Functional and metabolic adaptation of the heart to prolonged thyroid hormone treatment

H. DEGENS, A. J. GILDE, M. LINDHOUT, P. H. M. WILLEMSEN, G. J. VAN DER VUSSE, AND M. VAN BILSEN

Department of Physiology, Cardiovascular Research Institute Maastricht, Maastricht University, 6200 MD Maastricht, The Netherlands

Submitted 2 April 2002; accepted in final form 5 August 2002

Development of heart failure is accompanied by a variety of neuroendocrine changes. Recently, cardiac failure was shown to be associated with both a decline in circulating thyroid hormone (TH) levels (18, 24) and altered cardiac TH signaling, as evidenced by changes in myocardial expression of TH receptor isoforms (21, 22). As a result, a state of relative hypothyroidism may ensue, which has been held responsible for the decline in expression of TH responsive genes during hypertrophy and failure. These include the genes encoding α-myosin heavy chain (α-MHC) and sarcoplasmic reticulum Ca2+-ATPase (SERCA2a), both of which are important determinants of cardiac function (11, 38). The observation that short-term TH administration improves cardiac performance, both in animal models of cardiac dysfunction and in patients suffering from cardiac failure (11, 18, 24, 29, 31), agrees with this notion. This beneficial effect of TH was indeed associated with an elevation of α-MHC mRNA and protein levels. The effect on cardiac SERCA2a mRNA content was less consistent (11, 31). One should realize, however, that hyperthyroidism itself is often associated with impaired cardiac function (24, 35, 45) and that in rats TH treatment of myocardial infarction only transiently improves cardiac performance (28). This discrepancy may result from the duration of exposure to elevated plasma concentrations of TH. So far, detailed information on the nature of the cardiac adaptive response to TH supplementation as a function of time is virtually lacking.

In hyperthyroidism, cardiac hypertrophy is accompanied by an overall increase in metabolic rate and enhanced lipolysis (19). The absence of hypertrophy in heterotopic cardiac transplants in TH-treated rats suggests that cardiac hypertrophy does not result from direct effects of TH on cardiac muscle (23). The hypertrophic response rather results from the hyperdynamic circulatory state as a consequence of the enhanced metabolic rate, increased blood volume, and decreased peripheral resistance (23, 45). Each of these factors potentially increases myocardial energy demand. Accordingly, it has been reported that cardiac glucose metabolism (34) and fatty acid oxidation (37, 39) are enhanced in hyperthyroid animals. On the other hand, during pressure and volume overload-induced cardiac hypertrophy, cardiac substrate metabolism shifts from fatty acids to glucose (1), possibly due to a marked reduction in the expression of β-oxidation enzymes (4, 33, 40). This raises the question as to whether changes in cardiac energy metabolism as a consequence of TH supplementation are also associated with changes in the expression of genes involved in substrate handling, and if so, whether such changes depend on the duration of TH supplementation and the extent of cardiac hypertrophy.

The costs of publication of this article were defrayed in part by the payment of page charges. The article must therefore be hereby marked “advertisement” in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.
The main goal of this study was to investigate the time-related effects of elevated circulating TH levels on the development of cardiac hypertrophy, performance, and energy metabolism in rats. As a measure of cardiac performance, left ventricular (LV) ejection fraction (EF) was determined by means of echocardiography. MHC isoform distribution, collagen content, and SERCA2a and atrial natriuretic factor (ANF) mRNA levels were assessed to delineate the hypertrophic phenotype. Furthermore, TH-induced changes in cardiac metabolism were assessed at the biochemical and molecular level by measuring fatty acid oxidation and glycolytic capacity in homogenates as well as mRNA levels of genes involved in fatty acid and glucose uptake and metabolism. In addition, the mRNA levels of the uncoupling proteins (UCP2 and -3) were determined because these proteins may be responsible for the decrease in mitochondrial efficiency during hyperthyroidism (9). Finally, the expression of peroxisome proliferator-activated receptor-α (PPAR-α) was determined because this transcription factor is considered to play a pivotal role in the regulation of the expression of genes involved in cardiac lipid metabolism (44). Moreover, one study (5) reported that its expression was diminished in overload-induced cardiac hypertrophy.

