

Review Article

Antithyroid Drug Therapy for Graves' Disease and Implications for Recurrence

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Graves' disease (GD) is the most common cause of hyperthyroidism worldwide. Current therapeutic options for GD include antithyroid drugs (ATD), radioactive iodine, and thyroidectomy. ATD treatment is generally well accepted by patients and clinicians due to some advantages including normalizing thyroid function in a short time, hardly causing hypothyroidism, and ameliorating immune disorder while avoiding radiation exposure and invasive procedures. However, the relatively high recurrence rate is a major concern for ATD treatment, which is associated with multiple influencing factors like clinical characteristics, treatment strategies, and genetic and environmental factors. Of these influencing factors, some are modifiable but some are nonmodifiable. The recurrence risk can be reduced by adjusting the modifiable factors as much as possible. The titration regimen for 12–18 months is the optimal strategy of ATD. Levothyroxine administration after successful ATD treatment was not recommended. The addition of immunosuppressive drugs might be helpful to decrease the recurrence rate of GD patients after ATD withdrawal, whereas further studies are needed to address the safety and efficacy. This paper reviewed the current knowledge of ATD treatment and mainly focused on influencing factors for recurrence in GD patients with ATD treatment.

1. Introduction

Graves' disease (GD) is an organ-specific autoimmune disease characterized as overproduction of thyroid hormones in thyroid follicular cells resulting from the stimulation of circulating thyroid-stimulating hormone (TSH) receptor antibodies (TRAb) [1]. It is the most common cause of hyperthyroidism worldwide [1]. Current therapeutic options for GD include antithyroid drugs (ATD), radioactive iodine, and thyroidectomy [1, 2]. ATD treatment has many advantages, including normalizing thyroid function in a short time, hardly causing hypothyroidism, and ameliorating immune disorder while avoiding radiation exposure and invasive procedures, so it is generally well accepted by patients and clinicians [3, 4]. However, the high recurrence rate is a main limitation of ATD treatment, which varies greatly among patients with different treatment strategies, clinical characteristics, and environmental and genetic factors [2, 5–8]. In this paper, we review the current knowledge of ATD treatment and mainly focus on influencing factors for recurrence in GD patients with ATD treatment.

2. Treatment Strategies of ATD and the Recurrence Risk

2.1. Drug Selection. The commonly used ATD include methimazole and propylthiouracil [1]. Carbimazole is another ATD that is available in few regions [9–11]. Carbimazole exerts its pharmacological effect by converting to methimazole, so it has similar efficacy and features as methimazole [9–11]. Previous studies showed that methimazole had a better efficacy and restored the euthyroid state much faster than propylthiouracil, but the recurrence rate after withdrawal was comparable between the two drugs in GD patients [12–14]. The side effects of ATD is common (13%) but generally mild, including rash, pruritus, metallic taste, arthralgia, and liver damage [2]. A meta-analysis of 31 observational studies showed that rash was more common with methimazole treatment, whereas the predominant side effect of propylthiouracil was hepatic involvement [2]. ATD treatment also had some major side effects, including agranulocytosis and severe hepatotoxicity, which are life

threatening but rare (<0.5%) [2]. Methimazole has longer half-life and duration of action and fewer major side effects as compared to propylthiouracil [2, 4]. Thus, methimazole is recommended as the preferred drug for GD patients by ATA guidelines except for pregnant women during the first trimester [2, 4, 15].

2.2. Treatment Regimen. There are two regimens of ATD: titration-block and block-replace regimens [16]. The titration-block regimen means that the ATD dose is titrated from the initial dose to the lowest dose for maintaining a euthyroid state, and the block-replace regimen is initiated with a standard dose of ATD and the addition of levothyroxine [16]. The initial dose of ATD depends on the severity of hyperthyroidism [12]. Patients with mild hyperthyroidism begin with 10–15 mg daily of methimazole, while 20–40 mg daily is given to patients with severe hyperthyroidism [12]. A recent meta-analysis from Cochrane showed that the two regimens had similar recurrence rates, but the block-replace regimens caused relatively more side effects [16].

2.3. Treatment Duration. Just as important, the treatment duration of ATD also impacts the recurrence risk of GD patients [7, 11, 17, 18]. About 20 years ago, most GD patients were treated by ATD for 6 months [11, 17]. Recently, increasing evidence has demonstrated that ATD treatment for 12–18 months leads to a better prognosis than the 6-month treatment [7, 18]. The results from a recent meta-analysis showed that the 12-month titration regimen has a lower recurrence rate than the 6-month regimen, but extending treatment beyond 18 months failed to provide more benefits [16]. In addition, considering the high recurrence rate after drug withdrawal, some studies even advocated a continuous treatment with low-dose ATD for GD patients [19, 20]. They proposed that long-term maintenance of low-dose ATD had a persistent effect on preventing recurrence [19, 20]. However, because of the intermittent blood check and higher medical costs during long-term ATD maintenance, the titration regimen for 12–18 months is still considered as the optimal strategy of ATD.

2.4. Levothyroxine Administration after Successful ATD Treatment. Because increased TSH levels have been incriminated for promoting the production of TRAb, some studies have been conducted to evaluate whether levothyroxine administration after successful ATD treatment could decrease the recurrence risk of hyperthyroidism in GD patients [21–23]. Nevertheless, these studies demonstrated that levothyroxine does not prevent recurrence of hyperthyroidism in GD patients after successful ATD treatment [21, 22]. Mastorakos et al. even observed that levothyroxine administration after successful ATD treatment was associated with increased recurrence risk of GD patients [23]. Thus, levothyroxine administration after successful ATD treatment was not recommended.

