

Diagnosing Causes of Myopathy and Its Effective Treatment

Dr. Harshika Gupta

PT, Bachelor's of Physiotherapy (BPT), Master of Physiotherapy (MPT) in Neurology*, NIMS University, Jaipur Rajasthan (India) Email: harshika8874gupta@gmail.com

Abstract

This article is based on myopathy and its effective treatment and management process. This study focuses on myopathy which is one of the neuromuscular disorders that can affect the internal and external organs of the body. Accordingly, individuals can face difficulties in their daily life and household activities. On the other hand, this study has shed light on the diagnosis process of myopathy and it has been seen that CD4, CD8, and MHC classification make an impact on the disease, and these kinds of factors force to improve the treatment process also put the positive effects on the diagnosis process of this disease. This method also analyses the entire diagnosis reports and finds the symptoms of myopathy and this process of diagnosis also helps the individuals to get the proper treatment. Moreover, a biomedical genetic test is another process that can also recognize enzyme abnormalities and other complications. This article also highlights the tools that are generally used in the treatment process of myopathy. In order to check the level of enzymes, such as Creatine kinases (CK), Alanine Aminotransferase (ALT), Aspartate aminotransferase (AST) level helps to predict the damage to muscles tissues and biopsy is one of the processes of detection. myositis-specific antibody technique also checks the antibodies and their action. On the other hand, this study also discussed the treatment and management strategies of myopathy.

Keywords

MHC or Major Histocompatibility Complex, Electromyography, Creatinine Kinases (CK), Alanine Aminotransferase (ALT), Aspartate Aminotransferase (AST), STIR, Anti-Signal Recognition Particle (SRP)

INTRODUCTION

Myopathy is a neuromuscular disorder that causes muscle weakness due to the dysfunction of muscle fibre and includes muscle cramps, stiffness, and spasms. This disease indicates effectiveness on the muscles that are connected to skeletal muscle and creates a problem with daily chores appropriately. There are different types of myopathies that can be inherited or acquired due to the circumstances of the individual lifestyle. "Inherited myopathy" can be developed due to the inheritance of abnormal gene mutation from the parents that causes the development of congenital myopathy in the person's body [1]. The defect in the mitochondria leads to mitochondrial myopathy that affects organ systems such as the heart, brain, and gastrointestinal tract. "Metabolic myopathy" indicates that dysfunction in the genes creates a lack of code for the enzyme needed to maintain the normal function of the muscles and their movement. This type of Myopathy reflex exercise intolerance extensional muscle pain in the shoulder thighs and muscle fibre condition and leads to muscle weakness in the individuals and lack of utilization of normal muscle strength.

On the other hand, "Muscular dystrophy" includes progressive degeneration of muscle tissues that indicates insufficient structural support proteins present in the body. Limb-girdle muscular dystrophies are common in India and 0.8% per 100,000 people due to the lack of Lamin, myoferlin, and telethon protein in the individuals [2]. Infectious myopathy causes an attack on the body itself that reflects muscle dysfunction and toxic myopathy is the result of toxins from alcohol, toluene, and prescribed drugs. This disease causes people to get into trouble doing activities such as getting out of a chair, climbing stairs, and doing other activities in the household [3]. Myopathy causes people to feel shortness of breath in the exertion of completing their daily tasks efficiently. This disease can be treated with physiotherapy for developing muscle activity and recommended medication for minimizing the weakness of muscle tissues.

Muscle symptoms related to infection caused by bacteria, viruses, or other infectious organs can be improved with the treatment of antibiotics in the treatment process. The supportive and symptomatic treatment assists in developing the treatment process by providing support to the weakened muscle, including drug therapy, physiotherapy, and surgeries for better treatment of myopathy [4]. The objective of this study is to identify the causes of Myopathy in the diagnosis process and the effectiveness of its treatment procedure on the patient's health condition. This factor helps to know more about the specific symptoms of the individuals during myopathy and analyse the influence of medication and physiotherapy treatment on the individuals' health condition.



