Prevalence and ultrasound patterns of testicular adrenal rest tumors in adults with congenital adrenal hyperplasia

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Background: Testicular adrenal rest tumors (TARTs) are benign neoplasms affecting patients with congenital adrenal hyperplasia (CAH). The prevalence of TART in adult patients with CAH is not well known. Ultrasonography (US) is the main tool for diagnosing TART and the role of contrast-enhanced US (CEUS) is never investigated. The aim of this study was to evaluate the TART prevalence in adults with CAH, by stratifying patients according to disease phenotype and assessing the diagnostic performance of US, color Doppler (CD) US and CEUS.

Methods: Male patients >16 years old with certain diagnosis of CAH who underwent US for TARTs, between December 2015 and September 2019 were prospectively enrolled. The control group included patients without CAH affected by testicular lesions at US other than TARTs.

Results: TARTs were identified in 16 of 52 patients (31%), of whom 15 (93.8%) displayed the salt-wasting (SW) form ($P<0.001$). The prevalence of TARTs in patients with the SW form was 54%. One patient with the non-classic (NC) form (6%) showed TART (likely the first documented case). The mean age of patients upon detection of TARTs was significantly younger compared with the control group ($P<0.001$); moreover, TARTs were bilateral in 15/16 patients (93.8%; $P<0.001$) and the largest lesion was more frequent in the medium third of testis in the TART group (87.5%), statistically different from the control group ($P=0.013$).

Conclusions: TARTs almost exclusively affected patients with the SW form of CAH. Age at diagnosis and bilateralism are useful factors for achieving a correct diagnosis of TARTs in CAH adult patients.

Keywords: Testicular adrenal rest tumors (TARTs); congenital adrenal hyperplasia (CAH); ultrasonography (US)

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Introduction

Adrenal rest tumors represent a rare group of benign neoplasms that originate from cells with a biological profile similar to adrenal cells but are located in extra-adrenal sites (1-6). Among these tumors, testicular adrenal rest tumors (TARTs) are the most studied neoplasm due to their higher frequency.

TARTs were described for the first time by Wilkins in 1940 (7). Regardless of the different etiological theories (8-10), adrenocorticotrophic hormone (ACTH) has been recognized as the most important factor promoting the development of TARTs (1,11,12). Several diseases are characterized by an increase of ACTH; however, TARTs are discovered to exist almost exclusively in patients affected by congenital adrenal hyperplasia (CAH), excluding only rare cases (13-15).

CAH is a rare autosomal recessive disease that is generally (over 95% of all cases) caused by mutations in CYP21A2, the gene encoding the adrenal enzyme 21-hydroxylase (21OHD). CAH due to 21OHD deficiency is a continuum of disease severity that depends on the residual enzyme activity. The residual enzyme activity, which mainly depends on the genotype alterations (16), produces two different phenotypical categories defined as classic forms: (I) the salt-wasting (SW) form, which is the most severe type and is characterized by cortisol and aldosterone deficiencies due to absent residual enzyme activity; and (II) the simple virilizing (SV) form, in which the residual enzyme activity (1-2%) generally produces a sufficient aldosterone concentration but low cortisol and strongly elevated androgen concentrations. The incidence of classic CAH is estimated at 1:10,000 to 1:20,000 births worldwide (15,17). The non-classic (NC) form of CAH is characterized by a residual enzyme activity of 30-50%, only slightly elevated androgens, and generally sufficient cortisol and aldosterone concentrations. Therefore, it is considered the mildest form of CAH. The NC form is much more common, with an estimated prevalence of 1/1,000 births worldwide (18). There are other subtypes of CAH in which TARTs have been proven to occur, but these forms are extremely rare (19).

Different studies have demonstrated the possibility of adjusting hormonal replacement therapy for containing or even inducing the regression of TARTs in the early stages (20-23). Therefore, the correct diagnosis of TARTs is essential in the early stages of the disease. Furthermore, a correct diagnosis is also important for preventing misdiagnosis

and avoiding unnecessary testicular surgery (17,24-27).

