Introduction

The significance of cerebrovascular diseases as a public health problem is increasing worldwide due to the aging of the population. Stroke is an important cause of morbidity and mortality and one of the major factors contributing to loss of independence in older adults [1]. Thyroid disorders including subclinical hypo- and hyperthyroidism are common in the general population and their prevalence increases with age, although some studies suggest stability of thyroid function over time [2,3]. Complex pathways may be engaged in the relationship between stroke and thyroid function, reaching much further than the known correlations between hypothyroidism and atherosclerosis or hyperthyroidism and the risk of atrial fibrillation increasing the odds of a cardioembolic stroke [4-6]. Acute diseases, such as stroke, may cause euthyroid sick syndrome or nonthyroidal illness syndrome manifested by low serum triiodothyronine (T3) in individuals without previously diagnosed thyroid disease [7].

Moreover, studies suggests that thyroid status may influence recovery, prognosis, and outcomes of rehabilitation in stroke patients, but published data provide partially inconsistent results [8-12]. Therefore, the complex interrelationships between the thyroid, the heart and the brain remain an area of interest with a potential for future prevention and treatment approaches. Screening for thyroid disorders is currently one of the most popular and widely accessible blood tests, although thyroid hormones alterations are often underdiagnosed and the laboratory results might be biased by a number of factors [13,14]. The aim of the present study was to assess thyroid function in patients hospitalized due to acute ischemic stroke.

Material and Methods

The study was conducted in the neurology department of a tertiary care university hospital in Warsaw, Poland. The study group comprised of 50 consecutive patients (24 females and 26 males) with acute ischemic stroke admitted to the stroke unit. The age range of patients was 35-91 years, but the majority (80%) of patients were aged 60 years and over. Methods included: analysis of the medical documentation in terms of laboratory tests including thyroid hormones as well as coexisting diseases. Patients with preserved verbal communication were additionally...
questioned about history of thyroid diseases. Serum TSH was assessed in a routine diagnostic laboratory work-up in all patients, while serum fT3 and fT4 were available for 31 (64%) of patients.

**Results and Discussion**

The patients were divided into two clinical categories based on their ability to communicate verbally with the research team: Group A: normal consciousness and preserved effective verbal communication (N=35, including 19 F, 16 M, mean age 66.9±17 yr., age range 35-91 yr.), and Group B: poor or none verbal communication due to aphasia or altered mental status (N=15, including 5 F, 10 M, mean age 77.5±11.9 yr., age range 58-91 yr.). The main results of the study are presented in Table 1, including serum TSH, fT3 and fT4 levels with indicated reference ranges. In total, any abnormality of thyroid hormones was found in 8 (16%) of the patients with ischemic stroke and will be discussed below in detail.

**Table 1:** Characteristics of the patients with ischemic stroke including thyroid function and comorbidities.

| Characteristic                | Group A N=35 | Group B N=15 | Group A+B N=50 | Missing data |
|------------------------------|--------------|--------------|----------------|--------------|
| Serum TSH in ref. range 0.27-4.2 mIU/L | 33           | 14           | 47 (94)        | -            |
| Serum TSH above 4.2 mIU/L    | 2            | 0            | 2 (4)          | -            |
| Serum TSH below 0.27 mIU/L   | 0            | 1            | 1 (2)          | -            |
| Serum fT3 in ref. range 3.1-6.8 pmol/L | 21           | 8            | 29 (58)        | 19 (38)      |
| Serum fT3 above 6.8 pmol/L   | 0            | 0            | 0              | -            |
| Serum fT3 below 3.1 pmol/L   | 0            | 2            | 2 (4)          | -            |
| Serum fT4 in ref. range 12-22 pmol/L | 19           | 8            | 27 (54)        | 19 (38)      |
| Serum fT4 above 22 pmol/L    | 1            | 2            | 3 (6)          | -            |
| Serum fT4 below 12 pmol/L    | 1            | 0            | 1 (2)          | -            |
| History of AH                | 22           | 9            | 31 (62)        | -            |
| History of AF                | 5            | 4            | 9 (18)         | -            |
| History of DM                | 6            | 5            | 11 (22)        | -            |
| History of dyslipidemia      | 7            | 3            | 10 (20)        | -            |
| Thyroid disease reported by the patient | 8            | -            | -              | -            |