2.5. Additional Use of Immunosuppressive Drugs for Standard ATD Treatment. Considering the autoimmune nature of GD, several studies have attempted to evaluate the effect of additional use of immunosuppressive drugs on ATD

treatment in GD patients [24]. The immunosuppressive drugs for GD patients mainly included corticosteroid and noncorticosteroid drugs [25–29]. A recent meta-analysis demonstrated a strong reduction of the recurrence risk when immunosuppressive drugs were added to standard ATD treatment in GD patients, and the reduction of the recurrence risk was similar in studies using corticosteroid and noncorticosteroid immunosuppressive drugs [30]. In this meta-analysis, the overall recurrence rate in GD patients receiving the addition of immunosuppressive drugs was 23.5%, which was significantly lower than 59.1% in GD patients only treated with ATD [30]. Therefore, the addition of immunosuppressive drugs might be helpful to decrease the recurrence rate of GD patients after ATD withdrawal, whereas there are still some problems to mention. First, studies addressing immunosuppressive treatment for GD are always small, single-center, and with low to moderate quality and high risk of bias [25–29]. Second, the side effects of immunosuppressive drugs should have great importance [30]. The administration mode of immunosuppressive drug included oral and local administrations [25–29]. However, the side effects should not be disregarded, no matter the mode [30]. The side effects of corticosteroids include bone abnormalities, metabolic disturbances, and muscle wasting, and rituximab was associated with leucopenia, rash, minor infections, chills, and fever [28, 30]. Therefore, further large-scale randomized controlled trials are needed to address the safety, efficacy, optimal timing, and duration of immunosuppressive drugs in GD patients.

3. Influencing Factors for the Recurrence in GD Patients

3.1. General Situation. The incidence of GD gradually increases with age and then remains stable after the age of 30 [31]. It is generally accepted that younger GD patients have more severe immune disorders [32]. Previous studies showed that younger GD patients have a relatively poor response to ATD and often have a poor prognosis and higher recurrence risk [5, 33, 34]. A study by Allahabadia et al. found that GD patients younger than 40 years had a higher recurrence rate than older patients [34].

Females have a higher incidence of GD than males [31, 35]. The reason for a different incidence in gender is unclear and might be associated with varying sex hormones [36]. Estrogens influenced B-cell function and further regulated the immune system [36]. In GD patients, the increased estradiol level is related to the positivity of TRAb [37]. Although the incidence of GD is higher in women, male GD patients have a higher risk of recurrence after ATD withdrawal [6, 34, 38, 39]. There are some possible explanations for the higher recurrence risk of male GD patients. First, male GD patients had larger goiter size, which was associated with severe immune and biochemical disorders [39]. Second, a family history of thyroid or non-thyroid autoimmune diseases occurred more frequently in male GD patients [39]. So, the higher risk of recurrence in male GD patients might be associated with bigger goiter size

and genetic background. However, this is still a conflicting issue since other studies have inconsistent results [8, 40].

Several studies showed that smokers have a higher recurrence risk than nonsmokers in GD patients after ATD withdrawal [38, 41–43]. Meanwhile, quitting smoking has also been demonstrated to protect against recurrence in GD patients [40]. A previous study showed that smokers had significant higher TRAb levels than nonsmokers at 4 weeks after ATD withdrawal [44]. Therefore, smoking might promote immune disorder and elevate TRAb levels, contributing to the increased recurrence risk. Because there were some inconsistent results, the association between smoking and the recurrence rate still remains uncertain [8].

3.2. Biochemical Parameter. The severity of GD is associated with the recurrence risk in GD patients treated with ATD [8, 45–47]. The biochemical parameters partially represent the severity of GD [8, 45–47]. The key feature in untreated GD is the significant increase in the serum triiodothyronine (T3) level, which is caused by the elevated activity of intrathyroidal type 1 deiodinase [1]. Previous studies showed that the serum T3 levels and free T3 (FT3)/free thyroxine (FT4) ratios at the onset of GD were independent factors for predicting outcomes of ATD treatment in GD patients [8, 45–47]. Patients with mild hyperthyroidism can achieve remission after just treatment with beta blockers [48]. However, patients with higher serum T3 levels and FT3/FT4 ratios have a relatively higher recurrence risk, so they often need a higher initial dose and longer treatment duration [45–47]. Furthermore, a high T3/T4 ratio during ATD withdrawal also predicts a higher recurrence risk in GD patients [4]. The therapy duration should be prolonged in patients with a high T3/T4 ratio even after 12–18 months of ATD treatment.

In addition, the serum TSH level also should get more attention. As is known, the thyroid hormone can inhibit TSH via a negative feedback mechanism [49]. In some GD patients, the TSH level still sustained suppression and failed to return to a normal range with the normalization of thyroid hormone after ATD treatment [38, 49]. Previous studies have demonstrated that TSH suppression after drug withdrawal was a predictor for the recurrence of GD [8, 38, 50]. These studies suggested that GD patients with delayed TSH restoration should receive prolonged ATD treatment until their TSH levels reach the normal range.

3.3. Immune Parameters. As mentioned before, GD results from the overactivated TSH receptor in the thyroid follicular cells by TRAb [1]. TRAb is positive in about 95% of patients with newly diagnosed GD, and the higher TRAb levels hint a severe immune disorder [1, 5, 14, 51, 52]. In recent years, with increased accuracy of assay, TRAb has been supported as a useful predictive factor for the outcome of ATD treatment by many studies [5, 14, 51, 52]. Patients with higher TRAb levels at the time of GD diagnosis have significantly increased recurrence risk, while TRAb-negative patients often have a better prognosis and are prone to long-term remission [5, 14, 52]. Furthermore, a shift from positive to negative in TRAb may imply the alleviated immune disorder

following ATD treatment in GD patients [41, 43, 50]. The TRAb level at ATD withdrawal was also associated with the prognosis of GD patients [41, 43, 50]. The recurrence risk was higher in TRAb-positive GD patients at the time of drug withdrawal [41, 43, 50]. In addition, TRAb can be distinguished from the stimulating (TSAb) and blocking (TBAb) properties by using new assay techniques [53]. Antibodies of GD patients are predominantly TSAb [1]. Recently, TSAb has been demonstrated to have a superior predictive value for recurrence risk than TRAb in GD patients treated with ATD [53–55].

GD and Hashimoto's thyroiditis are the two main autoimmune thyroid diseases [56]. Prior epidemiologic studies have indicated that GD is possibly concomitant with Hashimoto's thyroiditis [57]. GD patients with Hashimoto's thyroiditis tend to be in remission after ATD treatment due to the progressive damage induced by Hashimoto's thyroiditis [57]. Since the positivity of peroxidase autoantibodies (TPOAb) and/or thyroglobulin antibody (TgAb) is the main feature of Hashimoto's thyroiditis, some studies have evaluated the association between the positivity of TgAb/TPOAb and the recurrence risk in GD patients [58, 59]. The reason for the inconsistent results might be that both TPOAb and TgAb also could be observed in some GD patients without Hashimoto's thyroiditis [59].