Physical examination may detect a TART as a palpable mass when the lesion has a diameter of at least 2 cm, with a detection rate between 5% and 35% (28,29). Ultrasonography (US) demonstrated a detection rate of 89-94%, even for smaller lesions (30-32).

TARTs have a prevalence of 37% in patients with CAH. Unfortunately, currently, there are no robust data concerning the prevalence of TARTs in adult patients with CAH due to the lack of dedicated studies with large populations. A recent review reported the TART prevalence in adults affected by CAH to be 46% compared with 25% in studies including only children (1). However, this result is affected by several limitations: (I) not every enrolled study stratified the patients according to CAH phenotype; (II) the studies analyzed in the review were not focused exclusively on adults; and (III) the NC subtype form of CAH was often not included.

The aim of this study was to assess the prevalence of TARTs in adults with CAH who are already under hormone therapy by stratifying the population according to the three different CAH phenotypes (SW, SV, and NC forms). The secondary aim was to assess the diagnostic performance of US, color Doppler (CD) US, and contrast-enhanced US (CEUS) in the evaluation of testicular lesions.

We present the following article in accordance with the STROBE reporting checklist (available at <http://dx.doi.org/10.21037/tau-20-998>).

Methods

The local institutional review board (No. RIF.CE: 4882) approved this prospective study, and written informed consent was obtained from all patients. This study was conducted according to the Declaration of Helsinki (as revised in 2013) for clinical studies.

Male patients with a certain diagnosis of CAH who underwent US for TARTs at our institution between December 2015 and September 2019 were eligible for this prospective study.

In all patients, the diagnosis of CAH was made on symptoms and laboratory tests and was confirmed by mutation analysis (33).

The inclusion criteria of this prospective study were as follows: (I) patients >16 years old; and (II) patients under treatment with hormonal replacement therapy based on glucocorticoids. Serum levels of ACTH were measured in all patients. ACTH was used as the hallmark in the evaluation

of hormonal control, and the adequate control of CAH was defined as ACTH serum levels less than 50 pg/mL.

Image technique and analysis

All the enrolled patients were evaluated with the same US machine: Canon-Toshiba Aplio 500TM (Ōtawara, Kanto, Japan).

Both testicles were evaluated, and the following features were recorded on a dedicated database:

- (I) Testicle echogenicity, recorded as homogeneous or heterogeneous;
- (II) The entire didymus volume, calculated by applying the formula for an ellipsoid (length × width × thickness × 0.52);
- (III) Additional didymal and scrotal features such as varicocele or hydrocele;
- (IV) Presence or absence and number of lesions. In cases with multiple lesions, each was evaluated as a single lesion;
- (V) Lesion's location, recorded as in the upper, medium, or inferior third of the didymus;
- (VI) Testicular lesion size measured as the maximum diameter;
- (VII) Characteristics of testicular lesions: echogenicity, margins, vascularization on CD-US and CEUS.

The echogenicity of the lesion was determined as hypo-, iso-, or hyper-echoic or heterogeneous.

The margins of the lesions were recorded as oval or irregular.

In the presence of testicular lesions, CD-US and CEUS were performed.

Two different patterns were recognized on the CD-US: (I) high, uniform, and diffuse signals or peripheral rim of increased signals; and (II) no difference in signals with the contiguous normal testicular parenchyma. The same probe used for CD-US was used for CEUS in a low mechanical index (range, 0.04–0.1) after the injection of 4.8 mL of contrast agent (SonoVue®, Bracco, Milano, Italy), followed by 10 mL of 0.9% saline. Contrast enhancement was evaluated for at least 2 minutes after the SonoVue injection, and different phases of contrast kinetics were stored as video clips. Low mechanical index mode and the gray scale imaging picture were simultaneously recorded on a split screen. The CEUS results were interpreted by comparing the brightness of the lesion to the surrounding testicular tissue after the injection of the contrast agent, with three different patterns identified: (I) hypereenhancement, (II)

isoeenhancement, and (III) hypoeenhancement.

The study design included the use of a control group of patients without CAH who were affected by testicular lesions other than TARTs. This control group was retrospectively selected among patients with testicular lesions of similar dimensions who underwent US examination at our institution. The same demographic information, clinical characteristics, and US features of the control group were recorded as those for the CAH patient population.

