Changes of Thyroid Hormones and Their Binding Proteins during Plasma Exchange for Polyneuropathy in a Patient with Substituted Hypothyroidism due to Hashimoto's Thyroiditis

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Keywords

Thyroid hormones · Plasma exchange · Substitution · Hypothyroidism

Summary

Background: Pharmacodynamic studies and data concerning adaptation of thyroid substitution in patients with substituted hypothyroidism during plasma exchange (PE) is not available. **Case Report:** We measured TSH, fT₃ and fT₄, total T₄, thyroxin binding globulin (TBG), and albumin before and after 5 PE procedures in a 37-year-old woman who underwent PE for a therapy-resistant polyneuropathy. Thyroxin was increased empirically by 8% resulting in a dose of 1.95 µg/kg per day. **Results:** Despite larger reductions of total T₄ and TBG over a series of 5 PEs (40–50% from baseline), only small reductions of 8% in fT₃ and fT₄ concentrations were documented with a concomitant increase in TSH level. Changes of fT₄, fT₃, and TSH remained within normal range. **Conclusions:** i) Despite a significant decrease in total thyroid hormone pool following PE, fT₄, fT₃, and TSH concentrations changed only slightly. ii) Based on this observation, a general increase in thyroid replacement therapy before PE cannot be recommended, but considered in case of a high normal TSH level.

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Introduction

Plasma exchange (PE) is the most commonly performed therapeutic apheresis procedure, according to data from international registries [1]. The basic premise of PE is that the removal of pathological or pathologically elevated substances will reduce further damage and may permit reversal of the pathologic process [2]. PE is recommended in several indications [3]. Primary hypothyroidism, mainly Hashimoto's autoimmune thyroiditis, is a frequent disorder with a prevalence of 0.3% for clinical and 4.3% for subclinical hypothyroidism [4]. Thyroxin (T₄) is mainly bound to thyroxin binding protein (TBG), albumin, and to a smaller extent to transthyretin. PE removes these binding proteins resulting in major changes of the total hormone pool (TT₄). In contrast to most other hormones, like insulin, cortisol and sex hormones that are rapidly cleared from the circulation and re-secreted again if needed, T₄ has an especially prolonged half-life time of around 7 days, a small fraction of free hormone (free thyroxine (fT₄) around 0.03%) and a quite stable plasma level throughout the day. In addition to the shift from bound to freely available thyroid hormone, the pituitary-thyroid axis is thought to compensate for a PE-induced reduction in TT₄ in a physiological condition [5]. However, pharmacodynamic studies or data about an adaptation of a given thyroid replacement therapy in view of a PE in a patient with hypothyroidism are currently not available.

We, therefore, aimed at investigating the effect of PE on thyroid hormone metabolism in a patient with therapy-resistant polyneuropathy who was treated for primary hypothyroidism (Hashimoto's thyroiditis).

Table 1. Immediate and steady-state changes of thyroid function tests and related binding proteins during 5 courses of PE and their related changes

	Immediate changes* (n = 5)		Steady-state changes** (n = 4)	
	absolute	relative	absolute	relative
TSH, mU/l	-0.37 ± 0.30	-45.9 ± 6.3%	0.41 ± 0.48	37.8 ± 25.0%
fT ₄ , pmol/l	-1.4 ± 1.0	-7.9 ± 5.7%	-0.75 ± 1.6	-4.5 ± 9.0%
TT ₄ , nmol/l	-36 ± 8.3	-45.4 ± 6.6%	-4.5 ± 10.5	-6.1 ± 14.4%
fT ₄ /TT ₄ , %	0.0183 ± 0.0082	80.3 ± 34.4%	0.00025 ± 0.0024	0.76 ± 10.1%
fT ₃ , pmol/l	-0.29 ± 0.12	-6.9 ± 2.1%	-0.33 ± 0.08	-8.5 ± 2.1%
TBG, mg/l	-11.0 ± 2.35	-62.0 ± 7.4%	-0.25 ± 4.35	-4.0 ± 28.4%
Albumin, g/l	-0.2 ± 3.1	-0.3 ± 8.2%	0.75 ± 1.5	1.9 ± 3.8%

*Immediate changes were calculated from pre to post plasma exchange.
**Steady-state values were calculated from pre plasma exchange up to the next pre plasma exchange (i.e. from full circle to full circle in fig. 1)

Case Report

An informed consent form was signed by the patient approving the use of material or information for scientific purposes.

We present the case of a 37-year-old woman with a severe painful peripheral polyneuropathy for 3 years. The etiology could not be determined although a chronic inflammatory demyelinating polyneuropathy was suspected. An initial treatment with oral steroids did not relieve symptoms. Also a therapy with intravenous immunoglobulins did not result in satisfactory symptomatic relief. Due to persistent symptoms a series of PEs was planned. The patient was known for primary hypothyroidism due to Hashimoto's thyroiditis since the age of 10 with documented elevated TPO antibodies. She was on stable thyroid replacement therapy.