Serum TSH was normal in the majority (47; 94%) of ischemic stroke patients. In the group A consisting of patients who could answer the question about previous thyroid disease, 8 patients confirmed history of thyroid disorders. Two patients in the group A had serum TSH levels above the limit, indicating possible subclinical hypothyroidism: 7.19 mIU/mL in an 81-year-old female with known thyroid disease, and 4.46 mIU/mL in a 58-year-old female without previous thyroid disorder. It is, however, worth noting that serum TSH in these patients did not exceed the value of 10 mIU/mL, accepted as the threshold of significant abnormality warranting treatment. Additionally, in the group A, two patients had serum fT4 levels slightly above the upper limit with normal values of TSH and fT3, none of them reported previous thyroid disorders.

Among patients in the group B who were not capable of responding to the question about previous thyroid disease, one case of hyperthyroidism was found in an 83-year-old female with known thyroid disease, and 4.46 mIU/mL in a 58-year-old female without previous thyroid disorder. It is, however, worth noting that serum TSH in these patients did not exceed the value of 10 mIU/mL, accepted as the threshold of significant abnormality warranting treatment. Additionally, in the group A, two patients had serum fT4 levels slightly above the upper limit with normal values of TSH and fT3, none of them reported previous thyroid disorders.

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Only one in eight patients reporting previous thyroid disease had abnormal value of TSH. It is, however, worth mentioning that brain research studies indicate that even treated hypothyroidism is a risk factor for significant cerebrovascular pathology [15]. Moreover, low T3 has been shown to correlate with a long-term post-stroke cognitive impairment [16]. In the present study, over half of the patients had the history of hypertension and one in five was previously diagnosed with diabetes, atrial fibrillation and/or dyslipidemia, commonly known risk factors for ischemic stroke (Table 1). A new consensus on cardiovascular care for stroke patients of the European Society of Cardiology underlines the role of interdisciplinary approach [17]. It is, therefore, deeply disappointing that this document does not mention the importance of thyroid function assessment.

In patients with stroke, various additional factors might influence thyroid function including iodinated contrast brain imaging, acute therapy of cardiovascular complications (e.g., amiodarone) or use of low molecular weight heparin preparation potentially causing abnormal thyroid tests [18-20]. Pre-admission communication (N=35, including 19 F, 16 M, mean age 66.9±17 yr., age range 35-91 yr.), and Group B: poor or none verbal communication due to aphasia or altered mental status (N=15, including 5 F, 10 M, mean age 77.5±11.9 yr., age range 58-91 yr). The main results of the study are presented in Table 1, including serum TSH, fT3 and fT4 levels with indicated reference ranges. In total, any abnormality of thyroid hormones was found in 8 (16%) of the patients with ischemic stroke and will be discussed below in detail.
medication lists and hospital treatment were not analyzed in the current study. The strengths of the present study include the assessment of consecutive patients with ischemic stroke, thus avoiding selection bias as well as availability of serum TSH level tested as a routine procedure upon admission to a stroke unit. Limitations of our study include relatively small number of patients, single assessment of thyroid hormones, missing data for fT3 and fT4 in 38% of patients, and lack of analysis of medications.

Conclusion

The majority of patients hospitalized with the diagnosis of acute ischemic stroke have normal thyroid function, but approximately one in six patients requires further assessment. Thyroid function and patients’ awareness of thyroid health should be assessed in stroke patients. Follow-up of thyroid function should be carefully planned in selected patients, as diagnostic procedures may increase the risk of thyroid disorders.

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