New Insights on the Mechanisms Affecting Fertility in Men with Non-Seminoma Testicular Cancer before Cancer Therapy

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Purpose: Patients with non-seminoma testicular cancer (NSTC) cancer can be subfertile or infertile, and present reduced sperm quality, but the underlying mechanisms are unknown. The aim of this study was to compare the sperm proteome of patients with NSTC, who cryopreserved their sperm before starting cancer treatment, with that from healthy fertile men.

Materials and Methods: Semen volume, sperm motility and sperm concentration were evaluated before the cryopreservation of samples from patients with NSTC (n=15) and the control group (n=15). Sperm proteomic analysis was performed by liquid chromatography-tandem mass spectrometry and the differentially expressed proteins (DEPs) between the two groups were identified using bioinformatic tools.

Results: A total of 189 DEPs was identified in the dataset, from which five DEPs related to sperm function and fertilization were selected for validation by Western blot. We were able to validate the underexpression of the mitochondrial complex subunits NADH:Ubiquinone Oxidoreductase Core Subunit S1 (NDUFS1) and ubiquinol-cytochrome C reductase core protein 2 (UQCRC2), as well as the underexpression of the testis-specific sodium/potassium-transporting ATPase subunit alpha-4 (ATP1A4) in the NSTC group.

Conclusions: Our results indicate that sperm mitochondrial dysfunction may explain the observed decrease in sperm concentration, total sperm count and total motile count in NSTC patients. The identified DEPs may serve as potential biomarkers for the pathophysiology of subfertility/infertility in patients with NSTC. Our study also associates the reduced fertilizing ability of NSTC patients with the dysregulation of important sperm molecular mechanisms.

Keywords: Cancer therapy; Male fertility; Non-seminoma; Sperm quality; Testicular cancer

INTRODUCTION

Testicular cancer (TC) is a rare tumor among men, representing only 1% to 2% of all tumors [1]. The USA National Cancer Institute reported 8,850 new cases in 2017 [2] and this number is estimated to increase to
9,310 new cases in 2018 [3]. The etiology of TC is not fully understood but several risk factors were identified, including genetic background, age, race/ethnicity, undescended testicle, human immunodeficiency virus infection and/or AIDS [4,5]. One of the main types of TC is non-seminoma testicular cancer (NSTC). It represents about half of the cases of germ cell tumors [6], which account up to 95% of all TC cases [7]. NSTC is commonly diagnosed in young men (25–35 years) [8,9], thus constituting a major threat for couples during reproductive years. Although the etiology of NSTC is largely unknown, its treatment is highly effective, showing a 5-year survival rate of 99% in early stages, and 48% to 91% in more advanced and metastatic stages [6]. Surgery, radiotherapy or chemotherapy are frequently used to treat this type of cancer [10]. However, these therapies are associated with secondary effects on male fertility, especially due to gonadal damage [11]. After the treatment, many patients show reduced sperm quality and impaired spermatogenesis, and often result in permanent infertility [12,13]. A 30% decrease in the probability to conceive by natural conception after cancer therapy has been reported [14]. Thus, in most of NSTC cases the clinicians advise the patients to undergo sperm banking before initiating any kind of cancer therapy [15,16]. NSTC tends to grow rapidly and spread to other tissues [17], highlighting the urgency to initiate the treatment.

Most of the cases of NSTC are found when couples are not able to have children and seek for medical assistance. In fact, the development of NSTC is facilitated by the presence of abnormal semen parameters [18] and subfertility/infertility [19-21]. There are many cases of men presenting poor semen quality and reduced fertilizing ability prior to NSTC diagnosis [12,22,23]. Part of the problem starts with the impairment of spermatogenesis, as the tumor affects the normal hormonal regulation [24]. It is of extreme importance for the field of male reproduction to investigate the mechanisms by which NSTC affects male fertility. In this context, proteomics has emerged as a valuable tool to explore alterations in sperm proteome triggered by certain health conditions [25-27]. Due to their post-translational modifications, spermatozoa are highly susceptible to proteome alterations during maturation throughout the male reproductive tract [28]. These changes are highly dependent on the individual health status and may be a useful source of biomarkers to identify certain disease conditions in its early stage. In this study, we aimed to analyze the sperm proteome of patients with NSTC who had cryopreserved semen samples before initiating cancer therapy and identify potential sperm protein biomarkers responsible for the altered reproductive function in these patients.