Methods

PE Procedures

The mobile centrifugal apheresis system Spectra Optia (TerumoBCT, Lake-wood, CO, USA) was used. Procedures were conducted by qualified nurses supervised by a trained physician. Within 14 days a total of 5 PEs were performed every 2–5 days. Each time the 1.2-fold of the patient's own plasma volume was replaced using a 5% albumin solution (CSL Behring AG, Bern, Switzerland) containing at least 96% albumin according to the manufacturer and physiologic saline (Sintetica SA, Couvet, Switzerland) in a ratio of 2:1.

We used citrate (ACD-A, Bichsel SA, Interlaken, Switzerland) as anticoagulant, following the manufacturers' instructions. Routinely, an intravenous continuous infusion of calcium chloride (Calcium-Sandoz 10%, Sandoz AG, Risch, Switzerland) throughout the PE was performed. The initial infusion rate corresponded to the administration of median 0.25 mg of ionized calcium (Ca²⁺) per milliliter of ACD-A (0.53 mmol Ca²⁺ / 10 mmol citrate) [6].

Substitution

The patient presented with a thyroid-stimulating hormone (TSH) level of 3.36 mU/l at baseline under a stable levothyroxine dose of 129 µg/day (1.81 µg/kg/day). As we anticipated a loss during PE, we empirically increased the substitution dose to 139 µg/day (+8%; 1.95 µg/kg/day) 3 weeks before the first PE. After the last PE the patient returned to her usual dose.

Laboratory Samples

Plasma samples were drawn before and immediately after PE. TSH, fT₃ (free liothyronine), and fT₄ were determined using electro-chemiluminescence immunoassay (Modular E170; F. Hoffmann-La Roche AG, Basel, Switzerland), and for albumin the bromocresol purple method was used (Modular P800; F. Hoffmann-La Roche AG) using routine laboratory. TT₄ and TBG was determined using the Chemiluminescent Microparticle Immunoassay, Architect

(Abbott Laboratories, Abbott Park, IL, USA) and a radio immunoassay, respectively, in an external laboratory (Laboratoire chimie clinique, Endocrinologie, Centre hospitalier universitaire Vaudois, Lausanne).

Statistical Analysis

Due to the small sample size only descriptive methods (arithmetic mean) for changes are reported. We calculated immediate differences (value post-PE minus value pre-PE) of all 5 procedures (both in absolute and relative terms, the mean ± SD of 5 plasmaphereses is given). The relative immediate differences reflect real removal and were calculated as: $100 - ((\text{value post-PE} \times 100) / \text{value pre-PE})$. Assuming a one-compartment model, a predicted removal can be calculated as: $100 - (((\text{value pre-PE} \times e^{-x}) \times 100) / \text{value pre-PE})$, where x is the plasma exchanged [7].

A new steady-state condition was assumed to be reflected by the values measured before each PE. We, therefore, calculated differences from the beginning of one PE to the next. The mean (± SD) of 4 steady-state differences in absolute and relative terms is reported. In order to determine whether or not the number of PE impacts on the different parameters of thyroid hormone metabolism the values before the first and the last procedure were considered.

Results

Clinical Course

Clinically, a focused history did not reveal any symptoms of a dysthyroid condition throughout the procedures. In electroneurography studies, minor improvements of polyneuropathy parameters could be detected. However, the patient reported no subjective improvement after the 5 PEs. The treatment was, therefore, discontinued after 5 procedures.

Laboratory Values

The minor dose increase of thyroid replacement therapy 3 weeks before the first PE resulted in a reduction of TSH levels slightly below the lowest limit of normal range (0.22 mU/l; normal range 0.27–4.5 mU/l). The changes of the thyroid function tests and the related binding proteins are shown in figure 1. Before the first and the last PE the TSH increased from 0.22 to 1.87 mU/l, fT₄ decreased from 19.7 to 16.7 pmol/l and fT₃ from 4.78 to 3.45 pmol/l (all within normal range). TT₄ was 95 nmol/l before the 1st and 77 nmol/l before the 5th PE (normal range 60–150 nmol/l). TBG completely recovered after each PE. Albumin remained constant, as it was substituted.

Fig. 1. Thyroid function tests and related binding proteins across 5 PEs in a patient with substituted primary hypothyroidism. Dashed lines indicate normal reference values. Full circles indicate values before and hollow circles values post plasma exchange.

In table 1 the PE-induced immediate changes are compared with steady-state changes. Assuming a one-compartment model, a removal of 69.9% would be expected. Total T₄ fell on average by $45 \pm 6.3\%$ during each PE, reflecting the reduction of TBG by $62 \pm 7.4\%$. The relative decrease in TSH concentrations was similar ($45.9 \pm 6.3\%$). The free hormones were only reduced by $7.9 \pm 5.7\%$ for fT₄, and by $-6.9 \pm 2.1\%$ for fT₃. Consequently, the ratio of fT₄ to TT₄ increased significantly.

