miR-22-3p is involved in gluconeogenic pathway modulated by 3,5-diiodo-L-thyronine (T2)

Rosalba Senese1,3, Federica Cioffi2,3, Giuseppe Petito1, Pieter de Lange1, Aniello Russo1,3, Fernando Goglia2, Antonia Lanni1* & Nicoletta Potenza1

The 3,5-diiodo-L-thyronine (T2) has emerged as an active iodothyronine and its beneficial effects on glucose metabolism including glucose tolerance and insulin resistance is well established. However, little is known about its molecular mechanisms. Given the emerging importance of microRNAs in various metabolic diseases, in this study a possible link between the effects of T2 on glucose metabolism and miRNA expression was investigated by using an in vivo model in which T2 was administered in rats receiving a high fat diet, a condition known to impair glucose homeostasis. The results showed that T2-treated rats had a better tolerance to glucose load and a better performance at the insulin tolerance test in comparison to high fat diet animals. Interestingly, in the serum of the animals treated with T2 there was a general decrease of miRNAs with miR-22a-3p, miR-34c-5p and miR-33a-3p significantly downregulated. Furthermore, miR-22a-3p had the largest variation pointing toward its preeminent role in T2 metabolic effect. In fact, in liver there was an up-regulation of its target (Transcription Factor 7) Tcf7, which had an important impact on gluconeogenesis. This study provide, for the first time, evidences that miRNAs are involved in the effects exerted by T2 on glucose homeostasis.

Thyroid hormones (THs; 3,5,3',5'-tetraiodo-L-thyronine,T4 and 3,5,3'-triiodothyronine, T3) are essential for many physiological processes such as growth, differentiation and metabolism1,2. In the last years, several data have shown that some THs derivatives (natural metabolites and analogues) play a relevant biological role on the metabolism (for reviews see3–8). Among these, many studies are focused on 3,5-diiodo-L-thyronine (T2), a biologically active iodothyronine9-11 (for review see7,12–15). In rats fed on a high-fat diet (HFD), the simultaneous administration of T2 (25 μg/100 g body weight for 4 weeks) prevented hyperlipidemia, liver adiposity, body weight gain and insulin resistance16–18 and, at that dose, no signs of thyrotoxicosis were detected16,17. Interestingly, it has been also shown that T2 administration to HFD-fed rats improved glucose tolerance, insulin sensitivity18 and gluconeogenesis in hepatocytes17. Furthermore, a possible physiological link between glucose metabolism16,20 and circulating T2 has been highlighted by findings from a healthy euthyroid population. To date, however, the mechanism by which T2 affects glucose homeostasis remains to be clarified.

Many reports have demonstrated the involvement of miRNAs in various diseases21–23, mainly in the etiology of cancer but also in metabolic disorders such as obesity and type 2 diabetes (T2D)24,25 which, in turn, are characterized by an impaired glucose homeostasis26,27. Indeed, these small non-coding RNA molecules modulate the levels of kinases and enzymes in the glucose metabolism28 by affecting posttranscriptional regulation of gene expression. Particularly, an altered hepatic miRNA signature for obese diabetic db/db mice was identified by comprehensive miRNome and in silico analyses29. This report showed an altered Wnt signaling pathways in diabetic liver29, and various lines of evidence suggest that Wnt/β-catenin signaling regulates hepatic glucose production30–32.

Based on those considerations, the aim of this study was to examine the potential involvement of miRNAs in the mechanisms of action by which T2 affects glucose homeostasis in rats fed on a high-fat-diet. This animal model was chosen since HFD treated rats showed an altered glucose homeostasis and some metabolic disorders, such as altered glucose tolerance and insulin-resistance, thus revealing the beneficial effect of a T2 treatment. In particular: a) a screening of circulating miRNA changes in the serum from HFD-T2 treated vs HFD rats was performed to identify miRNAs potentially involved in the effects of T2; b) the top-ranked microRNA from the