Collectively, the present findings indicate that TH exposure, although beneficial on a short-term basis, may become detrimental for the heart when continued for longer time intervals.

MATERIALS AND METHODS

Animals. At the start of the experiments, male Wistar rats (198 ± 13 g) were 7 wk old. Rats, two per cage, were kept at a 12:12-h light-dark cycle. Food (25% protein, 6% fat, and 38% carbohydrates; Hope Farms; Woerden, The Netherlands) and water were provided ad libitum. The rats were randomly assigned to TH-treated and corresponding sham groups. Body weights (BW) were determined weekly. TH rats daily received subcutaneous injections of 3,3',5-triiodo-L-thyronine (200 μg/kg BW) for 3, 7, or 28 days. TH treatment started 28, 7, or 3 days, respectively, before the terminal experiment. Accordingly, all animals were the same age (11 wk) at the time of death. TH (40 μg/ml) was dissolved in 1 mM NaOH, 0.9% NaCl. Sham rats received subcutaneous injections daily of the solvent at the same volume. All procedures were approved by the local Committee on Animal Experimentation of Maastricht University.

Echocardiography. In a subset of animals, echocardiography was performed at weekly intervals after initiation of TH or vehicle administration. The first measurements were performed within 6 h after the first injection of TH or vehicle. For echocardiography, the rats were anesthetized with pentobarbital sodium (50 mg/kg ip). M-mode images obtained from the short axis of the heart at the level of the papillary muscles with a 10-MHz probe (model LA14; Esaote Biomedical). From this image, heart rate (HR) was calculated. LV free wall thickness during diastole was determined and the LV EF measured from the internal ventricular diameter during diastole and systole. The ratio of LV free wall thickness to internal LV diameter during diastole was used as a measure of eccentric or concentric hypertrophy.

Terminal experiment. At the end of TH treatment, the rats were anesthetized with pentobarbital sodium (60 mg/kg ip). After thoracotomy, the heart was exposed and 1-ml blood samples were withdrawn from the LV cavity with a syringe containing 100 μl 3.8% sodium citrate. After 30 min, the blood samples were spun for 5 min at 6,000 revolutions/min on a tabletop centrifuge, and the supernatants were collected and stored at −80°C until use. Immediately after blood sampling, the heart was excised and adherent blood was removed in ice-cold saline. The heart was blotted dry and heart weight (HW) determined. After excision of the heart left tibia length (TL) was measured ex situ with a pair of calipers. The HW/TL ratio was used to assess the degree of cardiac hypertrophy because this ratio takes into account differences in BW, body composition, and/or growth that occur during TH treatment. After the atria were removed, a 3-mm thick cross-sectional slice of the entire base of the heart was placed on cork and frozen in liquid nitrogen for histological analysis. The right ventricle was then separated from the LV and septum, and both parts were frozen in liquid nitrogen. All tissue samples were stored at −80°C until RNA and protein extraction.

Metabolic capacity. In another subset of animals, a 5% homogenerate of the LV was made on ice in a buffer composed of 0.25 M sucrose, 2 mM EDTA, and 10 mM Tris (pH 7.4) for the measurement of palmitate oxidation and glycolytic rate. We choose homogenates because they allow one to assess whether the capacity for palmitate oxidation and glycolysis is affected, irrespective of changes in substrate availability and transport, and alterations in the content of cofactors during hypertrophy of the in situ heart.

Palmitate oxidation rates were determined in 100 μl of the LV homogenerate in a total volume of 0.5 ml, essentially as described previously (17). The final composition of the incubation medium was (in mM) 22.6 KCl, 114.5 Tris, 15 KHP04, 7.5 MgCl2, 1.9 EDTA, 87.5 sucrose, 5 ATP, 1 NaDP+, 0.1 CoA, 0.5 l-malate, 0.5-l-carnitine, and 0.025 cytochrome c. After preincubation (5 min), the reaction was started by the addition of unlabeled and [1-14C]palmitate bound to albumin in a 5:1 molar ratio (specific activity 40 Bq/nmol; final concentration 0.12 mM). Incubations were carried out in an airtight vial at 37°C. Reactions were stopped by addition of 200 μl 3 M perchloric acid, either just before (t = 0) or 10 or 20 min after addition of palmitate. The CO2 produced was trapped in 400 μl ethanolamine/ethylene glycol (1:2 vol/vol). To trap all CO2 produced, the vials were left overnight at 4°C. Acid-soluble products (citric acid cycle intermediates) were separated from palmitate by centrifugation. The 14CO2 trapped and 14C-labeled acid-soluble products were determined by liquid scintillation counting. Palmitate oxidation rate was calculated as the sum of the 14CO2 trapped and 14C-labeled acid soluble products and expressed as nanomoles per minute per grams wet weight of tissue.