3.4. Goiter Size. Large goiter size is a main clinical manifestation of GD patients [5, 8]. Previous studies have demonstrated that goiter size was a major predictor of a higher recurrence risk in GD patients after ATD withdrawal [5, 8]. A 5-year follow-up trial showed that patients with normal or mild goiter size had higher remission rates than patients with large goiters [49]. Moreover, GD patients with significantly decreased goiter sizes after ATD treatment also tend to have higher remission rates [41, 54]. These studies suggested that enlarged goiter size at the time of GD diagnosis and drug withdrawal is associated with a higher recurrence risk.

3.5. Graves' Orbitopathy. Graves' orbitopathy is present in around 30% of patients at the time of diagnosis of GD [40, 60]. The presence of Graves' orbitopathy often suggested a worse disorder of the immune system [60]. Previous studies have shown that patients with Graves' orbitopathy have a higher recurrence risk of GD after ATD withdrawal [38]. A study by Eckstein et al. even found that the remission rate of GD patients with severe Graves' orbitopathy was just 7% [61]. Even with the higher recurrence rate, ATD treatment is still a preferred therapeutic option for GD patients with Graves' orbitopathy due to a better outcome for Graves' orbitopathy, which might be associated with the stable euthyroid status and decreased levels of TRAb and inflammatory markers [62–65]. Recent studies have shown that a prolonged low dose of ATD treatment contributed to a better outcome in GD patients with Graves' orbitopathy [63, 66].

3.6. Genetic Factors. Genetic factors participate in the pathogenesis of GD [67]. Recent studies found that both cytotoxic T-lymphocyte-associated factor 4 (*CTLA4*) rs231775 and

rs231779 polymorphisms were associated with recurrence in GD patients after ATD withdrawal in Asians, while no association was observed in Caucasians [5, 41, 68]. In Caucasian patients with GD, the recurrence risk after ATD withdrawal was reported to be related to the polymorphisms of HLA *DQA2*, HLA *DRB1*03*, and HLA *DQB1*02* [5]. The HLA region contains some immune response genes and these HLA polymorphisms might influence the outcome of GD patients by regulating the immune system. In addition, a few studies also explored the association between the recurrence risk and polymorphisms, including T393C SNP of Galphas gene (*GNAS1*), CD40 (rs745307, rs11569309, and rs3765457), and E33SNP of thyroglobulin (*Tg*) (*Tg* E33SNP) in GD patients after ATD withdrawal [41, 42, 69].

3.7. Environmental Factors. The onset of GD is induced by some environmental factors in the individuals with the predisposed gene [70]. Stress is one of the environmental factors, and the association between stress and the recurrence of GD patients after ATD treatment was supported by most studies [71–73]. A Japanese study showed that the 2004 earthquake was associated with the recurrence of GD in patients [74]. In a prospective study that evaluated the recurrence risk of GD in patients, a total stress score of big life events was significantly higher in the recurrence group than in the remission group [75]. Another study showed that the recurrence group experienced more stressful events than did remission patients, and the overall number of stressful events was correlated with the number of recurrences [71]. It is worth mentioning that psychosocial stress is an important part of a stressful event. Previous studies showed that GD patients with psychiatric disorders such as depression and hypochondriasis had a higher recurrence risk than GD patients without such disorders [75–77]. Thus, reducing stress as much as possible is an important way to improve the prognosis of GD patients with ATD treatment.

Iodine intake is another environmental factor [78]. Iodine is a major substrate for the synthesis of thyroid hormone [78]. In thyrocytes, increased iodine content promoted ATD degradation and reduced ATD uptake [79]. Some previous studies showed that iodine supplementation elevated the recurrence rate of GD [80]. The administration of pharmacological doses of iodine caused the onset of hyperthyroidism in euthyroid GD patients after ATD withdrawal [81]. However, epidemiologic studies showed that the recurrence rate in iodine-sufficient regions is not higher than that in iodine-deficient regions in GD patients after ATD withdrawal [40, 82, 83]. In a recent Korean study, excessive iodine did not influence the outcomes of GD [84]. The study suggested that dietary iodine restriction might not be necessary for GD patients after ATD withdrawal [84]. In addition, a previous study showed that a dietary change from a low-iodine to a high-iodine intake increased the recurrence rate in GD patients after ATD treatment [85]. Thus, these results might suggest that only a sudden increase in iodine intake induces the recurrence of GD. Further large-scale intervention studies are needed to evaluate the effect of iodine uptake on the recurrence risk in GD patients after ATD withdrawal.

In addition, baseline vitamin D and selenium levels might impact the outcome of GD patients after ATD treatment. Vitamin D has been demonstrated as an immune modulator [86]. An animal study showed that vitamin D regulated TSH receptor immunization in BALB/c mice [87]. And vitamin D analog also exhibited the inhibitory effect on inflammatory response in human thyroid cells and T cells [88]. ATD treatment caused a greater decline in TRAb in GD patients with normal vitamin D levels compared to that in GD patients with decreased vitamin D levels [89]. Moreover, a decreased baseline vitamin D level was associated with greater thyroid goiter in female patients with newly onset GD [90]. Selenium is another component element of thyroid function [91, 92]. As a basic component of glutathione peroxidase and iodothyronine selenodeiodinase, selenium deficiency might impact the conversion of T4 to T3 and the production of free radicals [91, 93]. The association between inadequate selenium supply and GD has been found by many studies [91, 92]. Furthermore, decreased serum selenium levels were also related to severe immune disorders and the incidence of Graves' orbitopathy [94, 95]. High serum selenium levels were also shown to be associated with higher remission rates in GD patients [96]. Recently, more studies have found that selenium supplementation can enhance biochemical restoration of hyperthyroidism and improve the remission rate of GD patients [93, 97]. It is worth noting that there have been relatively fewer studies to evaluate the association between the recurrence risk and the serum levels of vitamin D and selenium, and they were also small-scale, single-center studies. Further large-scale prospective studies are needed to observe this association and whether the administration of vitamin D and selenium can bring benefit.