1Department of Environmental, Biological and Pharmaceutical Sciences and Technologies, University of Campania “L. Vanvitelli”, Caserta, Italy. 2Department of Sciences and Technologies, University of Sannio, Benevento, Italy. 3These authors contributed equally: Rosalba Senese and Federica Cioffi. *email: antonia.lanni@unicampania.it
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Table 1. T2 administration prevents HFD-induced changes in systemic metabolic parameters and fat accumulation. Body Weight (BW), percentage ratio liver weight/body weight (LW/BW); percentage ratio heart weight/body weight (HW/BW); percentage ratio gastrocnemius weight/body weight (GW/BW); percentage ratio white adipose tissue weight/body weight (WW/BW); and serum cholesterol levels in rats fed on to standard diet (N), to high-fat diet (HFD) and to HFD with contemporary administration of T2 (HFD-T2). *p < 0.05 vs HFD; #p < 0.05 vs HFD. Values are mean ± SEM of 5 independent treatments.

| Parameter          | N         | HFD       | HFD-T2     |
|--------------------|-----------|-----------|------------|
| BW (g)             | 405 ± 10  | 470 ± 12.6* | 410 ± 13.2* |
| LW/BW (%)          | 2.7 ± 0.23 | 3.18 ± 0.18  | 2.71 ± 0.22  |
| HW/BW (%)          | 0.3 ± 0.01 | 0.25 ± 0.01  | 0.309 ± 0.02 |
| GW/BW (%)          | 0.6 ± 0.04 | 0.5 ± 0.02   | 0.55 ± 0.03  |
| WW/BW (%)          | 2.8 ± 0.2  | 3.2 ± 0.32*  | 2.75 ± 0.15* |
| Cholesterol mg/dl  | 55 ± 2.5  | 80 ± 3.7*    | 60 ± 2.36*   |
| Triglycerides      | 125 ± 8.12| 250 ± 10.2*  | 124 ± 10.5*  |

In vivo T2-induced down-regulation of miR-22a-3p results in an up-regulation of Wnt-Responsive Transcription Factor Tcf7 with an impact on gluconeogenesis. miR-22 is abundantly expressed in the liver and its expression was increased in animals models of insulin resistance and type 2 diabetes. Moreover, miR-22 has been reported to have as direct target the Wnt-responsive transcription Factor Tcf7 in mouse and human cells, thus impairing gluconeogenesis and, as consequence, an increase in circulating glucose level. These data prompted us to investigate this pathway in our experimental rat model. First, miR-22 expression was measured in the liver, revealing that T2 administration to HFD-fed rats significantly reduced miR-22 expression when compared to HFD-fed rats. In HFD-fed rats, the miR-22a expression was increased, although not significantly when compared to N animals. Then, based on the prediction of a conserved miR-22 binding site on rat Tcf7 3’UTR, the Tcf7 hepatic expression was measured, showing a reversed pattern with respect to that of miR-22, as expected for a target of the miRNA.

Finally, in the same samples, we also analyzed the expression levels of genes of gluconeogenesis, i.e., glucose-6-phosphatase (G6Pase), fructose 1-6 bisphosphate (FBPase1) and phosphoenol pyruvate carboxykinase (PEPCK), whose expression increased after Tcf7 targeting siRNA transfection of cell culture. In our in vivo model a strong decrease in the expression of G6Pase and FBPase was observed in response to T2 administration, wherein Tcf7 resulted increased (Fig. 5). Furthermore, T2 treatment prevented the increase in glycolytic enzyme (liver pyruvate kinase (LPK)) induced by hyperlipidic diet (Fig. 5).
Discussion

It is well established that T2 exerts considerable effects on the metabolic activities. Indeed, several laboratories have demonstrated that it is able to influence both the energy metabolism and some biochemical pathways involved in lipid and glucose metabolism. However, the mechanisms by which it carries out these actions...
must still be clarified. About glucose metabolism it is known that T2 showed beneficial effect on glucose tolerance and insulin-resistance, as well as overweight/obesity and diabetes. Recently, many reports have shown the involvement of aberrant miRNA expression in metabolic disorders. Because of this, in an attempt to reveal a

Figure 3. Circulating miR-22-3p. Serum samples were collected at 1 months from normally-fed rats (N), HFD-fed rats (HFD) and HFD-fed T2-treated rats. Relative miR-22 level was determined by RT-QPCR as described and reported as mean ± SEM of five independent experiments (N = 5). **p < 0.01.