Statistical analysis

Data were reported as counts and percentages for categorical variables and the median and inter-quartile range for continuous variables. Categorical variables were compared by the Chi-square or Fisher's exact tests, as appropriate. Continuous variables were compared with the Kruskal-Wallis test. Comparisons for demographics and general clinical characteristics were performed between groups of patients with CAH with and without TARTs and between patients with TARTs compared with consecutive patients affected by testicular lesions other than TARTs.

Univariate and multivariate logistic regressions were performed to assess the demographic, pathological, and testicular ultrasound characteristics associated with TART diagnosis. The results were reported as odds ratios (ORs) with 95% confidence intervals (95% CIs). An OR with the entire 95% CI less than 1 indicated that the covariate reduced the probability of a TART diagnosis. Conversely, when the OR and entire 95% CI were greater than 1, the covariate increased the abovementioned risk. An OR with a 95% CI equal to 1 indicated that the covariate did not significantly influence this probability. The results obtained for the clinical endpoints from the multivariate analysis, in the presence of two or more covariates influencing the risk, were translated in graphic form through the use of special nomograms for logistic regression. The probability values are two-sided; a probability value of less than 0.05 was considered to indicate statistical significance. All computations were performed using SPSS 13 (SPSS Inc., Chicago, IL, USA).

Results

Fifty-two patients with CAH satisfied the inclusion criteria and were enrolled in this study. The median [range] age of this patient population was 22.5 [16–48] years.

Table 1 Demographics and clinical characteristics of patients enrolled in the study

	Patients with CAH			Patients without CAH	
	TART (n=16), n (%) or median [range]	No TART (n=36), n (%) or median [range]	P	Control group (other testicular masses; n=16), n (%) or median (range)	P (comparison with TART group)
Age (years)	23.5 [18–43]	22 [16–48]	0.538	36.5 [13–75]	0.010
SW	15 (93.8)	13 (36.1)	<0.001	–	–
SV	0	10 (27.8)	0.019	–	–
NC	1 (6.2)	13 (36.1)	0.025	–	–
ACTH (pg/mL)	190 [5–2,258]	34.5 [5–2,160]	0.019		
Age at diagnosis	19 [17–40]	–	–	36.5 [13–75]	0.003
Final diagnosis	TART	No TART	–	8 leydigoma; 5 seminoma; 3 mixed germ cell tumors [§]	–

[§], Mixed germ cell tumors: seminomatous and non-seminomatous germ-cell tumors. CAH, congenital adrenal hyperplasia; TART, testicular adrenal rest tumor; SW, salt-wasting; SV, simple virilizing; NC, non-classic; ACTH, adreno-cortico-tropic hormone.

The control group consisted of 16 patients, including 8 (50%) affected by leydigoma, 5 (31%) affected by seminoma, and 3 (19%) affected by mixed germ-cell tumors (seminomatous and non-seminomatous germ-cell tumors). The median [range] age of the control group was 36.5 [13–75] years, which was statistically older than that of the CAH patient population ($P<0.001$). The demographic and clinical characteristics of the enrolled patients are detailed in *Table 1*.

Twenty-eight patients with CAH (54%) had the classic SW form of CAH, 10 patients (19%) had the SV form, and 14 patients (27%) had the NC form.

Focal lesions compatible with TARTs were identified in 16 of 52 patients (31%), of whom 15 (93.8%) patients had the SW form ($P<0.001$). The prevalence of TARTs was 54% in patients with the SW form. Only 1 patient with the NC form (6%) demonstrated TARTs. No TARTs were detected in patients with the SV form.

The median [range] age at the detection of TARTs was 19 [17–40] years, which was statistically younger than that of the control group {36.5 [13–75] years; $P=0.003$ }.

In the CAH patient population, the ACTH values were significantly higher (median 190 pg/mL) in the TART group than in the non-TART group (median 34.5 pg/mL; $P=0.019$).

The ultrasound features of patients with TARTs and of the control group with other testicular masses are detailed in *Table 2*.

There were no statistical differences between the two groups in terms of testicular appearance, volume, and

extradidymal findings (*Figure 1*).