MATERIALS AND METHODS

1. Study population and ethics statement

This study was ethically approved by the Institution Review Board (IRB) of Cleveland Clinic. The control group included 15 healthy volunteers with proven fertility, i.e., men who fathered a child in the last two years before entering the study, whereas the NSTC group comprised 15 NSTC patients who cryopreserved semen samples before starting cancer therapy. All the participants signed an informed written consent at the time of sample collection at the Andrology Center, Cleveland Clinic.

2. Semen analysis and cryopreservation

Semen samples were collected after 2 to 3 days of abstinence and placed at 37°C for 20 to 30 minutes to allow liquefaction. Volume, sperm motility, and sperm concentration were evaluated according to World Health Organization (WHO) 2010 guidelines [29]. Total sperm count and total motile count were also calculated and results were expressed as mean±standard error of mean (SEM). Semen samples were then cryopreserved in TEST-yolk buffer (TYB; Irvine Scientific, Santa Ana, CA, USA) in a ratio 1:1 as previously described [30] and finally stored in liquid nitrogen at -196°C.

3. Sperm protein extraction and quantification

Cryopreserved samples were thawed and centrifuged at 4,000×g for 10 minutes to isolate spermatozoa. The pellet was washed four times with consecutive resuspensions in phosphate buffered saline (PBS; Irvine Scientific) and centrifugations at 4,000×g for 10 minutes, at 4°C. Radio-immunoprecipitation assay (RIPA; Sigma-Aldrich, St. Louis, MO, USA) buffer supplemented with Protease Inhibitor Cocktail, cOmplete™ ULTRA Tablets, ethylene diamine tetraacetic acid-free (Roche, Mannheim, Germany) was added to each sperm pellet (100 µL RIPA/10⁶ sperm) and left overnight at 4°C for cell lysis. Then, samples were centrifuged at 10,000×g for 30 minutes, at 4°C and the supernatant containing
the protein fraction was transferred to a new centrifuge tube. Protein quantification was performed using the Pierce BCA Protein Assay kit (Thermo Fisher Scientific, Waltham, MA, USA) according to the manufacturer’s instructions.

4. Liquid chromatography-tandem mass spectrometry

The present study was conducted in compliance with the Minimum Information about a Proteomics Experiment (MIAPE) guidelines of the Human Proteome Organization’s Proteomics Standards Initiative (HUPO-PSI) for reporting proteomics studies [31]. Three samples from control or NSTC groups (n=3/group) were pooled using the same amount of protein from each sample and analyzed by liquid chromatography-tandem mass spectrometry (LC-MS/MS). The samples in each pool were randomly selected within the experimental group and each pool was evaluated as an individual sample. Proteomic analysis was performed in a Finnigan LTQ-Obitrap Elite hybrid mass spectrometer using standard conditions and software [32,33]. The identification of the differentially expressed proteins (DEPs) between the control and NSTC groups was conducted via Scaffold (ver. 4.0.6.1; Proteome Software Inc., Portland, OR, USA) software. The abundance of each protein in the pool was classified as very low, low, medium or high based on the number of spectral counts. The normalized spectral abundance factor (NSAF) ratio was calculated to categorize the expression profile of DEPs as underexpressed, overexpressed or unique to one of the groups, as previously described [32].

5. Ingenuity Pathway Analysis

Bioinformatic analysis of the identified DEPs was performed through ingenuity pathway analysis (IPA; Qiagen, Hilden, Germany) software. This program allowed the evaluation of the top canonical pathways, diseases and bio functions, “Tox lists” and upstream regulators related to the DEPs. The criteria for the selection of DEPs for validation by Western blot (WB) included: 1) proteins involved in reproductive system development and function; 2) proteins involved in the top canonical pathways; 3) proteins with a higher difference of abundance between the experimental groups; 4) proteins with a well-described function in the literature. Proteins unique to the control group were not considered for protein validation.

