

Bioimpedance Analysis and Type 2 Diabetes: An Overview

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Annotation. Two anatomical structures are involved in the neurovascular conflict (NVK) in classical HTN: the trigeminal nerve root (CTN) and the artery (more often the superior cerebellar artery (BMA), less often the inferior anterior cerebellar artery (NPMA) and the basilar artery (BA). The occurrence of NVK occurs with a critical convergence of these structures and an increase in the traumatic effect of the artery on the CTN. The force of the artery hitting the CTN is subject to the physical laws of hydrodynamics. In turn, endothelial dysfunction contributes to atherosclerotic vascular modification with increased rigidity of the vascular wall. The existing methods of treating HTN reflect the evolution of ideas about its etiology and pathogenesis. Among the conservative methods of treatment of classical HTN, first-line drugs are anticonvulsants and, above all, carbamazepine, which suppresses cortical and stem foci of sensitization. In parallel with the increase in tolerance to the drug, the number of adverse events also increases.

Keywords: Bioimpedance Analysis (BIA), Body Composition, Type 2 Diabetes (T2D), Insulin Resistance, Visceral Adiposity/Fat, Skeletal Muscle Mass, Body Fat Percentage, Fat-Free Mass (FFM), Fluid Status, Metabolic Health, Risk Assessment, Diabetes Management

Introduction

Type 2 diabetes mellitus (T2DM) is one of the most prevalent chronic metabolic diseases globally, marked by insulin resistance and impaired glucose metabolism. Unlike type 1 diabetes, which involves an autoimmune destruction of insulin-producing beta cells, T2DM typically arises due to a combination of genetic, lifestyle, and environmental factors, leading to progressive insulin resistance. Obesity, particularly the accumulation of visceral fat, plays a significant role in its development and progression.

As T2DM progresses, it can result in various complications, such as cardiovascular disease, neuropathy, nephropathy, and retinopathy. These complications are primarily linked to poor glycemic control and chronic inflammation associated with obesity. Effective management of T2DM goes beyond mere blood glucose monitoring; it requires a comprehensive approach that includes monitoring body composition and fat distribution. This is where bioimpedance analysis (BIA) comes into play.

BIA is a non-invasive, accessible, and cost-effective method that assesses body composition, which includes measuring fat mass, lean body mass, and body water content. By examining these parameters, healthcare providers gain insights into a patient's metabolic health, potential risks, and progress in managing diabetes. Given the importance of muscle mass for glucose metabolism, fat distribution's effect on insulin sensitivity, and hydration status's impact on kidney function, BIA offers a valuable tool in the holistic management of T2DM.

In recent years, BIA has gained popularity as an essential method in clinical settings, where understanding and controlling body composition are increasingly recognized as critical aspects of treating diabetes. This analysis can provide ongoing information that complements traditional markers like HbA1c and fasting glucose levels, allowing healthcare professionals to create individualized care plans based on a patient's unique body composition. This not only aids in effective disease management but also supports preventive

measures that can help slow disease progression and mitigate complications associated with T2DM.

The Importance of Body Composition in T2DM

Understanding body composition is central to managing type 2 diabetes mellitus (T2DM) because it provides insight into factors that directly influence disease progression and management. Unlike simple weight or BMI measurements, body composition analysis reveals the proportions of fat, lean mass (muscle), and water within the body. These components are closely related to insulin sensitivity, metabolic health, and overall diabetes management, making them critical factors in assessing and treating T2DM effectively.

1. Visceral Fat and Insulin Resistance

One of the most significant insights from body composition analysis is the amount and distribution of fat, especially visceral fat. Visceral fat, stored deep within the abdominal cavity around organs, is metabolically active and linked to insulin resistance—a hallmark of T2DM. This type of fat releases free fatty acids, pro-inflammatory cytokines, and hormones that impair insulin signaling, thereby worsening glycemic control. Unlike subcutaneous fat, which is stored under the skin, visceral fat has a more direct and detrimental impact on glucose metabolism.

Studies indicate that individuals with higher visceral fat levels are more likely to experience rapid T2DM progression and an increased risk of cardiovascular complications. Therefore, monitoring visceral fat in T2DM patients allows for better-targeted interventions that focus on reducing this harmful fat to improve insulin sensitivity and metabolic health.

