

# To investigate Thyroid function abnormalities in individuals with chronic renal disease

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## Abstract

During our research, 50 CKD patients on conservative treatment were examined for thyroid function issues at the Department of Medicine at the Medical College and Hospital in China. The participants' ages range from 20 to 68. There were 34 men and 16 women among the patients, a ratio of 68% to 32%. The average duration of CKD symptoms is 9.84 months, although they can last anywhere from 4 months to 30 months. Between 6 and 32 ml/min, creatinine clearance occurs. The range of urea levels in the blood was 45-184mg/dl, with a mean of 102.12mg/dl. The range of serum creatinine concentrations was 3-14mg/dl, with a mean of 7.34mg/dl. In 10 patients, the serum calcium level was low; in 28 patients, it was normal; and in 12 patients, it was high. Twelve individuals had low serum phosphorous levels, whereas the 38 others were within the usual range. For example, serum T3 was measured between 0.2 and 2.0ng/ml, the average was 0, serum T4 was measured between 0.09 and 8.4g/dl, the average was 5.65, and serum TSH was measured between 0.06 and 38IU/ml, the average was 6.49 (normal range is 0.04 to 7.49 IU/ml). There were 29 individuals with low T3, 12 with low T4, and 4 with hypothyroidism in our research. 34 out of the 68 individuals had hypothyroidism-related symptoms. In our research, as people get older, the likelihood of developing Low T3 syndrome increases as well. As the severity of chronic renal disease grows, so does the number of individuals who have low T3. Renal failure severity is not related to the number of people with low T4 syndrome. It has been discovered that the serum T3 level is lower in those with poor GFR. The mean TSH levels in all phases of renal illness in individuals with low T3 syndrome are all within the normal range, and there was no connection between TSH levels and glomerular filtration rate (GFR). CKD patients' low T3 levels might be considered protective since they promote protein conservation.

**Keywords:** Chronickidney, pathophysiological, hyporeflexia, TSH, ESRD, asthenia

## Introduction

There are several pathophysiological mechanisms connected with chronic kidney disease, all of which lead to aberrant kidney function and a decrease in the kidney's ability to filter waste products.

An irreversible loss of renal function causes CKD, which manifests as an array of clinical symptoms such as metabolic, endocrine, excretory, and synthetic dysfunction. This leads to a buildup of non-protein nitrogenous compounds, which in turn causes metabolic dysfunction.

Patients who reach end stage renal illness have no option but to succumb to death since they will not be able to survive without replacement treatment.

Regardless of the underlying "cause, chronic kidney disease (CKD) is the last common pathway leading to permanent nephron loss, change of the "milieu interior," and a host of other health problems.

It's unclear how severe renal failure correlates with thyroid dysfunction. End-stage renal illness patients are at an increased risk of hypothyroidism in the range of 0-9 percent. Thyroid swelling (goitre) is more common in those with ESRD.

Taiwan's government, like that of many other nations, sees the value of information and communications technology (ICT) in advancing and improving education. There is a network service for all schools and educational research institutes in Taiwan, established up in 1980 by the Ministry of Education (MOE) (MOE 2006d). It's now possible for schools of all levels to collaborate easily and digitally with educational research institutions throughout the country in order to share information and resources. When it comes to

boosting educational advances, the MOE was more ambitious in the 1990s than it had been before. As a result, policies relating to information and communications technology (ICT) were developed between the middle of the 1990s and the end of the decade (MOE 2005). There have been internet connections in all elementary and secondary schools since 1999 as a result of the government's investment in ICT infrastructure and at least one computer lab has been established for teachers and students to use.

With its Blueprint for ICT Education at Primary and Secondary Schools, Taiwan's government took it a step further in 2001, shifting the emphasis away from teaching ICT as a separate topic to using ICT across the curriculum (MOE 2005). In 2002, the MOE launched the ICT Seed School Project (ICT SSP) up elementary schools, which tied in with the Blueprint. It was the goal of the ICT SSP to increase school use of new technologies by incorporating ICT into the lessons (MOE 2005).

The federal government provided training and financing to 600 ICT-capable schools known as 'ICT Seed Schools' under the ICT SSP in order to increase their usage of ICT across the curriculum. Their next responsibility was helping other schools improve their information and communication technology (ICT).