Figure 4. Hepatic expression of miR-22 and of its Tcf7 target. (A) RT-qPCR analysis of miR-22a in liver of N, HFD, HFD-T2 rats. (B) Predicted base-pairing between miR-22 and 3’UTR of Tcf7 mRNA. (C) Hepatic Tcf7 expression measured by RT-qPCR. Values are means ± SEM of five independent experiments (N = 5), *p < 0.05 vs N; ***p < 0.05 vs HFD; **p < 0.05 both vs N and HFD.

Figure 5. Hepatic expression of rate-limiting genes of gluconeogenesis. RT-qPCR analyses of glucose-6-phosphatase (G6Pase), fructose 1-6 bisphosphate (FBPase1) and phosphoenol pyruvate carboxykinase (PEPCK) were performed on RNA purified from the liver of N, HFD, HFD-T2 rats. Values are means ± SEM of five independent experiments (N = 5). **p < 0.05 vs N and HFD, *p < 0.05 vs N, ***p < 0.05 vs HFD.
Figure 6. A model depicting the miR-22-mediated beneficial effect of T2 in HFD rats. The effect of T2 is represented by bold lines: the down-regulation of miR-22 in response to T2 treatment results in a reduced inhibitory effect on the expression of Tcf7 whose hepatic level increases, with the consequence of a reduced expression of gluconeogenesis enzymes and gluconeogenesis impairing.

possible link between miRNA regulation and the beneficial effects of T2 in an HFD-context, a screening of circulating miRNAs was performed on serum samples from HFD and HFD-T2 animals. In the present study for the first time, we provide evidences that miRNAs are involved in the actions exerted by T2 on glucose homeostasis.

The results revealed that the T2-treated animals are protected against fat accumulation and the resulting insulin resistance. In fact, the diiodothyronine prevented body weight gain and the increase in white adipose tissue mass induced by high fat diet; in addition, the diiodothyronine was able to avoid a serum lipid increase (see Table 1). By this way, T2 contributed to improve glucose tolerance and insulin sensitivity (see Fig. 1). As already shown by Lanni et al. using the same experimental design, these effects were not ascribed to central actions of T2 and its actions were not associated with thyrotoxicosis16. Regarding miRNAs, our data revealed a general decrease of miRNAs in the serum of HFD-T2 versus HFD animals, with miR-22-3p, miR-34c-5p, and miR-133a-3p significantly downregulated, suggesting, for the first time, that some biological actions of T2 could be at least in part due to miRNA-based mechanisms. Consistent with our data obtained with an in vivo model, some papers already linked the three miRNAs with glucose metabolism. In particular, miR-34c-5p was found abundantly expressed in the circulating monocytes of type 2 diabetes patients16. From the same miRNA family, miR-34a was down-regulated in type 2 diabetes patients after therapeutic resveratrol treatment43; again its expression was altered in obese db/db mice and its overexpression was associated with increased lipid accumulation in HepG2 cells35. With regard to miR-133a-3p, it was shown that diabetes leads to its downregulation, with an impact on the regulation of target genes involved in cardiomyocyte hypertrophy, heart failure, cardiac arrhythmia and muscular dystrophy19,40,41. Finally, regarding to the most reduced miRNA in response to T2 treatment, miR-22 (~3.1-fold, HFD-T2 vs HFD), its increased hepatic expression has been reported in mouse models of insulin resistance and type 2 diabetes42,43. It is important to note that T2, a key transcription factor involved in the regulation of gluconeogenesis, being demonstrated as its target in mouse and human cells31,43-45. This pathway was further investigated in liver of our rat model, in the light of the effects of T2 in alleviating hyperglycemia, improving glucose tolerance and insulin sensitivity during HFD diet, a condition that impairs glucose homeostasis. We focalized our study principally on the liver because it is known that gluconeogenesis takes place mainly in the liver46,47, even if it cannot be ruled out that other insulin-resistant tissues could contribute to the level of circulating miR-22. The results showed that T2 administration to HFD-fed rats significantly reduced hepatic miR-22 expression when compared to HFD-fed rats alone. Then, based on the prediction of a conserved miR-22 binding site on rat Tcf7 3′UTR, the Tcf7 hepatic expression was measured, and accordingly with that reported for mouse and human cells37, revealed a reversed pattern with respect to that of miR-22, with a consequent reduced expression of Tcf7 targets (G6Pase and FBPase) that would result in a decrease of glucose release and improvement of glucose tolerance observed after T2 administration (Fig. 1). The reduction in miR-22 expression well correlates with the beneficial effects of T2 administration, since it has been found modulated in different metabolic disorders. Diniz GP et al. in 2017 showed that miR-22 is an important regulator of dyslipidemia as the loss of this miRNA attenuated the gain of fat mass and prevented dyslipidemia induced by hyperlipidic diet48. In addition, it has been shown the miR-22 involvement in cardiomyocyte hypertrophy and cardiac remodeling, probably by targeting SIRT1, a Nicotinamide Adenine Dinucleotide +-dependent histone deacetylase involved in a variety of biological processes, including energy metabolism, cell proliferation and chromatin dynamics49,50. Furthermore, we have already demonstrated that T2 prevented the increase in protein levels of LPK HFD-induced contributing to improve glucose tolerance37. Here we showed that T2 prevents that increase downregulating LPK mRNA levels suggesting that, in this way, the diiodothyronine could maintain unaltered protein levels of LPK. Again, the effect on improvement of glucose metabolism could be, at least in part, due to an increase in resting metabolic rate as already reported by Lanni et al.16.