4. Treatment Options for Recurrence GD Patients after ATD Withdrawal

The high recurrence rate is a major drawback of ATD therapy, and patients with recurrent GD often have a much higher recurrence risk than average [54, 82]. Most clinicians will recommend radioactive iodine or thyroidectomy for recurrent GD patients [98]. However, some recent studies have found that compared with radioactive iodine or thyroidectomy, the prolonged low-dose ATD treatment retained a stable euthyroid state while minimizing the risk of side effects [99, 100]. A recent prospective clinical study showed that ATD treatment also reached a higher remission rate in recurrent GD patients, and a much lower discontinuation drug dose of methimazole (2.5 mg qod) increased the rate of permanent remission [99]. In a recent study, during a long-term follow-up (up to 7 years), prolonged low-dose methimazole treatment was safe and effective and with fewer complications and less expense in recurrent GD patients [100]. It also contributes to a better outcome of Graves' orbitopathy and a lower frequency of thyroid dysfunction than radioactive iodine [100]. Thus, prolonged low-dose methimazole treatment might be a good alternative for recurrent GD patients who resist radioactive iodine or thyroidectomy.

5. Conclusions

Recurrence in GD patients with ATD treatment is associated with multiple influential factors such as clinical characteristics, treatment strategies, and genetic and environmental factors. Of these influencing factors, some are modifiable but some are nonmodifiable. The recurrence risk can be reduced by adjusting the modifiable factors as much as possible. If the recurrence evaluation based on the nonmodifiable factors strongly suggests a high risk of recurrence, a definitive treatment such as radioactive iodine or thyroidectomy is considered as an appropriate therapeutic approach. However, prolonged low-dose methimazole treatment might be a good alternative for GD patients with high recurrence risk due to its safety and efficacy. The addition of immunosuppressive drugs might be helpful to decrease the recurrence rate of GD patients after ATD withdrawal, whereas further studies are needed to address the safety and efficacy. Further large-scale prospective studies are also needed to observe whether the administration of vitamin D and selenium can bring benefit.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

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References

- [1] S. De Leo, S. Y. Lee, and L. E. Braverman, "Hyperthyroidism," *Lancet*, vol. 388, no. 10047, pp. 906–918, 2016.
- [2] V. Sundaresh, J. P. Brito, Z. Wang et al., "Comparative effectiveness of therapies for Graves' hyperthyroidism: a systematic review and network meta-analysis," *The Journal of Clinical Endocrinology and Metabolism*, vol. 98, no. 9, pp. 3671–3677, 2013.
- [3] C. Ma, J. Xie, H. Wang, J. Li, and S. Chen, "Radioiodine therapy versus antithyroid medications for Graves' disease," *Cochrane Database of Systematic Reviews*, vol. 2, article CD010094, 2016.
- [4] D. S. Cooper, "Antithyroid drugs," *The New England Journal of Medicine*, vol. 352, no. 9, pp. 905–917, 2005.
- [5] X. G. Vos, E. Ender, A. H. Zwinderman, J. G. Tijssen, and W. M. Wiersinga, "Predicting the risk of recurrence before the start of antithyroid drug therapy in patients with Graves' hyperthyroidism," *The Journal of Clinical Endocrinology and Metabolism*, vol. 101, no. 4, pp. 1381–1389, 2016.
- [6] K. W. Kim, Y. J. Park, T. Y. Kim, D. J. Park, K. S. Park, and B. Y. Cho, "Susceptible alleles of the CD40 and CTLA-4 genes are not associated with the relapse after antithyroid withdrawal in Graves' disease," *Thyroid*, vol. 17, no. 12, pp. 1229–1234, 2007.
- [7] H. Allannic, R. Fauchet, J. Orgiazzi et al., "Antithyroid drugs and Graves' disease: a prospective randomized evaluation of the efficacy of treatment duration," *Journal of Clinical Endocrinology and Metabolism*, vol. 70, no. 3, pp. 675–679, 1990.
- [8] L. Liu, H. Lu, Y. Liu, C. Liu, and C. Xun, "Predicting relapse of Graves' disease following treatment with antithyroid drugs," *Experimental and Therapeutic Medicine*, vol. 11, no. 4, pp. 1443–1458, 2016.
- [9] R. V. Garcia-Mayor, C. Paramo, R. Luna Cano, L. F. Pérez Mendez, J. C. Galofré, and A. Andrade, "Antithyroid drug and Graves' hyperthyroidism. Significance of treatment duration and TRAb determination on lasting remission," *Journal of Endocrinological Investigation*, vol. 15, no. 11, pp. 815–820, 1992.
- [10] J. Duprey, M. F. Louis, B. Ducornet et al., "Improvement of the prognosis of Basedow's disease by using high doses of carbimazole," *Presse médicale (Paris, France: 1983)*, vol. 17, no. 22, pp. 1124–1127, 1988.
- [11] A. P. Weetman, S. Ratanachaiyavong, and G. W. Middleton, "Prediction of outcome in Graves' disease after carbimazole treatment," *Quarterly Journal of Medicine*, vol. 59, no. 228, pp. 409–419, 1986.
- [12] H. Nakamura, J. Y. Noh, K. Itoh, S. Fukata, A. Miyauchi, and N. Hamada, "Comparison of methimazole and propylthiouracil in patients with hyperthyroidism caused by graves' disease," *The Journal of Clinical Endocrinology and Metabolism*, vol. 92, no. 6, pp. 2157–2162, 2007.
- [13] C. T. He, A. T. Hsieh, D. Pei et al., "Comparison of single daily dose of methimazole and propylthiouracil in the treatment of Graves' hyperthyroidism," *Clinical Endocrinology*, vol. 60, no. 6, pp. 676–681, 2004.
- [14] M. C. Peixoto, A. Buescu, M. R. B. Gonçalves, M. de Souza Albarnaz, C. M. Coeli, and M. Vaisman, "Antithyroid drugs for the treatment of Graves disease: a randomized clinical trial," *Endocrinologist*, vol. 16, no. 6, pp. 344–348, 2006.
- [15] D. S. Ross, H. B. Burch, D. S. Cooper et al., "2016 American thyroid association guidelines for diagnosis and management of hyperthyroidism and other causes of thyrotoxicosis," *Thyroid*, vol. 26, no. 10, pp. 1343–1421, 2016.
- [16] P. Abraham, A. Avenell, S. C. McGeoch, L. F. Clark, and J. S. Bevan, "Antithyroid drug regimen for treating Graves' hyperthyroidism," *Cochrane Database of Systematic Reviews*, vol. 20, no. 1, p. CD003420, 2010.
- [17] P. A. Dahlberg, F. A. Karlsson, B. Lindstrom, and L. Wide, "Studies of thyroid hormone and methimazole levels in patients with Graves' disease on a standardized anti-thyroid drug regimen," *Clinical Endocrinology*, vol. 14, no. 6, pp. 555–562, 1981.
- [18] C. Liu, Y. Duan, and X. Wu, "Prospective study on the relationship between treatment duration of antithyroid drug and remission rate of Graves' disease," *Zhonghua Yi Xue Za Zhi*, vol. 80, no. 11, pp. 835–837, 2000.
- [19] F. Morales García, A. Cayuela, N. García Hernández et al., "Long-term maintenance of low-dose antithyroid drugs versus drug withdrawal in patients with Graves' hyperthyroidism," *Endocrinología y Nutrición*, vol. 55, no. 3, pp. 123–131, 2008.
- [20] E. Mazza, M. Carlini, D. Flecchia et al., "Long-term follow-up of patients with hyperthyroidism due to Graves' disease treated with methimazole. Comparison of usual treatment schedule with drug discontinuation vs continuous treatment with low methimazole doses: a retrospective study,"