Glycolytic rates were determined using 100 μl of the LV homogenerate in a total volume of 0.5 ml. By following the method of Beatty et al. (6), the final composition of the incubation medium was (in mM) 70 KCl, 100 Tris, 8 KHP04, 5 MgSO4, 0.5 EDTA, 60 sucrose, 1 ATP, 1 ADP, 0.5 NaDP+, 0.2 NADP, 1 dichloroacetate, and 0.04 coenzyme A. Reactions were started by addition of unlabeled and [5-3H]glucose (3 Bq/nmol; final concentration 11 mM) and stopped by addition of 200 μl 3 M perchloric acid immediately before the addition of glucose (t = 0 min) or after 20 or 40 min. The medium was subsequently neutralized with 70 μl 10 M KOH, and 75 μl of the medium was filtered through an anion exchange resin (200–400 Mesh Dowex 1-X4; Sigma) pretreated with 0.4 M
potassium borate (7). The amount of $^3$H$_2$O, reflecting the glycolytic rate, was determined by liquid scintillation counting for $^3$H. [14C(U)]glucose was added to the incubation medium to correct for leakage of glucose through the column.

**Plasma glucose and fatty acid levels.** Plasma glucose and fatty acid levels were determined in ad libitum fed rats. Plasma (unesterified) fatty acids were determined by means of the NEFA C kit, following the instructions of the manufacturer (Wako; Neuss, Germany). Plasma glucose was determined with the use of a spectrophotometric analyzer (8) (Cobas Bio, Roche Analytical; Nutley, NJ).

**Histological analysis.** Cross sections (8 μm) of the LV of the heart were cut on a cryostat at −80°C and stored at −80°C until use. Sections were stained with Sirius red, which has a high affinity for collagen. Collagen-positive areas were quantified in 12 randomly selected fields by densitometrical analysis (Quantimet 570 Image Analyser; Leica, Cambridge, UK). Data were expressed as percentage cross-sectional area of the LV occupied by collagen (12).

**MHC composition.** Frozen LV tissue was pulverized with a mortar and pestle precooled in liquid nitrogen. Total RNA and protein were extracted from the pulverized left ventricles with the use of TRIZol reagent (GIBCO-BRL Life Technologies; Gaithersburg, MD). The protein precipitate was dissolved in 1% SDS. The protein content was determined by using a Silverstain plus kit (Bio-Rad; Hercules, CA) and scanned on a Fluor-S Imager (Bio-Rad), and the relative proportions of α- and β-MHC were determined using Quantity One (Bio-Rad).

**Microarray analysis.** Total RNA was isolated from the LV of sham and of 3-, 7-, and 28-day TH-treated rats. The microarray was performed with Incyte rat gene expression microarrays (GEM 2.20 and 3.17; containing 8,485 and 8,958 sequences, respectively; Genome Systems; St. Louis, MO), as described previously (3). The raw data set was filtered using various algorithms and Incyte software to eliminate spots with poor signal-to-noise ratio, leaving >15,000 sequences in total (some of which are present on both GEMs or represented more than once on the same GEM). Sequences of which the expression changed by at least 1.7-fold in duplicate assays were considered to be up- or downregulated.