2. Lean Body Mass and Glucose Uptake

Lean body mass, primarily muscle tissue, plays a vital role in glucose metabolism. Muscle tissue is one of the primary sites for insulin-mediated glucose uptake, which means that individuals with a higher proportion of lean mass generally have better glucose control. In contrast, low muscle mass is often associated with decreased insulin sensitivity and an increased risk of hyperglycemia.

In T2DM management, maintaining or increasing lean body mass is important for enhancing insulin sensitivity and reducing the severity of blood sugar fluctuations. Bioimpedance analysis (BIA) allows clinicians to monitor changes in lean body mass, helping them recommend resistance training or other muscle-building exercises that can support glucose uptake and improve glycemic control.

3. Body Water and Hydration Status

Hydration is an often-overlooked factor in diabetes management, yet it has substantial effects on kidney function, blood volume, and glucose regulation. Dehydration can contribute to hyperglycemia by concentrating glucose in the bloodstream, complicating blood sugar management. Furthermore, T2DM can disrupt hydration status due to osmotic diuresis, particularly when blood sugar levels are elevated.

BIA can differentiate between intracellular and extracellular water, offering an assessment of a patient's hydration status. This information helps healthcare providers make informed recommendations to maintain optimal hydration, which is essential for metabolic stability, especially in patients with impaired kidney function. Proper hydration can improve renal health, reduce the risk of acute kidney injury, and help stabilize blood glucose levels.

4. Individualized Treatment and Monitoring

Given the heterogeneous nature of T2DM, where patients exhibit a wide range of body compositions, an individualized approach to treatment is essential. Body composition analysis allows for a tailored treatment plan that addresses each patient's unique profile of muscle, fat, and water distribution. For instance, T2DM patients with high visceral fat may benefit from a different dietary and exercise regimen than those with low lean body mass.

BIA measurements provide continuous, trackable data, enabling healthcare providers to monitor a patient's progress over time. Regular assessments of body composition can help clinicians adjust treatment plans and interventions based on real-time changes in body composition, such as improvements in lean mass or reductions in visceral fat. This personalized approach enhances adherence and improves long-term health

outcomes in individuals with T2DM.

5. Reducing the Risk of Diabetes-Related Complications

Monitoring and improving body composition can help reduce the risk of diabetes-related complications. Higher levels of lean mass and lower levels of visceral fat are associated with lower rates of cardiovascular disease, neuropathy, and other diabetes complications. By maintaining a favorable body composition, patients with T2DM can minimize these risks and achieve better quality of life.

In summary, body composition analysis offers a comprehensive view of the factors affecting T2DM and provides actionable data for managing the disease. BIA serves as a critical tool in understanding and addressing the metabolic challenges of T2DM, supporting a more individualized and effective approach to diabetes care.

How Bioimpedance Analysis Works

Bioimpedance analysis (BIA) is a non-invasive, safe, and accessible method for assessing body composition. BIA works by passing a small electrical current through the body and measuring the resistance (impedance) as it encounters different types of tissues. Since body tissues like fat, muscle, and water conduct electricity differently, BIA can estimate the body's composition by interpreting how the current moves through each component. This method has become increasingly popular in clinical settings, especially for managing conditions like type 2 diabetes mellitus (T2DM), where precise monitoring of body composition is essential.

1. Basic Principle of BIA

The principle behind BIA relies on the fact that body tissues differ in their electrical properties:

- **Lean tissue (such as muscle)** is high in water content, making it a good conductor of electricity. Electrical currents pass more easily through muscle tissue because of its high water and electrolyte content.
- **Fat tissue**, on the other hand, contains less water and does not conduct electricity as efficiently. This causes higher resistance (impedance) to the electrical current.

The BIA device calculates body composition based on these variations in resistance. Using algorithms, the device estimates fat mass, lean body mass, and total body water.

2. Conducting the BIA Measurement

BIA measurement is simple and quick. The patient stands barefoot on a scale or holds electrodes connected to the device, which sends a low-level electrical current through the body. This current is typically imperceptible and safe for patients, including those with T2DM. The device records the resistance and reactance (the degree to which the current is slowed by tissues) and processes these values to determine body composition metrics.