Noteworthy is that the Ministry of Education's study (2006c) shows that although some ICT Seed Schools were highly effective in implementing and maintaining the ICT SSP, others were less so. The International Telecommunication Union (2003) found that Taiwan had the 9th-best overall ICT education and affordability rankings, with the 3rd best ranking in the developed Asia-Pacific region, but the results of a recent survey show that Taiwan has a significant digital divide between rural and urban education (MOE 2006a). Many industrialised nations have a similar but hard goal of narrowing the digital divide between rural and urban schools, as shown by the MOE's recent international ministerial conference on 'ICT for Better Education' in Taiwan (MOE 2007). There has been little study done on how rural schools, particularly those that are unable to improve further, handle pedagogical reform and innovations in ICT deployment despite this alarming indicator. ICT software and hardware are clearly not enough to support innovative pedagogical practises in educational settings if schools, particularly those in rural regions are to be successful in adopting and maintaining ICT integration.

## **Literature Review**

An isthmus connects the two lobes of the thyroid gland. The suprasternal notch and cricoid cartilage are located in front of the trachea, where it is located. The thyroid gland is larger in females than in boys, weighing between 12 and 20 grammes. The parathyroid glands, which release parathormone, are located near the rear of each thyroid lobe.

T3 (Triiodothyronine) and T4 are the two most important hormones released by the thyroid gland (Thyroxine). To keep metabolic balance in adults, thyroid hormones are required. "Thyroid hormones play an important role in cell differentiation throughout development. Thyroid Stimulating Hormone mostly regulates T3" and T4 production. Trenbolone and triiodothyronine (T3 and T4) are secreted in response to TRH. TSH and TRH production from the hypothalamus and pituitary are negatively regulated by free T4 and free T3.

Thyroid hormone production begins with iodide absorption. It is "oxidised" into iodine in the thyroid cell by iodide trapping, which occurs next. MIT and DIIODYROSINE are the byproducts of this reaction with tyrosine (DIT).

"When MIT and DIT are coupled, T3 is formed, but when two DIT are coupled, T4 is formed. In their bound state with thyroglobulin-7 until secretion, thyroid hormones T3 and T4 are stored. Catalyzes the oxidation, iodination, and coupling reactions are the functions of thyroid peroxidase enzyme. Most of the thyroid hormones T3 and T4 attach to plasma proteins called thyroxine-binding globulin, and thyroxine-binding

prealbumin, and albumin, respectively, once they are released into the blood. In contrast to T3, which attaches mostly to albumin, T4 mostly to thyroxine-binding globulin. They expand the circulating pool and aid in controlling hormone distribution to specific tissue locations because they delay hormonal clearance by binding to plasma proteins. Free T3 and free T4 are the other thyroid hormones that are delivered. Bound and unbound forms coexist peacefully in the marketplace.

T4 is converted to T3 by the enzyme 5' "Deiodenase in the periphery, whereas 5 deiodenase converts 45% of T4" to rT3. Thirty-three percent of T3 is generated by the thyroid gland, with the remaining eighty-seven percent derived from T4.

In terms of half-life, T4 has a longer half-life than T3, but T3 has a longer potency than T4. T4 has a half-life of seven days, while T3 has a half-life of between ten and twenty-four hours.

When it comes to thyroid hormones, the rate of secretion varies from person to person. Thyroxine is produced at "80 to 90 micrograms per day, whereas tri-iodothyronine is" at 4 to 5 microgram per day. When it comes to T3 levels, the blood has a concentration of 0.12 micrograms per deciliter, whereas the T4 level is 8.

### **Research Gap**

Individual correlations between CKD aetiology and thyroid dysfunction could not be examined in "patients with chronic kidney disease who had thyroid dysfunction. Patients on dialysis were not investigated" for thyroid dysfunction since dialysis changes the thyroid profile without affecting CKD.

### **Research Objective & Methodology**

To find out if individuals with chronic renal illness have abnormalities in their thyroid function. To see if there's a connection between thyroid dysfunction and kidney failure severity. Differentiate between primary hypothyroidism and chronic renal disease-related thyroid dysfunction

The current study involved 50 patients who were hospitalized to the Department of Medicine, china Medical College & Hospital, china, from July 2020 to July 2021 and were diagnosed with chronic renal disease under conservative treatment. Simple random sampling is used to select these samples. For the analysis, statistical parameters like mean, SD, and correlations are employed, as well as parametric and non-parametric tests.