- Journal of Endocrinological Investigation*, vol. 31, no. 10, pp. 866–872, 2008.
- [21] D. Glinoe, P. De Nayer, and M. Bex, “Effects of L-thyroxine administration, TSH-receptor antibodies and smoking on the risk of recurrence in Graves’ hyperthyroidism treated with antithyroid drugs: a double-blind prospective randomized study,” *European Journal of Endocrinology*, vol. 144, no. 5, pp. 475–483, 2001.
 - [22] R. Hoermann, B. Quadbeck, U. Roggenbuck et al., “Relapse of Graves’ disease after successful outcome of antithyroid drug therapy: results of a prospective randomized study on the use of levothyroxine,” *Thyroid*, vol. 12, no. 12, pp. 1119–1128, 2002.
 - [23] G. Mastorakos, A. G. Doufas, E. Mantzos, J. Mantzos, and D. A. Koutras, “T4 but not T3 administration is associated with increased recurrence of Graves’ disease after successful medical therapy,” *Journal of Endocrinological Investigation*, vol. 26, no. 10, pp. 979–984, 2003.
 - [24] A. Antonelli, S. M. Ferrari, A. Corrado, A. Di Domenicantonio, and P. Fallahi, “Autoimmune thyroid disorders,” *Autoimmunity Reviews*, vol. 14, no. 2, pp. 174–180, 2015.
 - [25] G. Obiols Alfonso, J. Mesa Manteca, R. Simo Canonge, L. García Pascual, C. García Alonso, and J. M. Tresánchez Trías, “Effect of carbimazole and dexamethasone on the TSH bond (TBI) inhibiting immunoglobulins in Graves-Basedow disease,” *Revista Clínica Española*, vol. 183, no. 8, pp. 401–404, 1988.
 - [26] L. Chen, H. Q. Wang, Y. Y. Gao et al., “Comparison of methimazole/hydrocortisone ointment with oral methimazole in patients with Graves disease: a prospective, randomized, open-label, parallel-group, 18-month study,” *Current Therapeutic Research, Clinical and Experimental*, vol. 69, no. 4, pp. 305–317, 2008.
 - [27] S. Kubota, H. Ohye, E. Nishihara et al., “Effect of high dose methylprednisolone pulse therapy followed by oral prednisolone administration on the production of anti-TSH receptor antibodies and clinical outcome in Graves’ disease,” *Endocrine Journal*, vol. 52, no. 6, pp. 735–741, 2005.
 - [28] D. El Fassi, C. H. Nielsen, S. J. Bonnema, H. C. Hasselbalch, and L. Hegedus, “B lymphocyte depletion with the monoclonal antibody rituximab in Graves’ disease: a controlled pilot study,” *The Journal of Clinical Endocrinology and Metabolism*, vol. 92, no. 5, pp. 1769–1772, 2007.
 - [29] B. Huszno, M. Trofimiuk, F. Golkowski, T. Plinta, and Z. Szybinski, “Assessment of early immunosuppressive therapy in the prevention of complications of Graves’ disease,” *Przegląd Lekarski*, vol. 61, no. 8, pp. 868–871, 2004.
 - [30] T. Struja, L. Guebelin, A. Kutz, H. Fehlberg, B. Mueller, and P. Schuetz, “Does immunosuppressive therapy improve outcomes in Graves’ disease? A systematic review and meta-analysis,” *Thyroid*, vol. 26, no. 5, pp. 634–640, 2016.
 - [31] A. Carle, I. B. Pedersen, N. Knudsen et al., “Epidemiology of subtypes of hyperthyroidism in Denmark: a population-based study,” *European Journal of Endocrinology*, vol. 164, no. 5, pp. 801–809, 2011.
 - [32] M. Kourime, S. McGowan, M. Al-Towati et al., “Outcome of thyrotoxicosis in childhood and adolescence in a geographically define area; a 24-year experience,” in *Hormone Research in Paediatrics*, M. Kourime, Ed., vol. 84, pp. 551–552, Abderrahim Harouchi Hôpital, Casablanca, Morocco, 2015.
 - [33] T. Yamada, T. Aizawa, Y. Koizumi, I. Komiya, K. Ichikawa, and K. Hashizume, “Age-related therapeutic response to antithyroid drug in patients with hyperthyroid Graves’ disease,” *Journal of the American Geriatrics Society*, vol. 42, no. 5, pp. 513–516, 1994.