Clinical Applications of Bioimpedance Analysis in T2DM Management

Bioimpedance analysis (BIA) is increasingly recognized as an essential tool in the clinical management of type 2 diabetes mellitus (T2DM). It provides valuable insights into body composition, which helps healthcare professionals optimize treatment strategies, monitor disease progression, and assess the impact of lifestyle interventions. BIA's ability to measure fat mass, lean body mass, hydration status, and visceral fat levels is particularly relevant for managing T2DM, as these factors are closely linked to metabolic health and glycemic control. Here, we explore the specific applications of BIA in T2DM management.

Visceral fat, located around the abdominal organs, is metabolically active and strongly associated with insulin resistance—a hallmark of T2DM. Elevated levels of visceral fat increase the risk of cardiovascular complications, poor glycemic control, and inflammatory responses, all of which worsen diabetes outcomes. BIA provides an estimation of visceral fat levels, allowing clinicians to monitor and manage this risk factor over time.

By tracking reductions in visceral fat, healthcare providers can gauge the effectiveness of dietary changes,

physical activity, and medications aimed at improving insulin sensitivity. Reducing visceral fat has been shown to significantly improve insulin action and lower blood glucose levels, underscoring the importance of this metric in T2DM management.

Sarcopenia, or the loss of muscle mass and strength, is a common issue in T2DM, particularly among older adults. Muscle tissue plays a crucial role in glucose metabolism, as it is a major site for insulin-stimulated glucose uptake. Loss of muscle mass can, therefore, exacerbate insulin resistance, making glycemic control more challenging. BIA provides a quick and reliable measure of lean body mass, which allows for early detection of sarcopenia.

Identifying sarcopenia early enables timely interventions, such as resistance training exercises and protein supplementation, to preserve or rebuild muscle mass. Addressing sarcopenia can improve muscle function, enhance insulin sensitivity, and contribute to better blood sugar management in T2DM patients.

Hydration status is crucial for patients with T2DM, as fluctuations in blood glucose levels can affect body water distribution. Hyperglycemia, for example, can lead to dehydration due to osmotic diuresis, putting patients at risk of acute kidney injury and electrolyte imbalances. BIA assesses total body water (TBW) and distinguishes between intracellular and extracellular water, providing a comprehensive view of hydration status.

For patients with T2DM, maintaining optimal hydration is important not only for kidney health but also for stabilizing blood glucose levels. BIA helps clinicians monitor hydration and make adjustments to fluid intake recommendations, which can reduce the risk of dehydration and related complications, especially in patients with diabetic nephropathy or autonomic dysfunction affecting thirst perception.

Lifestyle interventions, including dietary changes and exercise, are fundamental components of T2DM management. However, traditional metrics like weight and BMI may not fully capture the impact of these interventions on body composition. BIA provides a more detailed picture by assessing changes in fat mass, lean body mass, and visceral fat.

Regular BIA assessments during follow-up visits provide a clear picture of how patients respond to these interventions, allowing healthcare providers to tailor recommendations and keep patients motivated. Certain medications prescribed for T2DM, such as insulin and some sulfonylureas, can lead to weight gain, which may adversely affect body composition by increasing fat mass. Conversely, other medications, such as GLP-1 receptor agonists and SGLT-2 inhibitors, have been shown to reduce body weight and visceral fat. BIA can help clinicians evaluate how different pharmacological treatments impact body composition. By tracking changes in fat mass and lean body mass, healthcare providers can make informed decisions on adjusting or switching medications to achieve optimal metabolic control while minimizing weight gain. BIA can also highlight the need for additional lifestyle support to counteract any negative effects on body composition.

BIA offers valuable data that aids in risk stratification for T2DM-related complications. For example, patients with high visceral fat levels and low muscle mass are at an increased risk for cardiovascular diseases, neuropathy, and other complications. By assessing these factors, BIA enables clinicians to identify patients at higher risk and prioritize preventive measures. Regular BIA monitoring provides insights into which patients may require more intensive interventions, such as stricter glycemic targets, increased physical activity, or closer monitoring for early signs of complications. This proactive approach can help reduce hospitalizations, improve patient outcomes, and minimize healthcare costs.

