NT in the human body, dysfunction of GABA will result in destabilization of neuron function.

Pyridoxal-5-phosphate (PLP), the active form of vit. B6, acts as a cofactor in different reactions of amino acid metabolism.

PLP participation in amino acid decarboxylation is essential for the synthesis of NTs like GABA. A decrease or increase of vit. B6 in GABA metabolism should be considered as a possible cause of seizure activity.

Experimental work with PLP injection intra ventricularly, has resulted in the production of tonic clonic seizures in mice.

Kryzhanovski and Shandra have found that a combination of the vitamins nicotinamide, vit. B6 and alfa tocoferol (vit. E) functionally has a much better antiepileptic effect than the isolated vitamins. The enhanced efficacy of the combined treatment with nicotinamide, vit. B6 and vit. E together with an anti epileptic drug, is shown by the fact that a significant smaller dose of the anti epileptic drug is required.

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Table 1. Vitamin and Ca ion status

<table>
<thead>
<tr>
<th></th>
<th>Vit. B1 status (% TPP effect)</th>
<th>Vit. B6 status (activation coefficient)</th>
<th>Blood Ca(^{2+}) at pH 7.4 mmol/L</th>
<th>Plasma Folic acid ng/ml</th>
<th>Plasma vit. B12 pg/ml</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Epileptic patients</strong></td>
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<tr>
<td>newcomers</td>
<td>1.77±0.38 (n=20)</td>
<td>1.13±0.04 (n=20)</td>
<td>4.33±2.91 (n=18)</td>
<td>489±288 (n=18)</td>
<td></td>
</tr>
<tr>
<td>Chronic patients</td>
<td>1.66±0.51 (n=63)</td>
<td>1.12±0.04 (n=63)</td>
<td>4.61±3.32 (n=20)</td>
<td>538±40 (n=20)</td>
<td></td>
</tr>
<tr>
<td>whole group</td>
<td>48.8±26.4 (n=20)</td>
<td>1.69±0.48 (n=83)</td>
<td>4.47±3.05 (n=38)</td>
<td>514±308 (n=38)</td>
<td></td>
</tr>
<tr>
<td><strong>Control group</strong></td>
<td>14.5±12.4 (n=38)</td>
<td>1.27±0.11 (n=81)</td>
<td></td>
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<td></td>
</tr>
<tr>
<td><strong>Reference value range</strong></td>
<td></td>
<td></td>
<td>1.14-1.29</td>
<td>3.0-17</td>
<td>200-950</td>
</tr>
</tbody>
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Note: Values in mean x ± 1 SD
Table 2. Extrinsic factors in epileptics

<table>
<thead>
<tr>
<th>Education</th>
<th>Occupation</th>
<th>Daily Consumption</th>
<th>Stress Factors</th>
<th>Medication and Vit. Supplementation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Elementary</td>
<td>Low income -64%</td>
<td>Milk -10%</td>
<td>Mental + 70%</td>
<td>Phenobarbital -76.7%</td>
</tr>
<tr>
<td>Secondary</td>
<td>(laborers, farmers, mechanics)</td>
<td>Vegetables -50%</td>
<td></td>
<td>Diphantoin -35%</td>
</tr>
<tr>
<td>Academy</td>
<td>Fruit -30%</td>
<td></td>
<td></td>
<td>Carbamazepine -18.3%</td>
</tr>
<tr>
<td>University</td>
<td></td>
<td></td>
<td>Physical</td>
<td>Folic acid -20%</td>
</tr>
</tbody>
</table>

Moderate income-36% (shopkeepers, teachers, office workers) Vit. B complex -10%

Figure 1. Gamma aminobutyric acid synthesis and metabolism

Thus it is clear that unfavourable conditions like a deficiency in Ca²⁺, minerals, vitamins like B6, B1, Folic acid and malnutrition as a whole, will worsen the epileptic state.

Research work concerning the absorption of anti epileptic drugs and intake of food, has proven that carbamazepine absorption, both in healthy persons and patients, is unaffected by food, whether it is rich in fat or rich in protein.¹²

Neglected health care and an irresponsible way of life, will also weaken the body resistance, just like strenuous physical and mental conditions have a direct effect upon NT metabolism, both facilitating and inhibiting. Thus all these facts remind us that stress and other external factors are crucial problems for the epileptic patient.

Beside the extrinsic factors there are still the intrinsic factors to be considered, like the unstable synaptic transmission and the ectopic or pathologic pacemaker cells. An increased abnormal synaptic facilitation can result in seizure activity¹³,¹⁴ due to a lowered Mg²⁺ or to an increased K⁺ concentration. The
use of epileptic stimulants like picrotoxin is a frequent cause of seizures. The fact that seizure-like activity develops when synapticaptic between nerve cells is suppressed, shows that intrinsic membrane characteristics can determine seizure generation.15,16

Work on hippocampus slices have shown that certain cells, called pacemaker cells, in the hippocampus (the commissural associational neurons in the Ammon’s horn) areas CA1 and CA3 and in the brainstem, have the intrinsic capability to generate bursts. However, whether a full blown seizure will develop depends also on the presence of a critical mass of affected neurons in other areas. These so-called pacemaker cells possess the capability of synchronizing the activity of the whole population and the critical mass of affected neurons are recruited into ictal activity by recurrent excitatory collaterals, which is called biofeedback.

An interesting finding is the fact that intrinsic factors in epileptic patients are liable to alterations with the increase in age and with progressive polluting conditions. These clinical and structural changes are clearly observed in all neurophysiological recordings.

It is now generally accepted that both intrinsic as well as extrinsic factors are important milestones in the epileptic patient. An unstable personality with an undernourished condition is a vulnerable situation. We have confirmed the undernourishment and high stress conditions in the epileptic group and are of the opinion that the ideal treatment of epileptic patients is not only a pure medical problem but a well balanced package strategy, covering simultaneously the patients intrinsic and extrinsic shortcomings.

CONCLUSIONS

1. The extrinsic factors of the epileptic patients in the Cipto Mangunkusumo Hospital show a lot of shortcomings.
2. Epileptic patients show a deficiency in vit. B6, B1, Ca²⁺ and Folic acid (FA).
3. No difference in biochemistry values is found between the group of newcomers and the chronic epileptics, except for a slight decrease in the FA status in the chronic epileptics.
4. The combined treatment of extrinsic and intrinsic factors in epileptic patients in the Cipto Mangunkusumo Hospital should be more balanced and more dynamic.
5. An optimal epileptic treatment should not only include an appropriate drug strategy and its compliance but an overall dynamic, well balanced consideration of the whole patient’s lifestyle including dietary requirements.

REFERENCES