LITERATURE REVIEW

Diagnosis process of myopathy

Myopathy had been diagnosed with the help of diagnostic tests for analysing the impact of nerves and muscles and the heterogeneity of this disease for understanding clinical features to predict therapeutic outcomes. The inflammatory cells have muscle-bearing surface markers such as CD4, CD8, and major histocompatibility complex (MHC) classes for classification of the disease impact [5]. This factor assists to develop clinical thinking in proceeding with the diagnosis process of the patients. The health care professionals include genetic tests for identifying any issues related to the structure of genes with the help of collecting blood and saliva samples of the patient. This clinical report helps medical professionals identify the causes of myopathy symptoms happening in the individual's body and develop a better treatment procedure. The utilisation of biochemical genetic test guides to analyse the abnormality of enzymes from the samples collected of blood, urine, and amniotic fluid of the patient in the diagnosis procedure. This factor helps to examine the abnormal level of hormones that can cause stiffness and weakness of the muscle in the patient's body.

It has been seen that the older men had asymmetric, distal weakness in their muscle tissues and shown in the biopsy report for characterises the red-rimmed vascular of "Gomori trichrome stain". This factor helps to diagnose the impact of myopathy on the vascular system that indicates high serum CK levels for identifying the weaknesses of muscle tissues [6]. The autoimmune process guides the identification of the pathology of immune-mediated necrotizing myopathy in the diagnosis process and reflects the impact of lipid-lowering agents, myotonia, and periodic parallelism among the patients. Moreover, the diagnosis process assists the idiopathic inflammatory myopathy signs as the combination genetic and environmental factors that causes of inflammation in the muscles of patients. Myopathy diagnosis shows neuromuscular junction disorders such as Eaton-Lambert syndrome, and myasthenia gravis and includes inflammatory dystrophies as the result of a deficiency of protein in the muscles.

On the other hand, inflammatory muscle disease shows an increment in the serum muscle enzyme level that is identified in the characterised electromyography (EMG) findings and muscle biopsy demonstrating inflammation causes. This factor shows the self-reactive T-lymphocytes that have reacted to myocytes increase the expression of the MHC class and indicate the cause of myocyte death. The CD8+ T Cells are responsible for the myocyte's invasion of the polymyositis under the electroscope diagnosis process for compressing the muscle fibres [8]. Cytotoxic factor mechanism in polymyositis assists to identify the disease pathogen and formation of immunological synapses between MHC muscle fibre expression and CD8+ T Cells with the help of costimulatory support. The activation of complement C3 helps to identify the disease in the early stage and changes the vasculature and neurosis of the caterpillar, perivascular inflammation, and ischemic muscle fibre damage. This factor assists to reduce the number of caterpillars per myocytes in the patient's body. The mechanism of creating misfolded MHC glycoproteins in the automatism that leads to Endoplasmic reticulum stress and inflammatory response to the obstipation cell in defining the nature of muscle enzymes.

Tools utilized in the diagnosis process

The diagnosis process for identifying muscle weakness in the patients includes the analysis of patient history, physical examination routine, laboratory test, autoimmune serologies, imaging study, and neurologic evaluation EMG and muscle biopsy. The history and physical examination help to identify the causes of the myopathy happening in the patient's body and its impact on the patient's muscle tissue. This factor determines the onset of the disease, the pattern of presentation, and the impact of environmental factors on developing the treatment process of myopathy patients. A detailed neurology exam plays an important role in observing a neuropathic disease for bordering the differential of the neuropathic process such as metabolic process and dystrophy [9]. The determination of muscle enzyme levels such as "Creatinine kinases (CK)", "Aspartate aminotransferase (AST)" and "Alanine Aminotransferase (ALT)" for estimated damage of muscle tissues. This factor helps to examine the biopsy report before proceeding with the treatment procedure of myopathy among the patients.

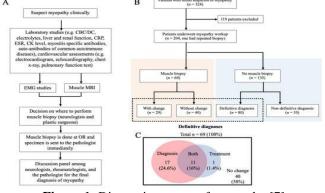


Figure 1: Diagnosis process of myopathy [7]

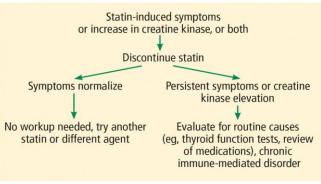


Figure 2: Stain-induced symptoms in myopathy[10]



The myositis-specific antibody technique is utilized for the determination of the presence or absence of antibodies with the immunosorbent assay methods for analysing the sensitivity and specificity of autoantibodies. On the other hand, the anti-signal recognition particle (SRP) assists to predict the future course and the progress of the treatment for schedule with the help of anti-tRNA synthesis . This factor acts as the part of antibodies for strong presentation in reducing the muscle enzyme impact from the patient's body and defines MCA status for the ant synthetase syndrome. This ant synthetase help to provide better treatment therapy to the muscle strengthening their muscles tissues and examining its impact on the recovery process. The imaging process utilises magnetic resonance imaging (MRI) for determining the location of muscle involvement, muscle loss, and severity of the disease activities. MRI process helps to understand the short tau inversion repeat (STIR) in observing the changes of water content and inflammation for increasing signal intensity that determines the autonomy of the disease.