One of the most impactful benefits of BIA in clinical practice is its ability to visually demonstrate changes in body composition to patients. Showing patients tangible improvements in muscle mass or reductions in visceral fat can increase motivation and adherence to treatment plans, especially when lifestyle changes are recommended. Patients are more likely to follow dietary and exercise regimens when they understand the connection between body composition and diabetes outcomes. For example, seeing a decrease in visceral fat can reinforce the importance of physical activity, while improvements in lean body mass can encourage

continued strength training.

While bioimpedance analysis (BIA) is a useful tool in the management of type 2 diabetes mellitus (T2DM), it has several limitations that can affect its accuracy and utility in clinical settings. Understanding these limitations is crucial for clinicians to interpret BIA results correctly and avoid over-reliance on the technology. One of the primary limitations of BIA is its sensitivity to hydration status. Since BIA measures electrical impedance, and water conducts electricity well, any changes in a patient's hydration level can affect readings. Dehydration tends to increase impedance, leading to an overestimation of body fat, while over hydration reduces impedance, which may underestimate body fat. For T2DM patients, hydration status can vary widely due to factors like osmotic diuresis, kidney dysfunction, or poor fluid intake. This variability in hydration can make it challenging to obtain consistent results over time, which may lead to inaccurate assessments of body composition if not carefully controlled.

Eating and physical activity prior to BIA measurements can significantly influence the results. Consuming food and beverages increases the water content in the gastrointestinal tract, potentially altering impedance readings and causing inaccurate measurements of fat and lean mass. Physical activity, especially intense exercise, can also affect results by redistributing water in the body, causing temporary shifts in impedance. Patients are usually advised to avoid food, beverages, and vigorous exercise for a few hours before a BIA measurement to minimize these effects. However, this requirement can be challenging in clinical practice, especially for patients who need frequent assessments or have unpredictable schedules.

Different BIA devices use varying algorithms to calculate body composition metrics, and these algorithms are often proprietary. Factors such as age, gender, height, and weight are incorporated into these calculations, but differences in algorithms between devices can result in inconsistencies in body composition measurements. This lack of standardization can make it difficult to compare results across different devices or clinical settings. For instance, a patient assessed with one type of BIA device may show different results when evaluated with another device, which could lead to challenges in tracking changes over time or interpreting longitudinal data accurately.

While some advanced BIA devices offer an estimate of visceral fat, they do not provide the precise, detailed measurements that other imaging methods, such as magnetic resonance imaging (MRI) or computed tomography (CT), can offer. This limitation is important for T2DM management because visceral fat plays a crucial role in insulin resistance and metabolic health.

BIA's visceral fat estimates are generally based on predictive models rather than direct measurements, making it less reliable for patients who require accurate visceral fat assessment. Consequently, BIA should be supplemented with other imaging techniques when precise visceral fat quantification is necessary for decision-making.

Edema, which is common in some T2DM patients, especially those with kidney disease or heart failure, can affect the accuracy of BIA measurements. BIA may not accurately differentiate between normal hydration and fluid retention, leading to overestimation of lean body mass and underestimation of fat mass in patients with edema.

This limitation is particularly relevant for T2DM patients with complications like nephropathy or cardiovascular disease, where fluid balance is often disrupted. For these patients, BIA measurements may be less reliable, and alternative methods, such as dual-energy X-ray absorptiometry (DXA), may be preferred for accurate body composition assessment.

Segmental BIA devices, which measure body composition in specific regions (e.g., arms, legs, torso), are more complex and can provide more detailed information. However, they require specific positioning and adherence to standardized measurement procedures, which can be challenging in a clinical setting. Inconsistent positioning or movement during measurement can lead to errors in segmental BIA readings. This limitation is particularly important when monitoring regional fat or muscle changes, as even slight inconsistencies in measurement protocols can affect accuracy.

BIA accuracy can be reduced in individuals with extreme body types, such as those who are very lean, very muscular, or obese. In obese patients, BIA may overestimate body fat due to the way electrical currents pass through excess adipose tissue, while in very muscular patients, it may underestimate fat mass and overestimate muscle mass.

This variability can make BIA less accurate in assessing body composition in patients with T2DM who fall outside typical body size ranges. For these patients, clinicians may need to rely on other assessment methods or interpret BIA results with caution.