Overall, there were 51 lesions compatible with TARTs in the CAH population: 8 (50%) patients had two TARTs and the remaining 50% had three or more lesions. No patients had only one nodule. There was a total of 17 lesions in the control group, of which only one patient presented with two lesions. This latter figure was significantly different compared with the TART group ($P<0.001$).

In the TART group, the lesions were bilateral in 15/16 patients (93.8%), differently from the control group ($P<0.001$). There were no differences in the incidence of lesions in the left didymus between the two groups. Conversely, the incidence of lesions in the right testis was significantly higher in the TART population than in the control group ($P=0.002$).

The largest lesion was significantly more frequently located in the medium third of the didymus in the TART group (87.5%) than in the control group ($P=0.013$).

In both groups, the most common US appearance of lesions was hypoechogenicity, which did not differ significantly between groups. No differences were found between the two populations in terms of lesion dimension and morphology or in the appearance on CD-US or CEUS.

The results of multivariate analyses (*Table 3*) for number of right testicular lesions and age at diagnosis are graphically reported in nomogram (*Figure 2*).

Discussion

The vast majority of TARTs are discovered in patients

Table 2 Ultrasound features of patients with TARTs compared to other testicular masses

	TART patients (n=16), n (%) or mean (SD)	Control group (other testicular masses) (n=16), n (%) or mean (SD)	P
Testicular US appearance			0.069
Homogeneous	13 (81.3)	16 (100.0)	
Heterogeneous	3 (18.7)	0	
Right testicle volume (cc)	10.7 (2.4)	10.6 (1.4)	0.466
Left testicle volume (cc)	11.5 (2.5)	11 (1.5)	0.347
Concomitant varicocele	4 (25.0)	7 (43.8)	0.264
Concomitant hydrocele	6 (37.5)	4 (25.0)	0.446
No. of testicular lesions per patient			<0.001
1	0	15 (93.8)	
2	8 (50.0)	1 (6.2)	
3	3 (18.8)	0	
4	1 (6.2)	0	
5	2 (12.5)	0	
6	2 (12.5)	0	
No. of right testicular lesions			0.002
1	10 (62.5)	5 (31.3)	
2	4 (25.0)	0	
3	1 (6.2)	0	
No. of left testicular lesions			0.070
1	10 (62.5)	12 (75.0)	
2	1 (6.2)	0	
3	3 (18.7)	0	
4	1 (6.3)	0	
5	1 (6.3)	0	
Bilateral lesions	15 (93.8)	1 (6.2)	<0.001
Location of largest lesion			0.013
Upper third	1 (6.25)	3 (18.8)	
Medium third	14 (87.5)	6 (37.5)	
Lower third	1 (6.25)	7 (43.7)	
Maximum lesion diameter (mm)	16.2 (16.9)	13.9 (1.1)	0.550
Testicular lesion echogenicity			0.827
Hypoechoic	10 (62.5)	11 (68.8)	
Hyperechoic	2 (12.5)	1 (6.2)	
Isoechoic	0	0	
Heterogeneous	4 (25.0)	4 (25.0)	

Table 2 (continued)

Table 2 (continued)

	TART patients (n=16), n (%) or mean (SD)	Control group (other testicular masses) (n=16), n (%) or mean (SD)	P
Testicular lesion morphology			0.723
Oval	8 (50.0)	9 (56.3)	
Irregular	8 (50.0)	7 (43.7)	
Intra-lesional signal on CD	12 (75.0)	11 (68.8)	0.694
CEUS appearance			1
Hyperenhancement	13 (81.2)	13 (81.2)	
Isoenhancement	3 (18.8)	3 (18.8)	
Hypoenhancement	–	–	

TART, testicular adrenal rest tumor; SD, standard deviation; US, ultrasound; CD, color Doppler; CEUS, contrast-enhanced US.