 - [34] A. Allahabadia, J. Daykin, R. L. Holder, M. C. Sheppard, S. C. Gough, and J. A. Franklyn, “Age and gender predict the outcome of treatment for Graves’ hyperthyroidism,” *The Journal of Clinical Endocrinology and Metabolism*, vol. 85, no. 3, pp. 1038–1042, 2000.
 - [35] M. L. Tanda, E. Piantanida, L. Liparulo et al., “Prevalence and natural history of Graves’ orbitopathy in a large series of patients with newly diagnosed Graves’ hyperthyroidism seen at a single center,” *The Journal of Clinical Endocrinology and Metabolism*, vol. 98, no. 4, pp. 1443–1449, 2013.
 - [36] J. A. Da Silva, “Sex hormones, glucocorticoids and autoimmunity: facts and hypotheses,” *Annals of the Rheumatic Diseases*, vol. 54, no. 1, pp. 6–16, 1995.
 - [37] L. O. Chailurkit, W. Aekplakorn, and B. Ongphiphadhanakul, “The relationship between circulating estradiol and thyroid autoimmunity in males,” *European Journal of Endocrinology*, vol. 170, no. 1, pp. 63–67, 2014.
 - [38] P. Anagnostis, F. Adamidou, S. A. Polyzos et al., “Predictors of long-term remission in patients with Graves’ disease: a single center experience,” *Endocrine*, vol. 44, no. 2, pp. 448–453, 2013.
 - [39] F. Magri, F. Zerbini, M. Gaiti et al., “Gender influences the clinical presentation and long-term outcome of Graves’ disease,” *Endocrine Practice*, vol. 22, no. 11, pp. 1336–1342, 2016.
 - [40] E. Mohlin, H. Filipsson Nystrom, and M. Eliasson, “Long-term prognosis after medical treatment of Graves’ disease in a northern Swedish population 2000–2010,” *European Journal of Endocrinology*, vol. 170, no. 3, pp. 419–427, 2014.
 - [41] P. W. Wang, I. Y. Chen, S. H. Juo, E. Hsi, R. T. Liu, and C. J. Hsieh, “Genotype and phenotype predictors of relapse of Graves’ disease after antithyroid drug withdrawal,” *European Thyroid Journal*, vol. 1, no. 4, pp. 251–258, 2013.
 - [42] J. Y. Hsiao, M. C. Hsieh, K. J. Tien, S. C. Hsu, S. J. Shin, and S. R. Lin, “Association between a C/T polymorphism in exon 33 of the thyroglobulin gene is associated with relapse of Graves’ hyperthyroidism after antithyroid withdrawal in Taiwanese,” *Journal of Clinical Endocrinology and Metabolism*, vol. 92, no. 8, pp. 3197–3201, 2007.
 - [43] D. Glinoe, “Clinical epidemiology of Graves’ disease: a multicentric prospective study in Belgium,” *Revue Médicale de Bruxelles*, vol. 21, no. 4, pp. 296–299, 2000.
 - [44] B. Quadbeck, U. Roggenbuck, O. E. Janssen et al., “Impact of smoking on the course of Graves’ disease after withdrawal of antithyroid drugs,” *Experimental and Clinical Endocrinology and Diabetes*, vol. 114, no. 8, pp. 406–411, 2006.
 - [45] Z. Gerenova and Y. Vulkov, “Clinical significance of serum T3/T4 ratio in the course of antithyroid drug therapy in Graves’ disease patients,” *Bulletin Médical*, vol. 8, no. 2, pp. 11–13, 2000.
 - [46] J. Gerenova, I. Tzvetanova, J. Valkov, and V. Tzoneva, “Changes in serum levels of free T3 and free T4 and ratio FT3/FT4 during treatment of Graves’ disease with methimazole,” *Bulgarian Medicine*, vol. 8, no. 6, pp. 9–11, 2000.
 - [47] C. M. Khanna, L. R. Shankar, C. B. Jaggi, J. K. Bansal, and P. Chugh, “Predictor of outcome of hyperthyroidism due to Graves’ disease: serum triiodothyronine/thyroxine ratio,” *The Journal of the Association of Physicians of India*, vol. 44, no. 2, pp. 98–101, 1996.