Environmental factors, such as temperature and humidity, can also impact BIA measurements. Since BIA relies on the body's electrical conductivity, environmental conditions that affect body temperature or skin moisture may influence results. Cold environments, for example, may constrict blood vessels and increase impedance, affecting body composition estimates.

In clinical practice, maintaining a controlled environment may be challenging, especially in outpatient settings. This variability can introduce minor inaccuracies that may be negligible in routine cases but can impact precision when tracking small changes over time.

Conclusion

Bioimpedance analysis (BIA) serves as a valuable tool in the clinical management of type 2 diabetes mellitus (T2DM), providing essential insights into body composition, including fat mass, lean body mass, hydration status, and visceral fat. These metrics are critical in personalizing care for T2DM patients, enabling healthcare providers to monitor disease progression, assess the effects of lifestyle and pharmacological interventions, and address factors that contribute to insulin resistance and metabolic health. However, BIA is not without limitations. Its accuracy can be affected by hydration status, food intake, physical activity, and the presence of conditions like edema, all of which are common concerns in T2DM management. Moreover, device variability, environmental factors, and limitations in estimating visceral fat mean that BIA should be interpreted cautiously and supplemented with additional assessments when needed. Despite these limitations, BIA's accessibility, cost-effectiveness, and non-invasive nature make it a practical choice for routine use in T2DM care. By understanding the technology's strengths and limitations, clinicians can use BIA as part of a comprehensive approach to diabetes management, supporting tailored treatment plans that address the unique body composition needs of each patient. With careful application and interpretation, BIA can enhance T2DM outcomes, improving both quality of life and long-term health for patients managing this complex condition.

References:

1. Ilhomjon ogli, M. U., Ibrohimjon ogli, S. Z., & Qurbonbek ogli, D. S. (2024). CLINICS AND RESULTS OF TREATMENT OF PATIENTS WITH CORONAVIRUS INFECTION COMPLICATED BY INTERSTITIAL PNEUMONIA IN THE FERGHANA REGION. MODELS AND METHODS FOR INCREASING THE EFFICIENCY OF INNOVATIVE RESEARCH, 3(30), 21-26.
2. Mahmudov, U. I. (2023). COMPARATIVE CHARACTERISTICS OF CLINICAL AND LABORATORY PARAMETERS OF PATIENTS OF THE DIABETIC FOOT DEPARTMENT, DEPENDING ON THE PRESENCE OR ABSENCE OF DIABETES MELLITUS. SO 'NGI ILMIY TADQIQOTLAR NAZARIYASI, 6(12), 355-360.
3. Mahmudov, U. I. (2024). MANAGEMENT OF THYROID NODULES. JOURNAL OF INNOVATIONS IN SCIENTIFIC AND EDUCATIONAL RESEARCH, 7(4), 1-7.
4. Saydaxmedov, Z. I., & Mahmudov, U. I. (2023). CLINICAL AND FUNCTIONAL STATUS OF THE CARDIOVASCULAR SYSTEM IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE WITH COVID-19. SCIENTIFIC ASPECTS AND TRENDS IN THE FIELD OF SCIENTIFIC RESEARCH, 2(16), 44-47.