The application of MRI assists to evaluate the extent of muscle damage and determine the appropriate site of muscle biopsy by looking into specific sides of interest in real-time and measuring the signs of active disease. The neuromuscular evaluation heads to differentiate between myopathy and neuropathy diseases with the help of Electromyography (EMG) that assists to measure the electrical activity of the patient's muscles [11]. This factor helps to understand nerve conduction by velocity testing that assists to distinguish the details of muscle disease from neurological disease in supporting the findings of myopathy diseases. Moreover, muscle biopsy assists medical professionals in using certain areas of tissues for examination of lymphocytic infiltration, fibre necrosis, degenerative and regenerative behaviour of the muscle fibre. The nerve conduction studies help to identify the places of electrodes in the skin for the stimulating process of nerves and recording the electrical activity that monitors the quality of muscle weakness [12]. This test assists to perform the lumbar puncture in extracting the cerebrospinal fluid with needles between two vertebrae in the lower back for analysis of the pattern of myopathy diseases.

Treatment and management of myopathy

The treatment procedure for myopathies includes different approaches such as strength therapy, supportive care, symptomatic therapy, and psychological support to the patients in maintaining the management procedures. The multimodal approach of conservative and supportive care helps these patients to improve their quality of life with the help of pharmacological treatment in reducing the issues of myopathy effectively. The strength therapies help to apply immunotherapy in dermatomyositis (DMY), polymyositis (PM), and immune-mediated necrotizing myopathy in developing muscle activities. The application of "Corticosteroids" assists to develop pulmonary and cardiac function with the help of high-dose medication [13]. This factor guides the stimulation of the muscle fibbers' activities in individuals through enzyme replacement therapy that manages the enzyme stimulation in the muscle tissues. The development of the simulation process helps to decrease the inflammation of the muscle tissues and improve the body's autoimmunity response with the help of drugs such as "methotrexate", "cyclosporine", "tacrolimus", and "azathioprine".

The implementation of pharmacological intervention assists healthcare providers in utilising creatine monohydrate as the counter supplement in improving the strength and function of dystrophies. This factor helps to enhance dermatomyositis and polymyositis and develop the tolerance of creatinine for fulfilling the needs of supplements. Exercise also helps to increase the flexibility of the muscle tissues and build the strength of muscles for doing daily activities of the individuals. The application of aerobic exercise in inflammatory myopathy leads to regulating PM and DMY for developing the maximum oxygen uptake, and isometric force with the help of randomised control trial process [14]. The improvement of peak torque guides to include resistive exercise in chronic myositis for developing the result in the inclusion of body myositis in strengthening the muscle groups. The studies found that low-intensity activity exercise helps to maximize the heart rate from 60% to 65% with the benefits of facioscapulohumeral dystrophy (FSHD) through Becker muscular dystrophy (BMD). This factor helps to strengthen the muscles with the help of an effective training process and provides supplements according to the requirement of the patient's diet.

The designing process of restrictive exercises Ashish to build the muscles with the help of repetitive contraction against the resistance force such as dumbbell, free weight, and resistive band. This factor helps myopathy patients to extend the distribution of weakness and develop baseline activity for enhancing the treatment procedure of patients. The guidance of a physiotherapist to understand the appropriate strengthening test for performing the strengthening evaluation in the maintenance of the daily log of activity and exercise performed by the patient. The documentation of physical tasks guides myopathy patients to determine the benefits of exercise programs in their health condition and evaluate its impact on strengthening muscle fibres effectively [15]. Moreover, the supported therapy helps to promote limiting limb contractures, respiratory compromise, swallowing, and gastrointestinal issues in the management process of myopathy treatment. The development of contraction with the help of strengthening and positioning assists to preserve the function of muscle tissues in maintaining the quality of health condition.