**Northern blots.** Northern blots were performed as described previously (41–43). In short, 10 or 20 μg total RNA was size fractionated on a denaturing gel (1% agarose, 1× MOPS, and 2% formaldehyde) and blotted by capillary transfer to a nylon membrane (Hybond-NX, Amersham; Slough, UK). The blots were hybridized with cDNA probes labeled with [α-32P]dCTP (3,000 Ci/mmol; Amersham) by random priming (Radprime, Life Technologies). The blots were then exposed to an imaging screen and scanned with a Personal FX Phosphor-Imager (Bio-Rad). Signals were quantitated with the use of Quantity One Software (Bio-Rad). In a blot, the signals were normalized to the 18S ribosomal RNA signal to correct for possible loading and transfer differences. Possible interblot differences were accounted for by subsequent normalization to corresponding sham samples that were present on each blot. The probes not described elsewhere (41–43) are presented in more detail below.

For markers for cardiac hypertrophy, we used a 0.7-kb HindIII/BamH I fragment of ANF, a 0.32 kb EcoRI/NsiI fragment of cardiac SERCA2a and a 1.3-kb BamH I/PstI fragment of collagen Iα (kindly provided by J. Cleutjens, Maastricht University, Maastricht, The Netherlands). We used glucose transporter 4 (GLUT4) hexokinase II (HKII), GAPDH, and a 3-kb EcoRI-PvuI fragment of mouse pyruvate dehydrogenase (PDH) E-1α-subunit (generous gift from H. H. Dahl, Royal Children’s Hospital; Melbourne, Australia) as markers for glucose metabolism. Fatty acid translocase (FAT/CD36), heart-type fatty acid-binding protein, acyl-CoA synthetase (ACS), muscle-type carnitine palmitoyl transferase I (mCPT-I), and long-chain acyl-CoA dehydrogenase (LCAD) were studied as markers of fatty acid metabolism. In addition, a 0.7-kb HindIII/Xba I fragment of glycogen phosphorylase and a probe for citrate synthase were used as markers of glycogen metabolism and the citric acid cycle, respectively. In addition, a probe for PPAR-α was used. Furthermore, blots were probed for rat UCP2 and UCP3.

**Statistics.** Data are presented as means ± SD. Statistical analysis was performed using INSTAT version 2.00 (GraphPad Software; San Diego, CA). For multiple comparisons, ANOVA followed by Tukey’s post hoc test was applied to locate the differences. Differences were considered significant at $P < 0.05$.

**RESULTS**

**Animal characteristics.** All rats were 11 wk old at the time of death. TLs were similar for all groups (Table 1), indicating that TH treatment for up to 28 days did not affect growth rate. BWs, however, were significantly lower in rats treated with TH for 28 days. The reduced BW/TL ratio in these rats most likely reflects the loss of adipose tissue due to enhanced lipolysis. Indeed, plasma fatty acid levels were more than doubled already after 3 days of TH treatment and remained elevated thereafter (Table 1). Plasma glucose levels were not significantly affected by TH treatment (Table 1).

**Cardiac dimensions and function.** TH induces a marked hypertrophy, both when expressed as absolute HW and when normalized to TL or BW (Table 1). After 3 days of TH treatment, the HW/TL ratio had increased by 10.22±0.33 on August 15, 2017 http://ajpheart.physiology.org/ Downloaded from

<table>
<thead>
<tr>
<th>Table 1. Animal characteristics</th>
</tr>
</thead>
<tbody>
<tr>
<td>BW, g</td>
</tr>
<tr>
<td>-------</td>
</tr>
<tr>
<td>Sham</td>
</tr>
<tr>
<td>TH, day 3</td>
</tr>
<tr>
<td>TH, day 7</td>
</tr>
<tr>
<td>TH, day 28</td>
</tr>
</tbody>
</table>

Values are means ± SD; n of experiments are shown in parentheses. TH, thyroid hormone; BW, body weight; TL, tibia length; HW, heart weight. TH-treated animals were given 200 μg/KG TH per day. Fatty acids and glucose were analyzed in plasma (see MATERIALS AND METHODS). *$P < 0.05$, significantly different from sham; †$P < 0.05$, significantly different from day 3; ‡$P < 0.05$, significantly different from day 7.

AJP-Heart Circ Physiol • VOL 284 • JANUARY 2003  •  www.ajpheart.org
increased by 21%. After 28 days, this ratio was increased by 47%. The echocardiographic measurements corroborated these observations and showed that LV free wall thickness markedly increased over time in TH-treated rats, reaching statistical significance after 7 days of TH treatment (Fig. 1A). The ventricular hypertrophy was concentric in nature as reflected by the increased ratio of end-diastolic LV wall thickness over LV inner diameter (data not shown).