Figure 1 This figure demonstrates two different patients, one affected by CAH with TARTs (A,B,C) and one with a testicular non-seminomatous germ-cell tumor (D,E,F). The lesions showed the same imaging features, such as a hypoechoic appearance on B-mode (A,D), intralesional signal on CD (B,E), and hypervascularity on CEUS (C,F). The only differences between the two cases were the bilateralism of lesions in the patient with TARTs (A) and the different location: in the middle third for TARTs (A) and in the inferior third in the non-seminomatous germ-cell tumor (D). CAH, congenital adrenal hyperplasia; TART, testicular adrenal rest tumor; CD, color Doppler; CEUS, contrast-enhanced ultrasonography.

affected by CAH (13-15). Unfortunately, currently, there are no robust data concerning the prevalence of TARTs in adult patients with CAH due to the lack of dedicated studies with significant series populations.

The present study population was comprised of 52 adult patients with CAH. The largest series in the literature were those of Engels *et al.* (34) who enrolled 141 adult

men with CAH, and of Finkelstein *et al.* (12) who enrolled 244 patients. However, the study of Engels *et al.* (34) involved 14 centers in six different European countries whereas that of Finkelstein *et al.* (12) included both children [170] and adults [74] from 36 states in the United States of America and from Puerto Rico, Canada, Ecuador, Turkey, and the Dominican Republic. Therefore, our study

Table 3 Uni- and multi-variate analyses for evaluating factors associated with TART diagnosis during testicular lesion evaluation

	Univariate analysis		Multivariate analysis	
	OR (95% CI)	P	OR (95% CI)	P
Age at diagnosis (years)	0.872 (0.793–0.959)	0.005	0.850 (0.731–0.989)	0.035
Right testicular volume (cc)	1.017 (0.708–1.461)	0.926		
Left testicular volume (cc)	1.113 (0.786–1.575)	0.564		
Concomitant varicocele	0.429 (0.095–1.925)	0.269		
Concomitant hydrocele	1.8 (0.394–8.215)	0.448		
Number of right testicular lesions	25.104 (2.749–229.271)	0.004	82.608 (2.690–2537.06)	0.012
Number of left testicular lesions	1.017 (0.957–1.081)	0.589		
Testicular lesion ultrasound echogenicity				
Hypoechoic	Reference	–		
Hyperechoic	2.2 (0.172–28.137)	0.544		
Isoechoic	–	–		
Heterogeneous	1.1 (0.215–5.609)	0.909		
Testicular lesion irregular morphology	1.288 (0.320–5.169)	0.723		
Presence of intra-lesion CD signal	1.363 (0.289–6.415)	0.695		
Hyperenhancement on CEUS	1 (0.169–5.903)	1		
Location of maximum diameter testicular lesions	0.485 (0.142–1.657)	0.248		

TART, testicular adrenal rest tumor; OR, odds ratio; CI, confidence interval; CD, color Doppler; CEUS, contrast-enhanced ultrasonography.

population was representative because it was prospectively recruited from a single tertiary center and included only adult patients.

Our study population had a median age of 22.5 years, in line with those reported by Engels *et al.* (34) and Finkelstein *et al.* (12). In fact, the mean ages reported by those authors were 28 and 31 years, respectively, in adults. Moreover, in the present study, the median age at the diagnosis of TARTs was 19 years, which was significantly lower than that of the control group of patients without CAH who were affected by testicular masses other than TARTs (36.5 years; $P=0.003$). This result, which was difficult to determine in other studies (12,34), appears important for identifying at what age a patient with CAH should undergo US surveillance for TARTs.

In the present study, the TART prevalence was 31%, and the lesions were discovered in patients with the SW form of CAH in the vast majority of cases (94%; $P<0.001$). The prevalence of TARTs was 54% in patients with the SW form, which represented the most common form of CAH in our sample (54%). No TARTs were detected in patients

with the SV form. Only 1 patient with the NC form (6%) displayed TARTs. The overall prevalence of TARTs in our series (31%) was lower than that of other studies on adult populations, such as the series of Finkelstein *et al.* (12) who reported a prevalence of 44%. This difference was only apparent because Finkelstein *et al.* (12) calculated this percentage as the number of TARTs only in “Classic men” (SW + SV). Therefore, if we calculate the percentage in the same manner, our prevalence was 39.5% in patients with only classic forms of CAH, similar to the abovementioned series. In another study by Dumic *et al.* (35), the overall TART prevalence was 29%, similar to our series. However, the study population by Dumic *et al.* (35), was composed of both children and adult patients. In fact, the prevalence of TART exclusively in adults was 44%, but the percentage of SW forms was not reported in adults.