5. Kamalovich, S. I. (2024). Congenital Esophageal Malformations in Children, Symptoms, Diagnosis and Treatment. *Miasto Przyszłości*, 53, 1241-1243.
6. Zakhriddinovich, I. B. (2024, June). Migraine in Children and its Causes, Symptoms and Treatment. In *Interdisciplinary Conference of Young Scholars in Social Sciences (USA)* (Vol. 7, pp. 29-32).
7. Zakhriddinovich, I. B. (2024, May). Febrile Seizure Disease and its Symptoms, Treatment. In *International Congress on Biological, Physical And Chemical Studies (ITALY)* (pp. 121-124).
8. Алимова, И. А., Райимова, З. М., Бабаджанова, Х. М., & АКТУАЛЬНОСТЬ, В. (2022). РАННЕГО ВМЕШАТЕЛЬСТВА В СЕМЕЙНЫЕ ПОЛИКЛИНИКИ ДЕТЯМ РАННЕГО ВОЗРАСТА. *JOURNAL OF CLINICAL AND PREVENTIVE MEDICINE*, 2, 5-11.
9. Alimova, I. (2021, January). BOLA TARBIYASIDA OTA-ONALARNING PSIXOLOGIK BILIMLARNI SHAKLLANTIRISHNING AHAMIYATI. In *INTERNATIONAL CONFERENCES ON LEARNING AND TEACHING* (Vol. 1, No. 1, pp. 131-132).
10. Анваровна А.И., Мелибаевна Б.Х., Мамамаджоновна Р.З., Захриддинович И.Б., Исломкулович У.М. (2023). Актуальность внедрения службы комплексного раннего вмешательства в семейных клиниках. *ВіоГеко Журнал новозеландської герпетології*, 12 (03), 1139-1145.
11. Anvarovna, A. I., & Melibaeva, B. K. (2022). JUVENILE IDIOPATHIC ARTHRITIS. *SCIENTIFIC JOURNAL OF RESEARCH IN MEDICINE (SJRM)*, 1(4), 6-8.
12. Melibayeva, B. X. (2023). Measures to Improve the Quality of Life of Patients with Comorbid Heart Pathology and Increase the Effectiveness of Their Treatment. *Scholastic: Journal of Natural and Medical Education*, 2(3), 34-36.
13. Kamalovich, S. I. (2024, May). CONGENITAL HEART DEFECTS IN CHILDREN. In *Proceedings of International Conference on Modern Science and Scientific Studies* (Vol. 3, No. 5, pp. 65-71).
14. Rayimov, G. N., Tillaboldiyev, A. R., Saloxiddinov, N., & Sh, D. S. (2022). Actical Errors in Surgical Treatment of Strengthened Abdominal Hernias. *The Peerian Journal*, 5, 130-135.
15. Isakjonovich, S. M. (2024). Effectivness of Aromatherapy in Post-Covid Syndrome. *Miasto Przyszłości*, 49, 1239-1242.
16. Nazirtashova, R. M. (2023). XALQ TABOBATIDA MAKKAJO „RINING O „RNI. *Journal of Chemistry of Goods and Traditional Medicine*, 2(1), 210-216.
17. Mamadaliyeva, N. R. (2023). INSONIYAT O'ZINI O'ZI ZAHARLAMOQDA. " GERMANY" MODERN SCIENTIFIC RESEARCH: ACHIEVEMENTS, INNOVATIONS AND DEVELOPMENT PROSPECTS, 9(1).
18. Nazirtashova, R. M., & Kirgizov, S. M. (2021). Research Of Pentosal Hydrolysis Products Of Plant Waste. *The American Journal of Applied sciences*, 3(04), 126-130.
19. Matyakubov, R., & Nazirtashova, R. M. (2021). Valuable Raw Materials For Producing Furfural. *The American Journal of Interdisciplinary Innovations and Research*, 3(06), 159-165.
20. Назирташова, Р. М. (2022). ДИНАМИЧЕСКОЕ ИССЛЕДОВАНИЕ КАРДИОРЕСПИРАТОРНОЙ СИСТЕМЫ УЧЕНИКОВ СПОРТИВНЫХ ШКОЛ К ОБУЧЕНИЮ В УСЛОВИЯХ ПОВЫШЕННОЙ СЛОЖНОСТИ. *BARQARORLIK VA YETAKSHI TADQIQOTLAR ONLAYN ILMIY JURNALI*, 90-94.
21. Анварова, З. (2024). СПИД/ВИЧ ИФИЦИРОВАНИЕ И ДЕТИ. *THEORY AND ANALYTICAL ASPECTS OF RECENT RESEARCH*, 2(22), 41-45.
22. Анварова, З. (2024). ЗАДЕРЖКА ВНУТРИУТРОБНОГО РАЗВИТИЯ ПЛОДА КАК ФАКТОР НАРУШЕНИЯ ГАРМОНИЧНОГО РАЗВИТИЯ ДЕТЕЙ. *THEORY AND ANALYTICAL ASPECTS OF RECENT RESEARCH*, 2(21), 234-237.
23. Qosimovna, A. Z. (2023). Factors that lead to asphyxia in babies. *American Journal of Pediatric Medicine and Health Sciences* (2993-2149), 1(10), 740-743.