6. Validation of selected differentially expressed proteins

Validation of key DEPs was performed by WB using six individual samples per group. Sperm proteins (25 µg/sample) were mixed with 4× Laemmli sample buffer in a ratio 1:3 in a total volume of 25 µL in PBS. The prepared samples were immediately boiled at 95°C for 10 minutes for protein denaturation and run into a 4% to 15% sodium dodecyl sulfate-polyacrylamide gel electrophoresis gel at constant voltage (90 V) for 2 hours. A molecular weight marker (Precision Plus Protein™ Dual Xtra Standards) was loaded in the first well of each gel. Transfer of proteins from the gels to methanol-activated polyvinylidene difluoride membranes was performed with constant voltage (18 V) for 30 minutes. Membranes were blocked for 90 minutes at room temperature, with a 5% non-fat milk solution prepared in tris-buffered saline tween-20. Membranes were incubated overnight (4°C) with specific primary antibodies followed by the respective secondary antibodies at room temperature, for 90 minutes (Supplemental Table 1). Images were acquired in a ChemiDoc™ MP Imaging System after 5 minutes-incubation with enhanced chemiluminescence reagent (GE Healthcare, Marlborough, MA, USA) protected from the light. Image Lab™ Software allowed quantification of bands density and the density of each band was divided by the corresponding total protein lane density. Results were expressed as fold change relative to the control group. All the reagents and equipment used for WB were purchased from BioRad (Hercules, CA, USA) unless specifically stated.

7. Statistical analysis

Semen parameters and WB results were analyzed by Mann-Whitney test for independent samples, using the MedCalc Software (ver. 17.8; MedCalc, Ostend, Belgium). The results were considered statistically significant for p<0.05.

<table>
<thead>
<tr>
<th>Table 1. Semen parameters of fertile men (control) and patients with non-seminoma testicular cancer (NSTC)</th>
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<tr>
<td>Parameter</td>
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<tr>
<td>Volume (mL)</td>
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<td>Sperm motility (%)</td>
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<tr>
<td>Sperm concentration (10⁶/mL)</td>
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<tr>
<td>Total sperm count (10⁶)</td>
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<td>Total motile count (10⁶)</td>
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<td>Values are presented as mean±standard error of mean (n=15 per group). Results were considered statistically significant for p&lt;0.05.</td>
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Belgium). All data are presented as mean±SEM and differences with p<0.05 were considered statistically significant.

Results

1. Semen parameters

After assessing the semen parameters of fertile men (control) and patients with NSTC (Table 1), we observed that all the samples were within the normal standards of WHO [29]. The volume of ejaculate and sperm motility were similar between the groups, however there was a reduction in total motile count in NSTC group to about half of that in the control group (p=0.0136). Besides, sperm concentration and total sperm count were significantly lower (p=0.0015, and p=0.0084, respectively) in NSTC patients in comparison to control group (Table 1).

2. Identification of the differently expressed proteins

A total of 1,149 proteins were identified in the control group and 1,123 proteins in the NSTC group. Comparative proteomic analysis resulted in the quantification of 1,201 proteins, from which 189 were found to be DEPs between the experimental groups (Fig. 1). Most of the DEPs were underexpressed (n=105; 55.6%), while 37.0% (n=70) were overexpressed in NSTC. Besides, 3.2% (n=6) were unique to NSTC group and 4.2% (n=8) unique to control group (Fig. 1).

3. Selection of key differentially expressed protein for validation

Through the IPA analysis, we observed that within the top ‘canonical pathways’ and ‘Tox lists’, many DEPs were involved in ‘mitochondrial dysfunction’. Among those, we selected the proteins NADH-ubiquinone oxidoreductase 75 kDa subunit (NDUFS1) and cytochrome b-c1 complex subunit 2 (UQCRC2), subunits of the mitochondrial complexes I and III, respectively. Besides, one of the top molecular and cellular functions in which the DEPs in this dataset were involved was ‘free radical scavenging’. NDUFS1 together with annexin A2 (ANXA2) were included in this category, playing a role in the synthesis and metabolism of reactive oxygen species (ROS).