- [48] J. L. Codaccioni, J. Orgiazzi, P. Blanc, M. Pugeat, R. Roulier, and P. Carayon, "Lasting remissions in patients treated for Graves' hyperthyroidism with propranolol alone: a pattern of spontaneous evolution of the disease," *The Journal of Clinical Endocrinology and Metabolism*, vol. 67, no. 4, pp. 656–662, 1988.
- [49] P. Laurberg, "Remission of Graves' disease during antithyroid drug therapy. Time to reconsider the mechanism?" *European Journal of Endocrinology*, vol. 155, no. 6, pp. 783–786, 2006.
- [50] M. Matsumoto, J. Yoshimura Noh, K. Mukasa et al., "Predictors of recurrence of Graves' disease after medical treatment," *Thyroid*, vol. 23, no. S1, p. A101, 2013.
- [51] S. Hwang, Y. Cho, D. Y. Shin, M. K. Song, and E. J. Lee, "High cut-off value of a chimeric TSH receptor (Mc4)-based bioassay may improve prediction of relapse in Graves' disease for 12 months," *Endocrine*, vol. 48, no. 1, pp. 89–95, 2015.
- [52] G. Barbesino and Y. Tomer, "Clinical review: clinical utility of TSH receptor antibodies," *The Journal of Clinical Endocrinology and Metabolism*, vol. 98, no. 6, pp. 2247–2255, 2013.
- [53] C. Giuliani, D. Cerrone, N. Harii et al., "A TSHR-LH/CGR chimera that measures functional thyroid-stimulating autoantibodies (TSAb) can predict remission or recurrence in Graves' patients undergoing antithyroid drug (ATD) treatment," *The Journal of Clinical Endocrinology and Metabolism*, vol. 97, no. 7, pp. E1080–E1087, 2012.
- [54] X. Liu, B. Shi, and H. Li, "Valuable predictive features of relapse of Graves' disease after antithyroid drug treatment," *Annales d'endocrinologie*, vol. 76, no. 6, pp. 679–683, 2015.
- [55] H. Kwon, W. Kim, E. Jang et al., "Usefulness of thyroid stimulating antibody at the time of antithyroid drug withdrawal in predicting relapse of Graves' hyperthyroidism," *Thyroid*, vol. 25, no. S1, p. A11, 2015.
- [56] O. Eber and W. Langsteger, "Clinical aspects of autoimmune thyroid diseases," *Acta Medica Austriaca*, vol. 21, no. 1, pp. 1–7, 1994.
- [57] T. Sato, I. Takata, T. Taketani, K. Saida, and H. Nakajima, "Concurrence of Graves' disease and Hashimoto's thyroiditis," *Archives of Disease in Childhood*, vol. 52, no. 12, pp. 951–955, 1977.
- [58] M. Stefanic and I. Karner, "Thyroid peroxidase autoantibodies are associated with a lesser likelihood of late reversion to hyperthyroidism after successful non-ablative treatment of Graves' disease in Croatian patients," *Journal of Endocrinological Investigation*, vol. 37, no. 1, pp. 71–77, 2014.
- [59] P. Morosini, G. Arnaldi, A. Tacaliti et al., "Antithyroid antibody tests for hyperthyroidism. Analysis of 315 patients," *Recenti Progressi in Medicina*, vol. 85, no. 11, pp. 521–525, 1994.
- [60] P. Laurberg, B. Nygaard, S. Andersen et al., "Association between TSH-receptor autoimmunity, hyperthyroidism, goitre, and orbitopathy in 208 patients included in the remission induction and sustenance in Graves' disease study," *Journal of Thyroid Research*, vol. 2014, Article ID 165487, 6 pages, 2014.
- [61] A. K. Eckstein, H. Lax, C. Losch et al., "Patients with severe Graves' ophthalmopathy have a higher risk of relapsing hyperthyroidism and are unlikely to remain in remission," *Clinical Endocrinology*, vol. 67, no. 4, pp. 607–612, 2007.
- [62] D. Y. Chen, P. F. Schneider, X. S. Zhang, X. Y. Luo, Z. M. He, and T. H. Chen, "Changes in Graves' ophthalmopathy after radioiodine and anti-thyroid drug treatment of Graves' disease from 2 prospective, randomized, open-label, blinded end point studies," *Experimental and Clinical Endocrinology & Diabetes*, vol. 122, no. 1, pp. 1–6, 2014.
- [63] P. Laurberg, D. C. Berman, S. Andersen, and I. Bulow Pedersen, "Sustained control of Graves' hyperthyroidism during long-term low-dose antithyroid drug therapy of patients with severe Graves' orbitopathy," *Thyroid*, vol. 21, no. 9, pp. 951–956, 2011.
- [64] P. Laurberg, G. Wallin, L. Tallstedt, M. Abraham-Nordling, G. Lundell, and O. Tørring, "TSH-receptor autoimmunity in Graves' disease after therapy with anti-thyroid drugs, surgery, or radioiodine: a 5-year prospective randomized study," *European Journal of Endocrinology*, vol. 158, no. 1, pp. 69–75, 2008.
- [65] A. Antonelli, S. M. Ferrari, A. Corrado, E. Ferrannini, and P. Fallahi, "Increase of interferon-gamma inducible CXCL9 and CXCL11 serum levels in patients with active Graves' disease and modulation by methimazole therapy," *Thyroid*, vol. 23, no. 11, pp. 1461–1469, 2013.
- [66] L. Elbers, M. Mourits, and W. Wiersinga, "Outcome of very long-term treatment with antithyroid drugs in Graves' hyperthyroidism associated with Graves' orbitopathy," *Thyroid*, vol. 21, no. 3, pp. 279–283, 2011.
- [67] M. Marino, F. Latrofa, F. Menconi, L. Chiovato, and P. Vitti, "Role of genetic and non-genetic factors in the etiology of Graves' disease," *Journal of Endocrinological Investigation*, vol. 38, no. 3, pp. 283–294, 2015.
- [68] P. W. Wang, I. Y. Chen, R. T. Liu, C. J. Hsieh, E. Hsi, and S. H. Juo, "Cytotoxic T lymphocyte-associated molecule-4 gene polymorphism and hyperthyroid Graves' disease relapse after antithyroid drug withdrawal: a follow-up study," *The Journal of Clinical Endocrinology and Metabolism*, vol. 92, no. 7, pp. 2513–2518, 2007.
- [69] D. Glowacka, C. Loesch, K. T. Johnson et al., "The T393C polymorphism of the Galphas gene (GNAS1) is associated with the course of Graves' disease," *Hormone and Metabolic Research = Hormon- und Stoffwechselforschung = Hormones et métabolisme*, vol. 41, no. 6, pp. 430–435, 2009.
- [70] R. Vita, D. Lapa, F. Trimarchi, and S. Benvenega, "Certain HLA alleles are associated with Graves' disease (GD) in patients in whom hyperthyroidism is preceded by stressful events (SE)," in *Thyroid*, R. Vita, D. Lapa, F. Trimarchi and S. Benvenega, Eds., vol. 25, p. A205, Department of Clinical and Experimental Medicine, University of Messina, Messina, Italy, 2015.
- [71] R. Vita, D. Lapa, F. Trimarchi, and S. Benvenega, "Stress triggers the onset and the recurrences of hyperthyroidism in patients with Graves' disease," *Endocrine*, vol. 48, no. 1, pp. 254–263, 2015.
- [72] L. Chiovato, M. Marino, G. Perugi et al., "Chronic recurrent stress due to panic disorder does not precipitate Graves' disease," *Journal of Endocrinological Investigation*, vol. 21, no. 11, pp. 758–764, 1998.
- [73] G. Effraimidis, J. G. Tijssen, J. F. Brosschot, and W. M. Wiersinga, "Involvement of stress in the pathogenesis of autoimmune thyroid disease: a prospective study," *Psychoneuroendocrinology*, vol. 37, no. 8, pp. 1191–1198, 2012.
- [74] K. Kamoi, M. Tanaka, T. Ikarashi, and M. Miyakoshi, "Effect of the 2004 mid Niigata prefecture earthquake on patients with endocrine disorders," *Endocrine Journal*, vol. 53, no. 4, pp. 511–521, 2006.