The absence of TARTs in the SV form of CAH in our series is in line with the vast majority of published series (34,36–38). Furthermore, in our series, only one patient with the NC form presented with TARTs. Although the NC form of CAH has a high prevalence in the general

Figure 2 Nomogram. Nomogram reporting a probability score for TART diagnosis expressed by the addition of single score for the number of right testicular lesions and the age at diagnosis. The number of right testicular lesions was associated with a score of 1–4, while the age at diagnosis varied from 0–4.5. TART, testicular adrenal rest tumor.

population, with a rate of 1 per 1,000 births, TARTs are reported to be exceptionally rare in these individuals. In fact, to the best of our knowledge, our case represents the sixth case of TARTs associated with NC in published series (1). In particular, Falhammar *et al.* (18), Engels *et al.* (34), and Kocova *et al.* (39) described the other five incidences of TARTs in patients with the NC form of CAH. However, the phenotype of these patients was not detailed, and based on mutation status, these patients might also belong to the more affected classic form of CAH. Conversely, our case was proven to be an NC form of CAH (this may be the first case of a patient proven to have the NC form of CAH affected by TARTs). It is likely that data concerning the prevalence of TARTs in the NC form of CAH could be underestimated due to the lack of specific studies on large populations. Moreover, the NC form of CAH often remains unknown because it is characterized by only a few symptoms or is completely asymptomatic, and there is no actual screening test for this form. Conversely, the high frequency of TARTs

in the SW form compared to the other forms of CAH may suggest the prominent role of the disease severity. In fact, compared to the other parameters such as age at detection, modality of diagnosis, and hormonal control, several studies have indicated that the SW phenotype is the main risk factor for developing TARTs (1,12,17,27,28).

In our CAH patient population, ACTH values were significantly higher in the TART group than in the non-TART group ($P=0.019$). This result supports the current perception that TARTs occur more frequently in patients affected by more severe forms of CAH because these patients are exposed to higher levels of ACTH, which is a possible causative factor for TART development (29,38,40,41). The first etiological theory of TARTs assumes that these lesions derived from ectopic adrenal cells and migrated to the gonads in fetal life, and their growth is under control of ACTH and angiotensin II (42,43). The second etiological theory sustains that TARTs may originate from pluripotent fetal Leydig cells that have ACTH and

luteinizing hormone receptors (10). In both etiological theories of TART, ACTH plays an important role, justifying our results.

There were 51 overall lesions compatible with TARTs in the CAH population in our series: 8 (50%) patients had two TARTs and the remaining 50% had three or more lesions. No patients had only one nodule. There were 17 testicular lesions in the control group of no-CAH patients with testicular lesions other than TART, and only one patient presented two lesions, differently from the TART group ($P<0.001$). The multiplicity of lesions per testis (more than one for didymus) has been described in the original series by Stikkelbroeck *et al.* (31), Wang *et al.* (44), and Chihaoui *et al.* (45), but with frequencies of 19%, 16%, and 25%, respectively, which were all lower than ours. According to the actual state of knowledge about TART histopathology, TARTs are thought to arise as small multiple nodules that tend progressively to coalesce into a single nodule. Thus, different numbers of nodules per testis are considered a reflection of a different pathological stage (31). The high frequency of the multiplicity of nodules per testis likely indicates that our cohort of patients was composed mainly of subjects with an early-intermediate stage of CAH disease.