We also selected two proteins related to ‘physiological system development and function’: acrosin precursor (ACR) and sodium/potassium-transporting ATPase subunit alpha-4 (ATP1A4). The function of these proteins is important for specific reproductive processes (Table 2) intricate to fertilization. The abundance and expression pattern obtained by the proteomic analysis regarding the five selected proteins is presented in Table 3.

4. Prediction of the upstream regulators

The IPA software can predict the upstream regulators of the DEPs in a certain dataset. The activation/inhibition of certain regulatory proteins may explain the

<table>
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<tr>
<th>Process</th>
<th>Protein</th>
<th>p-value</th>
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<tr>
<td>Cell movement of sperm</td>
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<td>Binding of sperm</td>
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<td>Function of reproductive system</td>
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<td>Morphology of germ cells</td>
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<td>Fertility</td>
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<tr>
<td>Penetration of zona pelludica</td>
<td>ACR</td>
<td>0.0314</td>
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ATP1A4: sodium/potassium-transporting ATPase subunit alpha-4, ACR: acrosin precursor.

Fig. 1. Number of proteins identified by proteomic analysis of spermatozoa samples obtained from fertile men (control) and men with non-seminoma testicular cancer (NSTC), and expression profile of the differentially expressed proteins (DEPs) identified after comparative analysis between the experimental groups. UE: underexpressed, OE: overexpressed.

Table 2. Specific functions of the differentially expressed proteins related to reproductive system development and function identified by the bioinformatic analysis when comparing the sperm proteome of patients with non-seminoma testicular cancer relative to fertile men.
altered protein expression in the spermatozoa of NSTC group and provide information about the underlying mechanisms. The underexpression of the mitochondrial subunits NDUFS1 and UQCRC2 was predicted to be due to the activation of the rapamycin-insensitive companion of mammalian target of rapamycin (RICTOR; z-score 2.4). Moreover, the inhibition of the metalloendopeptidase OMA1 (OMA1; z-score -2.0) was also related to the underexpression of UQCRC2.

5. Western blot validation

We identified the expression of all the selected DEPs through WB. We observed a significant decrease in the protein expression of NDUFS1 from 1.00±0.13 in the control group to 0.13±0.04-fold change to control in the NSTC group (Fig. 2). A similar decrease was detected in the expression of UQCRC2 in NSTC group (0.13±0.06-fold change to control) relative to control (1.00±0.14). WB results also showed a reduced expression of ATP1A4 and ANXA2 in patients with NSTC (0.25±0.05 and 0.57±0.11-fold change to control, respectively) when compared to the control group (1.00±0.14 and 1.00±0.17, respectively). On the other hand, there was a significant increase in the expression levels of ACR from 1.00±0.22 in the control group to 5.95±1.06-fold change to control in the NSTC group (Fig. 2).

**DISCUSSION**

Despite the significant advances in the field of male reproduction, the etiology of male subfertility/infertility is still poorly understood. About 15% to 30% of the infertility cases are diagnosed as idiopathic [34]. Patients with NSTC show similar reproductive disorders as those observed in infertile men [35]. It is important to investigate the molecular alterations at the spermatozoon level in patients with NSTC before initiating cancer therapy to understand the NSTC-related infer-
tility and allow a premature non-invasive diagnosis. This is the first study focusing on the identification of sperm protein alterations in patients with NSTC before any treatment compared with healthy fertile men.