Echocardiographic assessment of cardiac function did not reveal any acute effects of TH. Within hours after the first bolus of TH or vehicle, no significant differences in HR and LV EF were observed. However, at day 7, HR increased from 447 ± 9 to 536 ± 31 beats/min (P < 0.01) and EF was significantly elevated (Fig. 1B), suggesting enhanced cardiac function following TH treatment. Heart rate remained consistently increased during the remainder of the protocol. EF was highest between day 7 and 14 and gradually declined thereafter.

Phenotypic markers of TH-induced cardiac hypertrophy. The TH-induced increase in cardiac mass was not accompanied by changes in LV volume fraction of collagen (Fig. 2A), indicating that the hypertrophic response was not associated with fibrosis. In sham rats β-MHC protein could be detected in the majority of hearts analyzed (5.5 ± 6.8% of MHC isoforms; n = 16). β-MHC expression was undetectable after 28 days of TH treatment (Fig. 2B). mRNA levels of SERCA2a were not altered in hearts of TH-treated animals at any time point (Fig. 2C), which was consistent with the microarray analysis (data not shown). With the exception of M-band protein, the expression of none of the sarcomeric proteins represented on the gene chips changed significantly (data not shown). ANF mRNA levels of the LV of hyperthyroid animals were identical to those of the sham group before and during the first 7 days of TH treatment (Fig. 2D). After 28 days of treatment, however, ANF expression was found to be substantially elevated.

TH and metabolic remodeling of the heart. Fatty acid oxidation capacity, measured in LV homogenates under optimal assay conditions and normalized to wet weight, was not significantly affected by TH treatment at any time point (Fig. 3A). Likewise, the glycolytic capacity did not differ between the TH and sham groups (Fig. 3B). Consistent with these biochemical observations, cardiac mRNA levels of proteins involved in fatty acid uptake and metabolism (FAT/CD36, fatty acid binding protein, ACS, mCPT-I, and LCAD) (Fig. 4A) and glucose uptake and metabolism (GLUT4, HKII, GAPDH, and PDH) (Fig. 4B) were not specifically affected by TH. Similarly, the mRNA level of citrate synthase, a marker of mitochondrial citric acid cycle activity, did not change (Fig. 4C). Correspondingly, microarray analysis also did not reveal specific alterations in the expression of genes involved in FA metabolism (e.g., ACS, very long chain acyl-CoA synthetase (VLACS), mCPT-I, short chain acyl-CoA dehydrogenase (SCAD), LCAD, and very long chain acyl-CoA dehydrogenase (VLCAD)) or glucose handling (e.g., GLUT4, GAPDH, and LDH) (data not shown). Collectively, the biochemical and molecular data suggest that despite the marked degree of ventricular hypertrophy in the TH-treated animals and contrary to what has been reported for other forms of cardiac hypertrophy (1, 33), fatty acid and glucose metabolism was not specifically affected in this type of hypertrophy, but both followed the increase in muscular mass.

The mRNA level of UCP2 was more than doubled already after 3 days of TH treatment and remained elevated thereafter (Fig. 4C). Furthermore, after 28 days of TH treatment, the LV mRNA content of UCP3 was also increased (Fig. 4C). Finally, the mRNA level of PPAR-α was reduced to 60% after 3 days of TH treatment (P < 0.05) but was normal again after 7 and 28 days of treatment (Fig. 4C).

Because hypertrophy and failure were found to be associated with changes in the expression of TH receptor isoforms (21, 22), attention was paid to nuclear receptors and signaling pathways that may be activated secondary to TH treatment. Interestingly, in the microarray analysis, the expression of deiodinase, an enzyme involved in the inactivation of TH, was the only gene in this category that was transiently (1.7-fold
DISCUSSION

Clinical (18, 29) and experimental (11, 31) investigations have lent support to the contention that TH supplementation improves performance of the failing heart. Recent studies (21, 22) indicated that TH supplementation helps to overcome the otherwise impaired TH signaling that is associated with cardiac failure. At the same time, it is commonly acknowledged that hyperthyroidism ultimately may be detrimental to the heart. Indeed, in the present time-course study in rats, we show that TH supplementation leads to a massive hypertrophy, which is associated with an initial improvement in cardiac function. Prolonged TH treatment, however, may cause a pathological form of hypertrophy, as evidenced by enhanced expression of ANF and UCP2 and UCP3 in the LV tissue, in association with a decline in LV EF.