All patients gave their consent after being fully informed. Design of the investigation: cross-sectional study that is both observational and exploratory Patients with chronic renal disease must meet the inclusion criteria before they may be considered. Diabetic patients who met the diagnostic criteria for kidney disease but were receiving supportive treatment.

#### **Criteria for Chronic Kidney Disease**

1. "Presence of uraemic symptoms for 3 months or more"
2. "Raised blood urea, serum creatinine and reduced creatinine clearance. "
3. "Ultra sonogram evidence of chronic kidney disease"
4. "Bilateral contracted kidneys — size less than 9 cm. "
5. "Poor cortico-medullary differentiation. "
6. "Supportive laboratory evidence of CKD like anaemia, changes in serum electrolytes, etc., "

#### **Exclusion criteria**

1. "Patients on peritoneal dialysis or hemodialysis"
2. "Nephrotic range of proteinuria"
3. "Hypoalbuminemia"
4. "Other conditions like"
5. "Acute illness"
6. "Diabetes mellitus"
7. "Recent surgery"
8. "Trauma"
9. "Burns"
10. "Liver diseases"
11. "Drugs altering thyroid profile like amiodarone, phenytoin, beta-blocker, dopamine, steroids, estrogen pills and iodine- containing drugs. "

Comprehensive medical history and physical examinations were performed on patients with end-stage renal disease (ESRD) who were receiving conservative treatment. The following studies were carried out.

1. "Urine routine and microscopic examination"
2. "Peripheral smear for anemia"
3. "Blood urea"
4. "Serum Creatinine"
5. "Creatinine clearance (using Cockcroft — Gault formula) "
6. "Serum electrolytes"
7. "Serum calcium, phosphorous and uric acid"
8. "Serum cholesterol"
9. "24 hours urinary protein"
10. "Serum protein (Total protein / albumin / globulin) "
11. "USG abdomen for evidence of chronic kidney disease"

Five millilitres of blood are taken from individuals who meet the aforementioned requirements and sent for thyroid profile testing in non-heparinized serum bottles.

"Components of thyroid profile included in our study"

1. "Serum Triiodothyronine (T3) "
2. "Serum thyroxine (T4) "
3. "Serum thyroid stimulating hormone (TSH) "

"Enzyme Linked Immunosorbent Assay" is used to quantify T3, T4, and TSH.

#### **Data Analysis & Findings**

During our research, we looked at "50 patients with CKD who were on conservative treatment and met the

CKD criteria. Of them, 34 were men and 16 were women, ranging in age from 20 to 68.

Patients under the age of 30 comprised 18% of our study's 50 participants, those between the ages of 31 and 60 included 35 participants, and those 60 and beyond comprised 6 participants, each representing 12 percent of the total. Sixty-eight percent of patients were men, while 32 percent were women in the research.

Our study found that the average duration of CKD symptoms was between 7 and 12 months, ranging from 4 to 30 months. The average time frame was 9.84 months.

Thirty-three out of the fifty patients had a GFR less than 15 ml/minute, or 66 percent. There were 15 patients with GFRs ranging from 15 to 30 ml/minute and 2 patients with GFRs greater than or equal to 30 ml/minute, totaling 4%. Most of the individuals in the research had a creatinine clearance of less than 15 ml/min.

The urea level in the blood ranged from 45 to 184 mg/dl, with a mean of 102.12 mg/dl. Most of the individuals in the research have urea levels between 81 and 120 mg/dl.

In this study, the creatinine levels ranged from 3 to 14 mg/dl, with an average of 7.34 mg/dl found. In the study's patient population, the majority had blood creatinine levels between 4 and 8 milligrams/deciliter.

In 10 patients (20%), the serum calcium levels were determined to be low, whereas they were normal in 24 individuals (30%).

"The TSH values in our study ranged from 0.6-38 micro IU/ml, the mean value being 6.494. Four individuals exhibited values greater than 20 micro IU/ml, while the rest were within the normal range. Three of the high-risk patients were men, whereas one patient was a woman.