The stretching program guides the patient and their families to improve the joint range of motion in DMD with serial casting through the guidance of therapists. The symptomatic therapy treatment guides to fight against factors related to myopathy disease issues and resolves it with cardiac management, and treatment of Edema and pain effectively [16]. Psychological support helps to look after the



mental condition of the patients by providing medication and treatment for fighting against anxiety and depression to develop the quality of the patient's life.

METHODOLOGY

Research method

The research method assists to develop the constitution process of research questions for developing insights about the research topic and builds the research patterns for collecting the data properly. The explanatory method is utilised in this research process for explaining the diagnosis cause and the treatment procedure of myopathy and determining the response of the patient in the treatment process. This factor helps to provide insightful knowledge about the causes that increase fatigue in the muscle tissue and weaken the muscle fibre and result in paralysis in most cases [17]. The knowledge about the cause of myopathy helps the therapist and caregiver to enhance the planning process of the treatment procedure.

Research design

The well-constructed research design helps to ensure the usage of methods helps to fulfil the research aim with the help of the collection of high-quality data. This factor analyses the gathered data related to myopathy in the right way to answer the research question. The descriptive design helps the research process to develop the credibility of resources in gathering data related to the cause and the treatment procedure of myotherapy [18]. The utilization of proper resources assists to describe the myopathy impact on the individual's patients with the help of well-being in the health care services and motivates them to participate in the exercise and strength programs.

Research approach

The research approach guides provide a detailed plan for keeping track of the research process and progress with the help of appropriate data in managing the research process effectively. The deductive approach is used in this study that helps to analyse the possible relationship between concepts and variables related to myopathy causes and its treatment procedure. This factor assists to collect data from the online resources related to myopathy causes and treatment process that guides to the evaluation of the treatment programs efficiently [19]. These data access to answer the questions related to myopathy therapy and its causes that decreases the quality of life of the patients due to increment paralysis causes.

Data collection

The data collection process helps to collect data from reliable and valid resources such as PubMed, Google Scholar, and last year's articles and journals related to the research topic. The secondary data is collected to develop the research method and observe the research patterns with the help of collected data in the working process. This factor helps other researchers and readers to know more about the myopathy cause and treatment program's effect on the patient's health condition with the help of therapists and caregivers. The gathered data also helps to describe the myopathy sign and symptoms of the patient and explain the diagnosis process with the help of medical tools in the treatment process appropriately [20]. The knowledge about the diagnosis process helps to construct the research method to develop the treatment procedure of the individual patients by analysing their clinical and past history data for constructing proper treatment plans with the guidance of therapists.

Data analysis

The data analysis process helps to analyse and monitor the collected data related to the research topic in delivering better outcomes in the research work. The secondary data analysis process is utilised in this research process for developing an understanding of the relationship between the diagnosis and treatment process of myopathy. This factor helps to modify the insights about the cause of myopathy and its impact on the patient's health condition that enhance the therapist's approach to the patients. The development of the therapist approach helps to improve the construction of strength care, symptomatic care, and psychological support for participation in the treatment procedure. This factor assists to motivate the patients to take the treatment for improving their muscle strength with the help of the exercise treatment and take their prescribed medicine on time [21]. The maintenance of the treatment procedure helps the patient to notice a growth in their health condition and develop flexibility in the muscle tissues and build new muscle fibres effectively.

FINDINGS AND DISCUSSION

Analysing the diagnosis process of the myopathy

Myopathy diagnosis process assists the health care provider in observing the physical and clinical report of the patients in examining reflex activity, muscle strength balance, and sensation. This factor helps to understand the current condition of the patients and build a treatment plan according to their circumstances for reducing muscle weakness efficiently [22]. The blood test help to identify the muscle enzyme such as creatinine kinase that causes a breakdown in the muscle tissues and increases the impact of myopathy in the patient's body. The low level of electrolytes such as sodium, magnesium potassium, calcium, and phosphorus in the muscle tissues of patients indicates the cause of myocyte death in maintaining the straightness of the muscle fibre. On the other hand, the EMG report access to monitor the narc conduction with the electricity for understanding the type and degree of muscle damage [23]. The muscle biopsy helps to define the gene mutation causes in the patient's body for identifying the inflammatory issues in the muscle tissues.