TH-induced hypertrophy: physiological or pathological? Supplementation of TH is associated with a rapid and profound increase in cardiac mass. Notwithstanding the overt hypertrophy (almost 50% increase after 28 days), the absence of changes in (pro-)collagen mRNA levels and collagen surface area indicates that fibrosis did not occur. The cardiac α-MHC protein content progressively increases at the expense of β-MHC during TH treatment. Along with the marked rise in EF (seen within 7 days after the onset of treatment), these findings all favor the idea that initially the hypertrophic response is physiological rather than pathological in nature.

The shift in myosin isoform content in favor of α-MHC is consistent with transcriptional regulation of the gene via its TH-response element (10, 27). In view of the presence of a TH-response element in the SERCA2a promoter, a rise in SERCA2a mRNA levels in hearts of TH-treated rats was anticipated (10). The present findings, however, indicate that the myocardial SERCA2a mRNA content did not change irrespective of the duration of hormone supplementation. This is in striking contrast with the marked rise in SERCA2a mRNA observed in earlier studies of intact hearts (2, 30) and on neonatal cardiomyocytes (20; unpublished observations from our laboratory). At the present state of the art, one can only speculate regarding the discrepancy between the lack of effect of TH on SERCA2a expression in intact hearts and the upregulation in isolated cardiac myocytes (20). We propose that in vivo the stimulatory effect of TH on SERCA2a expression is counteracted by its extracardiac effects, i.e., the hyperdynamic circulatory state (19, 45). Indeed, this suggestion is supported by experiments with heterotopically transplanted hearts, showing that SERCA2a expression in unloaded hearts is substantially more elevated than in the hemodynamically loaded host hearts in response to TH treatment (30). Moreover, Ojamaa and colleagues (31) observed that after myocardial infarction, TH administration failed.
to restore the expression of SERCA2a in the viable region of the heart, stressing the notion that the relationship between TH and cardiac SERCA2a expression is highly complex.

In the clinical setting, the increased hemodynamic load may ultimately lead to high output failure (45). Because cardiac hypertrophy is a result of chronic hemodynamic overload, it is comprehensible that during prolonged TH treatment a pathological form of hypertrophy evolves. Consistent with this notion is the observation that TH supplementation is associated with activation of the intracardiac renin-angiotensin system (25, 26), which is commonly regarded as being involved in the development of pathological hypertrophy. Along with the present observations that TH supplementation only leads to a transient increase in cardiac performance and that ANF is expressed in ventricular tissue after prolonged TH treatment, the combined data suggest that TH-induced hypertrophy resembles physiological hypertrophy initially and gradually changes toward a more pathological form of hypertrophy.

Cardiac metabolism. On the basis of both extensive microarray analysis and on Northern blot data of a large panel of candidate genes, we found no evidence for specific changes in the expression of genes involved in the uptake and metabolism of either glucose or fatty acids in TH-induced hypertrophied cardiac tissue. Consistent with this, the flux of both fatty acids and glucose through their corresponding metabolic pathways as measured under optimal conditions in cardiac homogenates was not affected by TH at any time point. It should be emphasized, however, that the rate of glycolysis and fatty acid oxidation was normalized to the wet weight of LV tissue. This finding implies that the...
metabolic capacity as measured in homogenates follows the increase in tissue mass. The corollary of this notion is that in TH-induced hypertrophy the intracellular capacity of substrate utilization most likely remains in balance with the increase in cardiac energy demand. Earlier findings of Seymour and colleagues (34), indicating no change in the maximum activity of glycolytic enzymes in TH-treated hearts when normalized to wet weight of tissue are in line with the present observation.