In conclusion, our results suggest a miRNA-based action of T2, since different miRNAs changed their level in response to T2 administration; furthermore, our data indicate that one way through which T2 regulates glucose homeostasis could be the down-regulation of miR-22, resulting in an up-regulation of its Tcf7 target which, in turn, impairs gluconeogenesis by inhibiting the expression of gluconeogenesis enzymes (Fig. 6).

Materials and Methods

Animals and animal care. All animals received humane care according to the criteria outlined in the Guide for the Care and Use of Laboratory Animals prepared by the National Academy of Sciences and published by the National Institutes of Health. All animal protocols were approved by the Organism in charge of animal care.
welfare (OBPA) of the University of Campania “Luigi Vanvitelli” (Caserta-Italy) and the Italian Minister of Health (Permit Number:704/2016-FR). Every effort was made to minimize animal pain and suffering. Male Wistar Rats (250–300 g, aged 8 weeks) were individually caged in a temperature-controlled room at 28 °C (thermoreutral temperature for rats) under a 12-h light/12-h dark cycle. Before commencement of the study, a commercial mash (Charles River Laboratories, Calco, Italy) was available ad libitum and the animals had free access to water27,31. At the start of the study (day 0), after 7 days of acclimatization to thermoneutrality, the rats were divided into three groups of 5 and treated for four weeks as follows:

- The first group (N) received a standard diet (total metabolizable percentage of energy: 60.4 carbohydrates, 29 proteins, 10.6 fat J J⁻¹; 15.88 kJ gross energy g⁻¹; Muscedola, Milan, Italy) with a daily injection intraperitoneally of vehicle27,31.
- The second group (HFD) received a high-fat diet (280 g diet supplemented with 395 g of lyophilized lamb meat [Liomellin, Milan, Italy], 120 g cellulose [Sigma-Aldrich, St. Louis, MO], 20 g mineral mix [ICN Biomedical, Solon, OH], 7 g vitamin mix [ICN], and 200 g low-salt butter [Lurpak, Denmark]; total metabolizable percentage of energy: 21 carbohydrates, 29 proteins, 50 fat J J⁻¹; 19.85 kJ gross energy g⁻¹) with a daily injection intraperitoneally of vehicle27,31.
- The third group (HFD-T2) received the same HFD with a daily injection of T2 (25 μg 100 g⁻¹ body weight intraperitoneally) (Sigma-Aldrich)27,31.