Effectiveness of tools in myopathy diagnosis

Gene testing assists to analyse the DNA base mutation with the help of monitoring DNA samples for understanding the



variables of genetic nutrition that develops various types of myopathies in individuals. This factor indicates that the GNE gene in the patient's body is caused by weakness of the tibialis anterior muscle and decreases the stability of muscle fibres. Computerized tomography helps to develop the imaging process of muscle disease muscle declaration and muscle wasting and measure the amount of muscle damage in the diagnosis process. Magnetic resonance imaging (MRI) assists to identify the edema and fatty replacement of muscle tissues for the diagnosis process of inherited myopathy with the help of distinctive patterns in the muscle involvement [24]. This factor helps to reveal the diffuse hyperintense T2-weighted signal for including muscle atrophy and fatty replacement with the anti-SRP antibodies. Electromyography helps to analyse the individual's motor unit action potential (MUAP) duration from the close position muscle membrane and determine the muscle functions and fatigues.

Influence of treatment on myopathy patients

The treatment procedure of myopathy includes strength care, symptomatic care, supportive care, and psychological care for developing the quality of healthcare services and enhancing the patient's quality of life. The strength care process helps to enrol the patient in the standing program for obtaining the strength measurement in analysing the followup evaluation schedule in understanding the success and failure rate of the constructed programs. This program helps to develop the muscle's activity with the help of a caveat in reducing the forced vital capacity in dyspnoea for increasing the strength of muscle tissues [25]. The supportive care assists to develop the offering of supportive exercise with the help of Cervical orthoses for supporting the damaged muscle tissues in walking and straightening of limbs effectively. Symptomatic therapy assists therapists to reduce the secondary impact of obesity, pain, and edema due to the lack of muscle activity and develop the quality of treatment management of the patients. Psychological care helps to reduce the depression and anxiety impact on the patient with the encouragement to participate in the therapy process for developing their muscle strength with the help of exercise and medication efficiently.

Discussion

The above literature shows that myopathy refers to the disease of the muscles due to the lack of proper nutrients and minerals that causes damage to the muscle tissues. The common symptom is muscle weakness that indicates weakened muscles of the upper arm and upper legs and also creates issues in the respiratory muscles. Muscle atrophy begins the development of thinning out muscles and wasting away of muscles that create abnormal shapes of the bones. On the other hand, it also highlights a lack of energy, fatigue of muscle, and weakness in progressive muscle damage with exertion in the individual's patients [26]. The diagnosis process utilizers other blood such as erythrocyte sedimentation rate for measuring inflammation and antinuclear antibody test for monitoring autoimmune activity.

The application of electromyography helps to understand the muscle's consulate through the usage of needles for detecting different characteristics of muscle structure and movement.

The offering of exercise programs helps to develop the structural process of muscle tissues and increase the flexibility of muscles in the treatment procedure properly [27]. The treatment procedure also offers psychological treatment for motivating and encouraging the patient to participate in various activities related to muscle strengthening and developing their mental health effectively.

CONCLUSION

In this above study, it has been described that myopathy is one kind of neuromuscular disorder that is responsible for muscle weakness. There are some classifications of myopathy that can be acquired or inherited due to the specific circumstances of the individual's lifestyle. From this study, it has been seen that Myopathy has the ability to affect other organs of the body and also make an impact on the health condition and psychological condition of a person. This disease affects the brain, heart, gastrointestinal tract, and other essential internal parts of the body. On the other hand, myopathy disease creates barriers to leading a healthy lifestyle and it also reduces the activity level of a person who faces difficulties in climbing stairs, walking, holding something by hand, and other household activities. This study has also discussed the diagnosis process of myopathy and also discusses the tools that are used in the treatment of myopathy. Accordingly, this study has shed light on the management and treatment process of myopathy.

From this study, it has been recognized that this disease has various factors that force it to change and develop clinical thinking and it makes an impact on the process of treatment and also helps the individual to get better treatment. This research study has utilised the secondary qualitative method for getting authentic data and also uses the data for the study. This process helps to get lots of information on myopathy and its treatment process. These processes help to provide value to the research paper. The finding of the study reflects that the observation of blood report and EMG report guides the therapist to provide a proper treatment plan to the myopathy patients and analyse the muscle functionality. The diagnosis process of myopathy utilizes muscle biopsy for analysing the gene mutation and determining the type and degree of muscle damage efficiently.