In the present series, in the TART group, the lesions were bilateral in the vast majority of patients (93.8%), unlike that observed in patients without CAH who were affected by testicular tumors other than TART ($P<0.001$). There were no differences in the incidence of lesions in the left didymus between the two groups. Conversely, the incidence of TARTs in the right testis was significantly higher in the TART population than in the control group ($P=0.002$). Bilateralism is traditionally considered one of the best clues in the diagnosis of TARTs. In fact, different published series reported bilateral nodules compatible with TART in all of their cases (10,30,46). Nevertheless, different series have reported that bilateralism was not present in all cases (17,20,28,34). In particular, the lowest percentage of bilateralism (31%) was detected by Claahsen-van der Grinten *et al.* (20). It is interesting to note that all the nodules detected by Claahsen-van der Grinten *et al.* (20) were less than 10 mm in size. In our case series, the mean maximum diameter was greater than 10 mm. Our results sustain the hypothesis that TARTs may always have a bilateral process, but in patients with early stages of TARTs, indicated by the small dimension of the lesions, synchronous contralateral lesions may be too small to be detected.

The largest lesions were significantly more frequently located in the medium third of the didymus, near the rete

testes, in the TART group (87.5%) compared with the control group ($P=0.013$). This is a hallmark of TARTs that is reported constantly in the literature (1,29,30). A single case of our series displayed involvement of the epididymis by a nodule compatible with TARTs. Although no pathological proof of its nature was achieved, the epididymis has been described as an unusual location for TARTs (2).

In the vast majority of our cases, the TARTs appeared hypoechoic on US, with intra-lesional signals on the CD-US and hypervascularization on CEUS. However, these features were not specific for TARTs compared with the control group. Furthermore, no statistical differences were found in terms of didymal and extra-didymal features in patients with CAH and TARTs in respect to the subjects without CAH who were affected by testicular lesions other than TARTs. These data do not differ significantly from other original studies (28,31,44).

The role of CD-US in TART characterization is controversial due to a great discordance present in the literature. Several authors (28,31,34) failed to identify an increased vascular signal on CD-US in their series, unlike other experiences (30,44,45).

Differently from CD-US, there are no published series exploring the role of CEUS in TARTs. Therefore, our results are the first that reported imaging findings of CEUS in patients affected by TARTs.

Parenchymal echogenicity of testes has been reported in a few studies, such as those by Bouvattier *et al.* (17) and Cabrera *et al.* (47), with results very similar to ours.

One unicity of the present study is the comparison of CAH patients with TARTs with non-CAH subjects with testicular lesions other than TARTs. Finally, none of the US, CD-US, and CEUS features of the lesions and of the didymus or extradidymus showed statistically significant differences between TART and non-TART testicular lesions. Therefore, in our opinion, the imaging figures are helpful for detecting TARTs in high-risk patients such as adults affected by CAH, rather than performing a differential diagnosis.

Another novelty of this study is the introduction of the nomogram for the risk estimation of TARTs by the radiologist based on the number of right testicular lesions and age at diagnosis. The nomogram is a two-dimensional diagram that allows the approximate graphical calculation of a function, such as a multivariate analysis. These graphical instruments are easy to use and enable rapid calculation of the pathological risks for a single patient in the era of tailored medicine.

This study is affected by several limitations. First, a histological diagnosis of suspected TART lesions in CAH patients was not achieved. However, the biopsy is not indicated in the clinical suspect of TARTs. The main limitation is the difficulty in recruiting a large study population, due the rarity of the disease. However, this work could represent a good model for a single tertiary center study of the prevalence of TARTs in CAH. Furthermore, due to the limited sample size of our study, the multivariate analysis is affected by an overfitting variables bias. However, the results of the present multivariate analysis should be interpreted with caution but are still valuable for addressing further larger studies in order to validate our results.

In conclusion, TARTs affected approximately one-third of the adult patients with CAH, principally those affected by the SW form. The age at diagnosis and the right testicular lesions appear very useful figures to achieve a correct diagnosis of TARTs in CAH patients. No specific US figures were demonstrated to be helpful in differential diagnosis with other testicular lesions; therefore, US is mandatory only for lesion detection. If our results are confirmed by other studies, it will be possible to plan US surveillance programs in patients with the SW form of CAH starting from adulthood.

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Footnote

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Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at <http://dx.doi.org/10.21037/tau-20-998>). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related

to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The local institutional review board (No. RIF.CE: 4882) approved this prospective study, and written informed consent was obtained from all patients. This study was conducted according to the Declaration of Helsinki (as revised in 2013) for clinical studies.

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