- [75] A. Fukao, J. Takamatsu, Y. Murakami et al., "The relationship of psychological factors to the prognosis of hyperthyroidism in antithyroid drug-treated patients with Graves' disease," *Clinical Endocrinology*, vol. 58, no. 5, pp. 550–555, 2003.
- [76] A. Fukao, J. Takamatsu, S. Kubota, A. Miyauchi, and T. Hanafusa, "The thyroid function of Graves' disease patients is aggravated by depressive personality during antithyroid drug treatment," *BioPsychoSocial Medicine*, vol. 5, no. 1, p. 9, 2011.
- [77] K. Yoshiuchi, H. Kumano, S. Nomura et al., "Psychosocial factors influencing the short-term outcome of antithyroid drug therapy in Graves' disease," *Psychosomatic Medicine*, vol. 60, no. 5, pp. 592–596, 1998.
- [78] F. Azizi, "Environmental iodine intake affects the response to methimazole in patients with diffuse toxic goiter," *The Journal of Clinical Endocrinology and Metabolism*, vol. 61, no. 2, pp. 374–377, 1985.
- [79] T. Nakashima, A. Taurog, and G. Riesco, "Mechanism of action of thioureylenes antithyroid drugs: factors affecting intrathyroidal metabolism of propylthiouracil and methimazole in rats," *Endocrinology*, vol. 103, no. 6, pp. 2187–2197, 1978.
- [80] W. D. Alexander, R. M. Harden, D. A. Koutras, and E. Wayne, "Influence of iodine intake after treatment with antithyroid drugs," *Lancet*, vol. 2, no. 7418, pp. 866–868, 1965.
- [81] E. Roti, E. Gardini, R. Minelli et al., "Effects of chronic iodine administration on thyroid status in euthyroid subjects previously treated with antithyroid drugs for Graves' hyperthyroidism," *The Journal of Clinical Endocrinology and Metabolism*, vol. 76, no. 4, pp. 928–932, 1993.
- [82] P. W. Wang, R. T. Liu, S. C. Tung et al., "Outcome of Graves' disease after antithyroid drug treatment in Taiwan," *Journal of the Formosan Medical Association*, vol. 97, no. 9, pp. 619–625, 1998.
- [83] N. B. Tutuncu, T. Tutuncu, A. Ozgen, and T. Erbas, "Long-term outcome of Graves' disease patients treated in a region with iodine deficiency: relapse rate increases in years with thionamides," *Journal of the National Medical Association*, vol. 98, no. 6, pp. 926–930, 2006.
- [84] S. M. Park, Y. Y. Cho, J. Y. Joung, S. Y. Sohn, S. W. Kim, and J. H. Chung, "Excessive iodine intake does not increase the recurrence rate of Graves' disease after withdrawal of the antithyroid drug in an iodine-replete area," *European Thyroid Journal*, vol. 4, no. 1, pp. 36–42, 2015.
- [85] B. L. Solomon, J. E. Evaul, K. D. Burman, and L. Wartofsky, "Remission rates with antithyroid drug therapy: continuing influence of iodine intake?" *Annals of Internal Medicine*, vol. 107, no. 4, pp. 510–512, 1987.
- [86] F. Baeke, T. Takiishi, H. Korf, C. Gysemans, and C. Mathieu, "Vitamin D: modulator of the immune system," *Current Opinion in Pharmacology*, vol. 10, no. 4, pp. 482–496, 2010.
- [87] A. Misharin, M. Hewison, C. R. Chen et al., "Vitamin D deficiency modulates Graves' hyperthyroidism induced in BALB/c mice by thyrotropin receptor immunization," *Endocrinology*, vol. 150, no. 2, pp. 1051–1060, 2009.
- [88] E. Borgogni, E. Sarchielli, M. Sottili et al., "Elocalcitol inhibits inflammatory responses in human thyroid cells and T cells," *Endocrinology*, vol. 149, no. 7, pp. 3626–3634, 2008.
- [89] D. Shin, S. Hwang, D. Kim, K. Kim, and E. Lee, "Baseline vitamin D level could be a short-term prognostic marker in patients with Graves' disease," in *Thyroid*, D. Shin, S. Hwang, D. Kim and E. Lee, Eds., vol. 21, p. A48, Department of Internal Medicine, Yonsei University, College of Medicine, Seoul, South Korea, 2011.
- [90] T. Yasuda, Y. Okamoto, N. Hamada et al., "Serum vitamin D levels are decreased and associated with thyroid volume in female patients with newly onset Graves' disease," *Endocrine*, vol. 42, no. 3, pp. 739–741, 2012.
- [91] J. Reglinski, W. E. Smith, R. Wilson, D. J. Halls, J. H. McKillop, and J. A. Thomson, "Selenium in Graves' disease," *Clinica Chimica Acta*, vol. 211, no. 3, pp. 189–190, 1992.
- [92] I. Bulow Pedersen, N. Knudsen, A. Carle et al., "Serum selenium is low in newly diagnosed Graves' disease: a population-based study," *Clinical Endocrinology*, vol. 79, no. 4, pp. 584–590, 2013.
- [93] L. Wang, B. Wang, S. R. Chen et al., "Effect of selenium supplementation on recurrent hyperthyroidism caused by Graves' disease: a prospective pilot study," *Hormone and Metabolic Research*, vol. 48, no. 9, pp. 559–564, 2016.
- [94] J. J. Khong, R. F. Goldstein, K. M. Sanders et al., "Serum selenium status in Graves' disease with and without orbitopathy: a case-control study," *Clinical Endocrinology*, vol. 80, no. 6, pp. 905–910, 2014.
- [95] L. H. Duntas, "The evolving role of selenium in the treatment of Graves' disease and ophthalmopathy," *Journal of Thyroid Research*, vol. 2012, Article ID 736161, 2012.
- [96] T. Wertenbruch, H. S. Willenberg, C. Sagert et al., "Serum selenium levels in patients with remission and relapse of Graves' disease," *Medicinal Chemistry*, vol. 3, no. 3, pp. 281–284, 2007.
- [97] J. Calissendorff, E. Mikulski, E. H. Larsen, and M. Moller, "A prospective investigation of Graves' disease and selenium: thyroid hormones, auto-antibodies and self-rated symptoms," *European Thyroid Journal*, vol. 4, no. 2, pp. 93–98, 2015.
- [98] A. P. Weetman, "Graves' disease," *The New England Journal of Medicine*, vol. 343, no. 17, pp. 1236–1248, 2000.
- [99] X. Liu, W. Qiang, L. Liu et al., "A second course of antithyroid drug therapy for recurrent Graves' disease: an experience in endocrine practice," *European Journal of Endocrinology*, vol. 172, no. 3, pp. 321–326, 2015.
- [100] D. Villagelin, J. H. Romaldini, R. B. Santos, A. B. Milkos, and L. S. Ward, "Outcomes in relapsed Graves' disease patients following radioiodine or prolonged low dose of methimazole treatment," *Thyroid*, vol. 25, no. 12, pp. 1282–1290, 2015.