The offering of supportive and psychological care helps the patient to participate in exercise therapy. This factor helps to use supplement tools for developing the flexibility of muscle tissues in the hand and legs in the treatment process. Psychological care helps the patient to fight against the present anxiety in the treatment procedure to accept the failures in the myopathy treatment program.



REFERENCES

- [1] Bugiardini, E., Khan, A.M., Phadke, R., Lynch, D.S., Cortese, A., Feng, L., Gang, Q., Pittman, A.M., Morrow, J.M., Turner, C. and Carr, A.S., 2019. Genetic and phenotypic characterisation of inherited myopathies in a tertiary neuromuscular centre. *Neuromuscular Disorders*, 29(10), pp.747-757.
- [2] Rare disease India, 2021. *Muscular Dystrophy*. Available at:http://www.rarediseasesindia.org/musculardystrophy [Accessed on:25th December 2022]
- [3] Cashy, C., De Leon, A., Anderson, J. and Boes, T., 2020. Fusobacterium: A Rare Case of Septicemia in a Patient with Multiple Abscesses. *Ohio Chapter/Air Force Chapters*, p.26.
- [4] Korb, M., Peck, A., Alfano, L.N., Berger, K.I., James, M.K., Ghoshal, N., Healzer, E., Henchcliffe, C., Khan, S., Mammen, P. and Patel, S., 2022. Development of a standard of care for patients with valosin-containing protein associated multisystem proteinopathy. *Orphanet journal of rare diseases*, 17(1), pp.1-15.
- [5] Ahmed, S.T., Craven, L., Russell, O.M., Turnbull, D.M. and Vincent, A.E., 2018. Diagnosis and treatment of mitochondrial myopathies. *Neurotherapeutics*, 15(4), pp.943-953.
- [6] Bugiardini, E., Morrow, J.M., Shah, S., Wood, C.L., Lynch, D.S., Pitmann, A.M., Reilly, M.M., Houlden, H., Matthews, E., Parton, M. and Hanna, M.G., 2018. The diagnostic value of MRI pattern recognition in distal myopathies. *Frontiers in neurology*, 9, p.456.
- [7] Paoletti, M., Pichiecchio, A., Cotti Piccinelli, S., Tasca, G., Berardinelli, A.L., Padovani, A. and Filosto, M., 2019. Advances in quantitative imaging of genetic and acquired myopathies: clinical applications and perspectives. *Frontiers in neurology*, 10, p.78.
- [8] Olivier, P.A., De Paepe, B., Aronica, E., Berfelo, F., Colman, R., Amato, A., Dimitri, D., Gallardo, E., Gherardi, R., Goebel, H.H. and Hilton-Jones, D., 2019. Idiopathic inflammatory myopathy: Interrater variability in muscle biopsy reading. *Neurology*, 93(9), pp.e889-e894.
- [9] Tsamis, K.I., Boutsoras, C., Kaltsonoudis, E., Pelechas, E., Nikas, I.P., Simos, Y.V., Voulgari, P.V. and Sarmas, I., 2022. Clinical features and diagnostic tools in idiopathic inflammatory myopathies. *Critical Reviews in Clinical Laboratory Sciences*, 59(4), pp.219-240.
- [10] Tran, A., Walsh, C.J., Batt, J., Dos Santos, C.C. and Hu, P., 2020. A machine learning-based clinical tool for diagnosing myopathy using multi-cohort microarray expression profiles. *Journal of translational medicine*, *18*(1), pp.1-9.
- [11] Lehmann Urban, D., Mohamed, M., Ludolph, A.C., Kassubek, J. and Rosenbohm, A., 2021. The value of qualitative muscle MRI in the diagnostic procedures of myopathies: a biopsy-controlled study in 191 patients. *Therapeutic Advances in Neurological Disorders*, 14, p.1756286420985256.