The absence of changes in mCPT-I expression is consistent with previous findings indicating that, unlike liver, heart mCPT-I mRNA levels do not change in response to TH (13). However, in intact isolated cardiac myocytes from hyperthyroid rats treated with a fivefold higher dose of TH compared with the present study, CPT-I enzyme activity was enhanced in association with an increased rate of fatty acid oxidation. The discrepancy between these and our findings may be related to differences in the experimental models used (cardiac homogenates and isolated myocytes) and doses of TH applied. The use of cardiac homogenates in the present study precludes any confounding secondary effects due to differences in cardiac work between euthyroid and hyperthyroid animals, and changes in substrate supply and concentrations of intracellular cofactors. However, possible differences in transsarcolemmal transport rate of substrates cannot be appreciated in this preparation. In addition, it should be stressed that protein (enzyme) levels may be changed even in the absence of changes in tissue content of their corresponding mRNA.

Previous studies have shown that in pressure- and volume-overload cardiac hypertrophy, substrate preference is shifted from fatty acids utilization to glucose (1, 16, 46), which is generally considered a hallmark of the return to the fetal gene program of the hypertrophied heart (4, 15). This shift was reported to go along with diminished expression of a set of β-oxidation genes (32, 33, 46). Prolonged TH treatment, resulting in a substantial increase in cardiac mass, obviously did not evoke the shift in fuel selection from fatty acids to glucose. In contrast, the chronically elevated plasma fatty acid levels may even favor the utilization of fatty acids in the intact heart due to increased availability of these substrates.

It is of note that similar to what has been observed by others (9), TH was found to increase cardiac mRNA levels of UCP2 and UCP3, which are believed to reduce the mitochondrial efficiency by dissipating the electrochemical proton gradient. In this way, the elevated expression of UCP might hamper energy conversion in the hypertrophied heart, and consequently cardiac function. However, the significance of this TH-mediated effect in muscle has been questioned because ATP production by skeletal muscle mitochondria of hypertrophied rats was found to increase, rather than decrease, under these conditions (36). Van der Lee et al. (42) and Young et al. (47) showed that the expression of UCPs is stimulated by fatty acids, most likely in a PPAR-dependent manner. Similarly, the rise in plasma fatty acid concentration induced by fasting was accompanied by an enhanced expression of other PPAR-α-responsive genes in the heart (43). At first sight, the enhanced expression of UCP2 and UCP3 in the hearts of TH-treated animals seems consistent with the marked rise in circulating fatty acid levels. However, the time course of changes in UCP expression, in particular UCP3, does not parallel the rapid rise in plasma fatty acid levels. Moreover, because we observed a transient decrease in PPARα mRNA, a direct relationship between the expression of this transcription factor and that of the UCPs is neither apparent. Finally, the expression of other PPAR-regulated genes, among which FAT, ACS, CPT-I, and LCAD is not changed at all. This seems to exclude a specific role of PPARα in the regulation of the expression of cardiac genes by elevated TH.

In summary, the present biochemical and molecular data indicate that irrespective of the duration of TH supplementation and the severity of the ensuing hypertrophy, the increase in cardiac mass is met by a parallel increase in glycolytic and fatty acid oxidative capacity. In this respect, the TH-induced hypertrophy compares favorably to hypertension-induced cardiac hypertrophy. However, several typical markers of physiological hypertrophy are not present, and cardiac performance is only transiently enhanced. Furthermore, with prolonged TH supplementation, ANF and UCP2/3 expression becomes elevated. This suggests that with time the beneficial effects of TH are overruled by detrimental (most likely extracardiac) effects of the hormone.

The authors thank N. Herben for the collagen analysis, P. Leenders for instruction on echocardiography on rats, Drs. G. Porter (Incyte) and C. Evelo for microarray analysis, and Claire Bollen for help in preparing the manuscript. We highly appreciate the critical reading of the manuscript by Dr. Robert S. Reneman and the helpful discussions.

This study was supported by Netherlands Heart Foundation Grant 97.092. M. van Bilsen is an established investigator of the Netherlands Heart Foundation (D98.015).

REFERENCES

7. Belke DD, Larsen TS, Lopaschuk GD, and Severson DL. Glucose and fatty acid metabolism in the isolated working mouse
THYROID HORMONE AND CARDIAC METABOLISM