In the steady state the TBG changes were largely abolished ($-6.1 \pm 14.4\%$), as were TT4 changes ($-4.5 \pm 9.0\%$), but for TSH an increase was found ($37.8 \pm 25.0\%$). The reductions in free hormone concentrations were comparable to the immediate changes ($-4.5 \pm 9.0\%$ for fT₄, $-8.5 \pm 2.1\%$ for fT₃). The free-to-bound ratio remained constant over the course of the PE.

Discussion

The main results from this report are three-fold: i) A PE induces an immediate decrease in TT₄ and TBG below normal range that recovers and stabilizes in the lower half of normal range after 5 consecutive procedures. ii) Five PE within 14 days resulted in a steady decrease in fT₄ and fT₃ concentrations and an increase in TSH levels, albeit all within normal range. iii) Based on these findings, an increase in the dose of thyroid replacement therapy before PE cannot be generally recommended. A thyroid replacement dose inducing a TSH in the lower normal range can be considered before starting the first PE.

Acute changes of thyroid parameters included a decrease in TT₄ pool in parallel with a decrease in TBG. Interestingly, the values

immediately after the PE were lower than the normal range. This is not surprising since 99.97% of T_4 is bound, with about three quarters bound to TBG. In fact, TT_4 relates to the 3,400-fold of the free hormone pool. The removal of TBG (–62%) was close to the expected value (–69.9%), whereas TT_4 was somewhat lower (–45%). The lower than expected values could be due to TBG and T_4 included in the albumin preparation, a similar effect has been shown for haptoglobin [7]. In contrast, fT_4 and fT_3 decreased only slightly – far off from a logarithmical removal – indicating a shift from the bound T_4 pool to the free T_4 fraction. This is reflected by an increase in the fT_4/TT_4 ratio and may explain the lack of significant clinical symptoms and signs throughout the procedures. There is a tendency for an increase in TT_4 during the last PE, while TBG remained stable throughout the procedures. Apart from a possible increase in production rate of TBG, possibly there was also a reverse flow of extravascular T_4 as shown in a rabbit model [8].

PE has been described as an effective method of last therapeutic resort for therapy-resistant thyrotoxicosis [9] and thyroxine intoxication [10]. In both situations substantially higher fT_4 levels are reported based on an enhanced endogenous production or intake. Due to the ongoing endogenous production, PE is rather considered a temporal relief therapy before definitive surgery. Taking in account the current observation of only small reductions in fT_4 and fT_3 and only insignificant reductions of fT_4 in one retrospective study [9] following PE, the utility of such a procedure has to be questioned, although it may be a life-saving procedure for thyroid storm in the case of intoxication [11].

There is a steady decrease in fT_4 and fT_3 with a concomitant increase in TSH after 5 procedures, albeit within normal range. This indicates that – in contrast to TT_4 and TGB – the compensatory mechanisms are not capable to guarantee completely stable fT_4 and fT_3 levels throughout several PEs, and more than 5 PEs may eventually result in a hypothyroid condition. Compensatory mechanisms include the shift from bound to free thyroid hormone pool and/or TSH-stimulated production of thyroid hormone. The current study was not designed to differentiate between these mechanisms. However, the observation of an increase in the fT_4/TT_4 ratio and TSH levels suggest that both mechanisms were probably involved. Interestingly, in euthyroid patients without Hashimoto's thyroiditis, PE was reported not to result in significant changes of free hormones despite the fact that similar changes for TT_4 and TBG were observed [5].

Limitations

The main limitation of this report is that it is restricted to one subject with only 5 instances of PE. Clinically more relevant changes may become apparent in patients with more than 5 procedures.

Strength

The setting has been highly standardized with a sufficient temporal resolution and additional measurement of clinically now rarely used parameters.

Recommendation

Based on these data it appears reasonable to aim for a TSH level in the lower normal range before plasma exchange is initiated. At least for 5 consecutive PEs, a general increase in thyroid substitution dose cannot be recommended. However, if more than 5 procedures are planned in a series, an adaption of thyroid hormone replacement dose may be considered. After plasma exchange the original substitution regimen can be re-adopted. Frequent measurements of TSH, fT_3 , and fT_4 are mandatory to further adapt the substitution dose.

Conclusion

Despite a significant decrease in TT_4 following plasma exchange, fT_4 , fT_3 and TSH concentrations changed only slightly.

Based on this observation, a general increase in thyroid replacement therapy before plasma exchange cannot be recommended.

An increase in substitution dose can be considered in case of a high normal TSH level.

Plasma exchange in patients with substituted hypothyroidism may serve as a system's perturbation tool to investigate the physiology of thyroid hormones and their binding proteins. Results from such observations may help quantify expected losses for future patients.

Disclosure Statement

The authors declare that they do not have conflicts of interest.

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