- [12] Dos Santos, A.M., Misse, R.G., Borges, I.B.P., Perandini, L.A.B. and Shinjo, S.K., 2021. Physical exercise for the management of systemic autoimmune myopathies: recent findings, and future perspectives. *Current opinion in rheumatology*, 33(6), pp.563-569.
- [13] Hervier, B. and Uzunhan, Y., 2020. Inflammatory myopathy-related interstitial lung disease: from pathophysiology to treatment. *Frontiers in Medicine*, *6*, p.326.
- [14] Cheung, K., Rathbone, A., Melanson, M., Trier, J., Ritsma, B.R. and Allen, M.D., 2021. Pathophysiology and management of critical illness polyneuropathy and myopathy. *Journal of Applied Physiology*, 130(5), pp.1479-1489.
- [15] Kruse, R.L., Albayda, J., Vozniak, S.O., Lawrence, C.E., Goel, R., Lokhandwala, P.M., Ness, P.M., Tobian, A.A., Bloch, E.M. and Crowe, E.P., 2022. Therapeutic plasma exchange for the treatment of refractory necrotizing autoimmune myopathy. *Journal of Clinical Apheresis*, 37(3), pp.253-262.
- [16] Nadin, T., Haque, A., Akil, M. and Hughes, M., 2019. Management of the idiopathic inflammatory myopathies. *Prescriber*, 30(5), pp.28-33.
- [17] Vill, K., Müller-Felber, W., Landfarth, T., Köppl, C., Herzig, N., Knerr, C., Holla, H., Steidle, G., Harms, E., Hohenfellner, K. and Interdisciplinary Cystinosis Group, 2022. Neuromuscular conditions and the impact of cystine-depleting therapy in infantile nephropathic cystinosis: A cross-sectional analysis of 55 patients. Journal of Inherited Metabolic Disease, 45(2), pp.183-191.
- [18] Veziari, Y., Kumar, S. and Leach, M.J., 2022. An exploration of barriers and enablers to the conduct and application of research among complementary and alternative medicine stakeholders in Australia and New Zealand: A qualitative descriptive study. *PloS one*, *17*(2), p.e0264221.
- [19] Rashid, Y., Rashid, A., Warraich, M.A., Sabir, S.S. and Waseem, A., 2019. Case study method: A step-by-step guide for business researchers. *International journal of qualitative methods*, 18, p.1609406919862424.
- [20] Hazarika, A., Dutta, L., Barthakur, M. and Bhuyan, M., 2018. A multiview discriminant feature fusion-based nonlinear process assessment and diagnosis: application to medical diagnosis. *IEEE Transactions on Instrumentation and Measurement*, 68(7), pp.2498-2506.
- [21] Wang, Q., Li, Y., Ji, S., Feng, F. and Bu, B., 2018. Immunopathological characterization of muscle biopsy samples from immune-mediated necrotizing myopathy patients. *Medical Science Monitor: International Medical Journal of Experimental and Clinical Research*, 24, p.2189.
- [22] Sainio, M.T., Aaltio, J., Hyttinen, V., Kortelainen, M., Ojanen, S., Paetau, A., Tienari, P., Ylikallio, E., Auranen, M. and Tyynismaa, H., 2022. Effectiveness of clinical exome sequencing in adult patients with difficult-to-diagnose neurological disorders. *Acta Neurologica Scandinavica*, 145(1), pp.63-72.



- [23] Rodriguez, B., Larsson, L. and Z'Graggen, W.J., 2022. Critical Illness Myopathy: Diagnostic Approach and Resulting Therapeutic Implications. *Current treatment options in neurology*, pp.1-10.
- [24] Angelini, C., Marozzo, R., Pegoraro, V. and Sacconi, S., 2020. Diagnostic challenges in metabolic myopathies. *Expert Review of Neurotherapeutics*, 20(12), pp.1287-1298.
- [25] Packer, M., 2019. Disease–treatment interactions in the management of patients with obesity and diabetes who have atrial fibrillation: the potential mediating influence of epicardial adipose tissue. *Cardiovascular diabetology*, *18*(1), pp.1-6.
- [26] Danielak, D., Karaźniewicz-Łada, M. and Główka, F., 2018. Assessment of the risk of rhabdomyolysis and myopathy during concomitant treatment with ticagrelor and statins. *Drugs*, 78(11), pp.1105-1112.
- [27] Rietveld, A., Lim, J., de Visser, M., van Engelen, B., Pruijn, G., Benveniste, O., van der Kooi, A. and Saris, C., 2019. Autoantibody testing in idiopathic inflammatory myopathies. *Practical neurology*, 19(4), pp.284-294.