In spermatozoa, mitochondria are crucial for cellular energy production, survival and function [36]. Additionally, the mitochondrial respiratory chain is the main source of ROS. ROS play a key role in fertilization as they regulate sperm capacitation, hyperactivation, and acrosome reaction [37]. The mitochondrial complexes I and III are the principal sites of ROS generation [38]. They are constituted by several subunits, which are required for the functional assembly of the complexes [39]. For instance, NDUFS1 is the largest subunit of complex I, which by NADH oxidation provides electrons for the respiratory chain [40]. Besides, UQCRC2 is an essential component of complex III, another electron pump that contributes for oxidative phosphorylation (OXPHOS) [41]. Our proteomic data showed the underexpression of NDUFS1 and UQCRC2 in the NSTC group (Table 3), and these results were confirmed by the WB analysis (Fig. 2). The underexpression of NDUFS1 in the spermatozoa of NSTC men may lead to complex I dysfunction, which is one of the most common mitochondrial dysfunctions observed in humans [39]. Additionally, underexpression of UQCRC2 was associated with poor fertilization rates due to insufficient adenosine triphosphate (ATP) production and reduced sperm kinematics parameters [42]. Downregulation of NDUFS1 and UQCRC2 suggest mitochondrial dysfunction and subsequently abnormal ROS production that may result in oxidative stress. This has a major impact in sperm function, especially on the ability of spermatozoa to undergo capacitation and bind to the oocyte [43]. Oxidative stress is a common condition found in patients with NSTC and patients with other infertility-related disorders such as varicocele [44]. In fact, an underexpression of NDUFS1 and UQCRC2 was also observed in men with varicocele [44]. Furthermore, infertile varicocele patients with higher ROS production showed decreased sperm concentration and motility [45]. Thus, reduced total motile count, concentration and total sperm count observed in NSTC group (Table 1) may be explained by reduced ATP production and uncontrolled ROS production induced by mitochondrial dysfunction. Mitochondria are also susceptible to ROS attack, thus releasing pro-apoptotic molecules that will increase sperm cell death [46].

The underexpression of NDUFS1 and UQCRC2 was predicted to be due to activation of RICTOR, a component of the mammalian target of rapamycin (mTOR) pathway. This upstream regulator deserves special attention as the mTOR signaling cascade is responsible for the regulation of important cellular events, including cell growth and metabolism [47]. More specifically, RICTOR is a subunit of mTOR complex 2 (mTORC2), which is involved in cytoskeleton organization and insulin signaling [48,49]. The mechanisms by which mTOR regulates male fertility are not fully understood, but both complexes (mTORC1 and mTORC2) have been implicated in the regulation of blood-testis-barrier dynamics during spermatogenesis [50]. Though, a higher expression of RICTOR was observed in the early stages of the seminiferous epithelial cycle, while a progressive decline was observed at late stages [51]. Thus, activation of RICTOR in mature spermatozoa of NSTC patients may lead to a dysregulation of the mTOR system, affecting sperm quality. Inhibition or absence of mTOR has been reported to result in lower sperm production and decreased motility [50].

Our results indicate reduced sperm concentration, total sperm count and total motile count (Table 1) in NSTC group and these findings may be explained by an altered mTOR system in these patients. In fact, mTOR deregulation is common to many types of cancer [52] and should be explored as a mechanism of male fertility impairment in patients with NSTC. Furthermore, another upstream regulator, OMA1 was also associated with the underexpression of UQCRC2. OMA1 is a proteolytic enzyme located in the inner membrane of the mitochondria that is involved in the regulation of mitochondrial dynamics [53]. OMA1 deficiency severely affects the mitochondrial fusion-fission equilibrium, having important implications in cellular metabolic homeostasis [54]. Thus, the inhibition of OMA1 in spermatozoa of NSTC patients may result in mitochondrial dysfunction, further evidenced by the underexpression of not only UQCRC2, but also NDUFS1.

Among the identified DEPS, ATP1A4 was selected due to its participation in many reproductive processes, including sperm motility, function and fertilization, as well as function of reproductive system (Table 2). ATP1A4 is involved in the exchange of sodium and potassium ions across the plasma membrane, which are essential ions for sperm maturation [55]. Proteomic analysis demonstrated an underexpression of ATP1A4
in NSTC group and this result was validated by the WB technique (Fig. 2). As ATP1A4 function is dependent on the availability of ATP [56], the downregulation of this protein is in accordance with the observed decrease in the expression of mitochondrial subunits NDUFS1 and UQCRC2, which consequently result in reduced ATP production. Previous studies reported that ATP1A4 plays a key role in human sperm motility [57,58] and fertilizing ability in mice [55]. The underexpression of this protein may also be related to the reduction in total motile count seen in NSTC patients compared to fertile men (Table 1).