The creatinine clearance was less than 15 ml/s in all four of the individuals with primary hypothyroidism. It's a measure of how severe hypothyroid individuals' renal failure is.

"The mean TSH levels in various stages of renal failure were found to be within the normal range in our research of CKD patients with low T3 syndrome. In our research, there was no connection between TSH levels and glomerular filtration rate.

## **Conclusion**

Fifty CKD patients on conservative treatment were included in my research. Sixty-six percent of the individuals in this group had low levels of the thyroid hormone T3. Patients with CKD may benefit from changes in T3 and T4 serum levels because they are protective and help conserve protein. Low T3 and T4 syndrome patients grow in number as the degree of renal failure increases. Patients with chronic renal illness are more likely to have hypothyroidism. With the exception of hypothyroidism, 58 percent of patients have low T3 levels, whereas only 24 percent have low T4 levels. Low T3 syndrome occurs more frequently in CKD patients as they get older. Serum T3 levels were shown to be lower in patients with poor GFR. Thus, GFR and T3 levels are connected in a linear fashion.

## **References**

1. Andrew S. Levey, Josef Coresh, Ethan Balk, Annamaria T. Kausz, Ronald D. Perrone. National Kidney Foundation Practice Guidelines for Chronic Kidney Disease: Evaluation, Classification, and Stratification. *MD Ann Intern Med.* 2003;139:137-147.
2. Joanne M. Bargman, Karl S. Korecki. Chronic kidney disease. In: Dan L. Lango, Anthony S. Fauci, Dennis Kasper et al. *Harrison's Principles of Internal Medicine, Vol. 2, 18th edn., 2011; McGraw Hill, USA, pp. 2289-2293; 2308- 2313.*

3. Feinstein EI, Kaptein EM, Nicoloff JT & Massry SG. Thyroid function in patients with nephrotic syndrome and normal renal function. *American Journal of Nephrology* 1982 2 70-76.
4. Kaptein EM, Quion-Verde H & Massry SG. Hemodynamic effects of thyroid hormone. *Contributions to Nephrology* 1984 41 151-159.
5. Kaptein EM. Thyroid function in renal failure. *Contributions to Nephrology* 1986 50 64-72.
6. Robert W Schrier. Abnormalities in the thyroid gland and hypothalamo pituitary thyroid axis in patients with CKD – *Diseases of the kidney and urinary tract*, eighth edition 2007; volume 3: page number 2518.
7. P Iglesias and J JDí'Ez. Thyroid dysfunction and kidney disease. *European Journal of Endocrinology* (2009) 160: 503-515.
8. MWJ Strachan, BR Walker. Endocrine disease. In: Nicholas A.Boon, Nicki R.Colledge et al. *Davidson's Principle and Practice of Medicine*, 20th edn., 2006; Churchill Livingstone, Elsevier, Philadelphia, pp. 744-754.
9. J. L. Jameson and A. P. Weetman, *Harrison's Principles of Internal Medicine* 18th Edition, Disorders of the Thyroid Gland.
10. Custro N et al. Prospective study on thyroid function anomalies in seriously ill patient. *Ann Ital Med Mt*, 1992; 7:13-8.
11. Degroot. *The thyroid and its diseases*, 6th Edition. Non-Thyroidal illness.
12. Hasegawa K et al. Abnormal response of thyrotrophin and growth hormone to thyrotrophin releasing hormone in chronic renal failure. *ActaEndocrinol*, 1975; 79: 635-43.
13. Ramirez G et al. Thyroid dysfunction in uraemia. Evidence for thyroid and hypophyseal abnormalities. *Ann Inter Med*, 1976; 84: 672-6.
14. Silverberg DS et al. Effect of chronic hemodialysis on thyroid function in chronic renal failure. *Can Med An*, 1973; 109: 282-6.
15. Weissel M et al. Basal and TRH stimulated Thyroid and Pituitary hormones in various degree of renal insufficiency. *ActaEndocrinol*, 1979; 90 23-32.
16. C.CraigTisher, Kirsten M, Madsen. *Anatomy of the Kidney*. In: Brenner and Recters : *The Kidney* 16th edition.
17. K/DOQI Clinical Practice Guidelines For Chronic Kidney Disease:.Evaluation, Classification and Stratification .*Am J Kidney Dis* 39 :S1 S266,200.