All animals continued to have free access to water. At the end of the treatment, the rats were anesthetized by intraperitoneal injection of chloral hydrate (40 mg 100 g⁻¹ body weight) and then decapitated. Liver, heart, gastrocnemius muscle, and abdominal white adipose tissue were excised, weighed, and immediately frozen in liquid nitrogen and stored at −80 °C for later processing27,31.

**Measurements of metabolic parameters.** The serum levels of cholesterol and triglycerides were determined by standard procedures. For the oral glucose tolerance test (OGTT), rats were fasted overnight and then orally dosed with glucose (3 g/kg body wt) dissolved in water17. For the insulin tolerance test (ITT), rats were fasted for 5 h and then injected intraperitoneally with insulin (homolog rapid-acting, 10 units/kg body wt in sterile saline; Novartis, Basel, Switzerland)17. Samples of blood were collected before the oral glucose tolerance test and insulin tolerance test and at various times thereafter (as indicated in the Fig. 1), and glucose and insulin values were determined by means of a glucose monitor (BRIOS, Ascensia, NY), calibrated for use with rats.

**RNA purification.** 200 μl of serum were used to extract total RNA, including small RNAs, by using miRNeasy Mini kit (Qiagen) according to the supplementary protocol provided for miRNA isolation from serum. A 5-μl aliquot of 5 mmol/l Syn-cel-miRNA-39 miScript miRNA Mimic was spiked into each sample before nucleic acid preparation to monitor the efficiency of miRNA recovery and to normalize miRNA expression in the subsequent real-time PCR analyses52.

Total RNA, including small RNAs, was also extracted by miRNeasy Mini kit (Qiagen) from liver tissues homogenized by polytron. The RNA was spectrophotometrically quantified by NanoDrop 2000c, ThermoScientific.

**miRNA profiling.** Reverse- transcription of RNA was performed by miScript II RT kit (SABiosciences); then, aliquots of cDNA preparation were used for real-time PCR profiling of mature miRNAs using Rat Serum&Plasma miRNA PCR Arrays (SABiosciences) in combination with miScript SYBR Green PCR kit (SABiosciences) and the MyiQ2 (Bio-Rad, Hercules, CA) instrument. Data analysis was performed with the web-based software package for the miRNA PCR array system (http://pcrdataanalysis.sabiosciences.com/miRNA/arrayanalysis.php). In brief, ΔCt value for each miRNA profiled in a plate is calculated using the formula ΔCt = CtmiRNA – Ctcel-miR-39. ΔΔCt for each miRNA across the two groups of samples (HFD and HFD T2 treated animals) is calculated using the formula: ΔΔCt = ΔCt of HFD-T2 treated group − ΔCt of HFD group. Expression fold-change was then obtained as 2−ΔΔCt (the normalized gene expression (2−ΔCt) in the T2 treatment group divided the normalized gene expression (2−ΔCt) in the HFD group. Data are reported as fold-regulation, where fold-regulation is equal to the fold-change for fold-change values greater than one (up-regulation) or is the negative inverse of the fold-change for fold-change values lower than one (down-regulation)52. The replicate 2−ΔCt values for each miRNA in the HFD and HFD-T2 treatment groups were statistically analyzed by Student’s t-test and the criteria of differential expression were p < 0.05.

**Real-time PCR analyses.** miR-22 was detected and quantified by RT-QPCR with TaqMan® miRNA assays from Applied Biosystems according to the manufacturer’s protocol. Serum miR-22 level was normalized to that of spiked-in cel-miR-39 as described above; the expression level of hepatic miR-22 was normalized to that of RNU6B used as reference gene in the 2−ΔCt method53.