ANXA2 belongs to a family of calcium-dependent proteins that bind to negatively charged phospholipids in the cellular membrane [59]. This protein was identified in prostasomes, which are membrane-bound vesicles produced by the prostate epithelial cells [60] that are also present in the acrosome and tail of spermatozoa [61]. Prostasomes are rich in calcium and an increase in sperm intracellular calcium concentration was seen after fusion with prostasomes [62]. Calcium is very important for sperm motility acquisition and hyperactivation [63]. ANXA2 is believed to play a role in calcium transfer from prostasomes to spermatozoa, and is therefore important for sperm function [60].

Our proteomic analysis showed an overexpression of ANXA2 in NSTC group relative to control (Table 3). On the contrary, our WB results revealed a significant decrease in the expression levels of ANXA2 (Fig. 2) in patients with NSTC. We believe that our WB results are more reliable for the following reasons: i) use of a higher sample size for WB evaluation; ii) the intensity of the bands in each group (Fig. 2); and iii) the significant decrease in total motile count found in NSTC group (Table 1). In fact, a decreased mRNA expression of ANXA2 was associated with reduced motility [64]. In our study, the underexpression of ANXA2 in patients with NSTC suggests an insufficient calcium transfer from prostasomes, thereby contributing to the decrease in sperm quality.

The protein ACR was selected for validation due to its association with several reproductive processes, including binding of sperm, fertilization, morphology of germ cells, fertility, and penetration of zona pellucida (Table 2). The precursor form (inactive) of this protein - proacrosin, was identified by the proteomic analysis as underexpressed in patients with NSTC. However, quantification of WB data showed a high overexpression of this protein in NSTC group (Fig. 2). Once again, we have contradicting results between the two techniques (proteomics vs. WB), but using a larger sample size for WB, we could observe by the representative bands (Fig. 2) that ACR is highly expressed in NSTC group. During acrosome reaction, proacrosin is converted to acrosin (active form) that plays a key role in sperm binding to the oocyte [65]. The overexpression of proacrosin in the spermatozoa of patients with NSTC may lead to an abnormally high production of acrosin, resulting in premature or defective acrosome reaction and impaired sperm-oocyte binding. Though, ANXA2 and ACR results need further confirmation with a larger sample size or an additional technique.

CONCLUSIONS

Altogether, our results indicate that patients with NSTC present an altered expression of proteins that are essential for sperm function and fertilization, especially those involved in sperm mitochondrial function (NDUFS1 and UQCRC2), motility (ATP1A4 and ANXA2), and fertilization (ATP1A2 and ACR). We validated the results obtained through the proteomic analysis by WB, showing the underexpression of NDUFS1, UQCRC2 and ATP1A4 in the spermatozoa of patients with NSTC relative to fertile men. Once these findings are reproduced by others in the field, we speculate that these three proteins may serve as potential biomarkers for the early and non-invasive clinical diagnosis of NSTC, as well as for new treatment approaches. In future, the analysis of the expression levels of these three proteins could be included into the routine evaluation of an infertile men as a screening for TC. Our data also highlight the importance of mTOR signaling pathway in the regulation of male fertility and that its possible dysregulation in spermatozoa of NSTC patients is related with reduced sperm quality and fertility potential. Although we found discordant results regarding the expression levels of ANXA2 and ACR, the alterations in these two proteins need further investigation in patients with NSTC as they may contribute to the understanding of the molecular basis of impaired fertility in these patients.

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Disclosure

Authors have no potential conflicts of interest to disclose.

Author Contribution

Agarwal A and Sharma R designed the study and helped in data interpretation. Dias TR processed the samples, conducted the experiments and wrote the first draft of the manuscript. Pushparaj PN performed the bioinformatic analysis and Ahmad G helped in the collection/processing of samples. All authors read and approved the final manuscript.

Supplementary Materials

Supplementary materials can be found via https://doi.org/10.5534/wjmh.180099.

REFERENCES