24. Абдуллаев, С. (2024). АКТУАЛЬНОСТЬ ПРОБЛЕМ РАЗВИТИЯ ОСТРЫХ ПНЕВМОНИЙ У ДЕТЕЙ. THEORY AND ANALYTICAL ASPECTS OF RECENT RESEARCH, 2(22), 29-33.
25. Mukhtarzhanovna, I. G. (2024, May). Development of Principles of Study and Treatment of Vaginal Dysbiosis During Pregnancy. In International Congress on Biological, Physical And Chemical Studies (ITALY) (pp. 112-115).
26. Mukhtorjonovna, I. G. (2024). Modern Surgical Methods of Placental Aggregation. Web of Semantics: Journal of Interdisciplinary Science, 2(5), 412-416.
27. Solijon o'g'li, A. S. (2024). BACTERIAL, VIRAL AND MUCOPLASMA PNEUMONIA IN CHILDREN. American Journal of Pediatric Medicine and Health Sciences (2993-2149), 2(1), 273-280.
28. Абдуллаев, С. (2024). ПСИХОЛОГИЧЕСКИЕ ОСОБЕННОСТИ УЧЕБНЫХ ИГР В ПОДГОТОВКЕ СТУДЕНТОВ МЕДИЦИНСКИХ ИНСТИТУТОВ. FORMATION OF PSYCHOLOGY AND PEDAGOGY AS INTERDISCIPLINARY SCIENCES, 2(25), 222-224.
29. Александровна, А.Е. (2023). ОСНОВНЫЕ АСПЕКТЫ РЕСПИРАТОРНОЙ РЕАБИЛИТАЦИИ ПОСЛЕДСТВИЙ НОВОЙ КОРОНАВИРУСНОЙ ИНФЕКЦИИ У ДЕТЕЙ С БРОНХОЛЕГОЧНЫМИ ЗАБОЛЕВАНИЯМИ. Всемирный бюллетень социальных наук , 18 , 81-83.
30. Abdullaev, S. S. (2023). TO THE QUESTION OF COMMUNITY-ACCOMPANIED PNEUMONIA IN YOUNG CHILDREN. Journal of Social Sciences and Humanities Research Fundamentals, 3(05), 51-53.
31. Худайназарова, С. Р., Курьязова, Ш. М., & Охунова, М. Ж. (2023). ОСОБЕННОСТИ БРОНХООБСТРУКТИВНОГО СИНДРОМА ПРИ ВНЕБОЛЬНИЧНОЙ ПНЕВМОНИИ У ДЕТЕЙ РАННЕГО ВОЗРАСТА. Interpretation and researches, 1(6).
32. Анварова, З. (2024). СПИД/ВИЧ ИФИЦИРОВАНИЕ И ДЕТИ. THEORY AND ANALYTICAL ASPECTS OF RECENT RESEARCH, 2(22), 41-45.
33. Анварова, З. (2024). ЗАДЕРЖКА ВНУТРИУТРОБНОГО РАЗВИТИЯ ПЛОДА КАК ФАКТОР НАРУШЕНИЯ ГАРМОНИЧНОГО РАЗВИТИЯ ДЕТЕЙ. THEORY AND ANALYTICAL ASPECTS OF RECENT RESEARCH, 2(21), 234-237.
34. Alexandrovna, A. E. (2023). Clinical and functional features of the bronchopulmonary system in chronic kidney disease. Texas Journal of Medical Science, 16, 57-59.
35. Qosimovna, A. Z. (2023). Factors that lead to asphyxia in babies. American Journal of Pediatric Medicine and Health Sciences (2993-2149), 1(10), 740-743.
36. Kamoldinovich, X. D. (2024). INTRAVENOUS ADMINISTRATION OF CONTRAST AGENTS AND ITS CHARACTERISTICS. Miasto Przyszłości, 48, 119-131.
37. Kamoldinovich, X. D. (2024, May). MODERN CONTRAST AGENTS IN MEDICINE. In Proceedings of Scientific Conference on Multidisciplinary Studies (Vol. 3, No. 5, pp. 132-144).
38. Kamoldinovich, X. D. (2024, May). ULTRASOUND LEVELING AND ITS ADVANTAGES. In Proceedings of International Conference on Modern Science and Scientific Studies (Vol. 3, No. 5, pp. 59-64).