The hepatic expression levels of the other transcripts were determined by RT-QPCR. In particular, 1 μg of total RNA was used to generate cDNA strands in a 20-μl reaction volume using the Super Script First Strand Synthesis System for RT-PCR (Invitrogen)17. An equivalent of 25 ng total cDNA was subsequently used in the amplification. Real-Time quantitative RT-PCR (QRT-PCR) was carried out with 50 nM gene-specific primers and IQ SYBR Green supermix (Biorad) using standard cycle parameters on a MyiQ2 (BioRad). A melting curve analysis was performed with the web-based software package for the miRNA PCR array system. Data analysis was performed with the web-based software package for the miRNA PCR array system (http://pcrdataanalysis.sabiosciences.com/miRNA/arrayanalysis.php). In brief, ΔCt value for each miRNA profiled in a plate is calculated using the formula ΔCt = CtmiRNA – Ctcel-miR-39. ΔΔCt for each miRNA across the two groups of samples (HFD and HFD T2 treated animals) is calculated using the formula: ΔΔCt = ΔCt of HFD-T2 treated group − ΔCt of HFD group. Expression fold-change was then obtained as 2−ΔΔCt (the normalized gene expression (2−ΔCt) in the T2 treatment group divided the normalized gene expression (2−ΔCt) in the HFD group. Data are reported as fold-regulation, where fold-regulation is equal to the fold-change for fold-change values greater than one (up-regulation) or is the negative inverse of the fold-change for fold-change values lower than one (down-regulation)52. The replicate 2−ΔCt values for each miRNA in the HFD and HFD-T2 treatment groups were statistically analyzed by Student’s t-test and the criteria of differential expression were p < 0.05.
PCK primers were designed by using the Primer 3 program (Untergasser et al., 2012), and synthesized and verified by sequencing at Eurofins Genomics (Ebersberg, Germany). Primers used were the following:

- rβ-ACT sense 5′-GGG TCA TGG TGT GAT ACT GGT TGC-3′
- rβ-ACT Antisense 5′-GGA GAT TAC TGC CCT GGC TCC TA-3′
- rG6Pase sense 5′-CAT TGC TGG GAG ACT GGT TCT-3′
- rG6Pase antisense 5′-CTG GCC TTG AAT GCT TCT-3′
- rLPK sense 5′-GAA TGC TGG GAG ACT GGT TCT-3′
- rLPK antisense 5′-TGT CCA AAG ATA GCG AGA-3′
- rPEPCK sense 5′-CTG GCC TTG AAT GCT TCT-3′
- rPEPCK antisense 5′-CCG GCT AGA TGG CAG ATG TA-3′
- rTCF7 sense 5′-AGG TCA GAT GGG TTG GAC TG-3′
- rTCF7 antisense 5′-GCA TGG CAG ACT GGG TTT AG-3′
- rFBPase sense 5′-ATG TGG GAC CAT GCAC CTTC TG-3′
- rFBPase antisense 5′-TTT TCA ATTT TCCC GTG TGG GAA TCC-3′

Statistical analysis. Results are expressed as mean ± SEM. The statistical significance of differences between N, HFD and HFD-T2 groups (n = 5) was determined using one-way ANOVA followed by a Student-Newman–Keuls test. Differences were considered significant at p < 0.05. The statistical significance of differences between HFD and HFD-T2 groups (n = 5) was determined using Student’s t-test. Differences were considered significant at p < 0.05.

Statement of ethics. Animal experiments conform to internationally accepted standards. All animals received humane care according to the criteria outlined in the Guide for the Care and Use of Laboratory Animals prepared by the National Academy of Sciences and published by the National Institutes of Health. All animal protocols were approved by the Organism in Charge of Animal Welfare (OBPA) of the University of Campania “Luigi Vanvitelli” (Caserta–Italy) and the Italian Minister of Health (Permit Number: 704/2016-PR).

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Author contributions
A.L., A.R. and F.G. conceived and designed the experiments, P.d.L. supervised the animal studies; R.S., F.C. and N.P. carried out all the animal studies; R.S., F.C., N.P. and G.P. performed the experiments; R.S., F.C., N.P. and G.P. analyzed the data; R.S., F.C. and N.P. wrote the manuscript; all the authors revised the manuscript.

Competing interests
The authors declare no competing interests.

Additional information
Correspondence and requests for materials should be addressed to A.L.
