GEORGIAN MEDICAL MEWS

ISSN 1512-0112

NO 7-8 (364-365) Июль-Август 2025

ТБИЛИСИ - NEW YORK



ЕЖЕМЕСЯЧНЫЙ НАУЧНЫЙ ЖУРНАЛ

Медицинские новости Грузии საქართველოს სამედიცინო სიახლენი

GEORGIAN MEDICAL NEWS

Monthly Georgia-US joint scientific journal published both in electronic and paper formats of the Agency of Medical Information of the Georgian Association of Business Press. Published since 1994. Distributed in NIS, EU and USA.

GMN: Georgian Medical News is peer-reviewed, published monthly journal committed to promoting the science and art of medicine and the betterment of public health, published by the GMN Editorial Board since 1994. GMN carries original scientific articles on medicine, biology and pharmacy, which are of experimental, theoretical and practical character; publishes original research, reviews, commentaries, editorials, essays, medical news, and correspondence in English and Russian.

GMN is indexed in MEDLINE, SCOPUS, PubMed and VINITI Russian Academy of Sciences. The full text content is available through EBSCO databases.

GMN: Медицинские новости Грузии - ежемесячный рецензируемый научный журнал, издаётся Редакционной коллегией с 1994 года на русском и английском языках в целях поддержки медицинской науки и улучшения здравоохранения. В журнале публикуются оригинальные научные статьи в области медицины, биологии и фармации, статьи обзорного характера, научные сообщения, новости медицины и здравоохранения. Журнал индексируется в MEDLINE, отражён в базе данных SCOPUS, PubMed и ВИНИТИ РАН. Полнотекстовые статьи журнала доступны через БД EBSCO.

GMN: Georgian Medical News – საქართველოს სამედიცინო სიახლენი – არის ყოველთვიური სამეცნიერო სამედიცინო რეცენზირებადი ჟურნალი, გამოიცემა 1994 წლიდან, წარმოადგენს სარედაქციო კოლეგიისა და აშშ-ის მეცნიერების, განათლების, ინდუსტრიის, ხელოვნებისა და ბუნებისმეტყველების საერთაშორისო აკადემიის ერთობლივ გამოცემას. GMN-ში რუსულ და ინგლისურ ენებზე ქვეყნდება ექსპერიმენტული, თეორიული და პრაქტიკული ხასიათის ორიგინალური სამეცნიერო სტატიები მედიცინის, ბიოლოგიისა და ფარმაციის სფეროში, მიმოხილვითი ხასიათის სტატიები.

ჟურნალი ინდექსირებულია MEDLINE-ის საერთაშორისო სისტემაში, ასახულია SCOPUS-ის, PubMed-ის და ВИНИТИ РАН-ის მონაცემთა ბაზებში. სტატიების სრული ტექსტი ხელმისაწვდომია EBSCO-ს მონაცემთა ბაზებიდან.

WEBSITE

www.geomednews.com

К СВЕДЕНИЮ АВТОРОВ!

При направлении статьи в редакцию необходимо соблюдать следующие правила:

- 1. Статья должна быть представлена в двух экземплярах, на русском или английском языках, напечатанная через полтора интервала на одной стороне стандартного листа с шириной левого поля в три сантиметра. Используемый компьютерный шрифт для текста на русском и английском языках Times New Roman (Кириллица), для текста на грузинском языке следует использовать AcadNusx. Размер шрифта 12. К рукописи, напечатанной на компьютере, должен быть приложен CD со статьей.
- 2. Размер статьи должен быть не менее десяти и не более двадцати страниц машинописи, включая указатель литературы и резюме на английском, русском и грузинском языках.
- 3. В статье должны быть освещены актуальность данного материала, методы и результаты исследования и их обсуждение.

При представлении в печать научных экспериментальных работ авторы должны указывать вид и количество экспериментальных животных, применявшиеся методы обезболивания и усыпления (в ходе острых опытов).

- 4. К статье должны быть приложены краткое (на полстраницы) резюме на английском, русском и грузинском языках (включающее следующие разделы: цель исследования, материал и методы, результаты и заключение) и список ключевых слов (key words).
- 5. Таблицы необходимо представлять в печатной форме. Фотокопии не принимаются. Все цифровые, итоговые и процентные данные в таблицах должны соответствовать таковым в тексте статьи. Таблицы и графики должны быть озаглавлены.
- 6. Фотографии должны быть контрастными, фотокопии с рентгенограмм в позитивном изображении. Рисунки, чертежи и диаграммы следует озаглавить, пронумеровать и вставить в соответствующее место текста в tiff формате.

В подписях к микрофотографиям следует указывать степень увеличения через окуляр или объектив и метод окраски или импрегнации срезов.

- 7. Фамилии отечественных авторов приводятся в оригинальной транскрипции.
- 8. При оформлении и направлении статей в журнал МНГ просим авторов соблюдать правила, изложенные в «Единых требованиях к рукописям, представляемым в биомедицинские журналы», принятых Международным комитетом редакторов медицинских журналов http://www.spinesurgery.ru/files/publish.pdf и http://www.nlm.nih.gov/bsd/uniform_requirements.html В конце каждой оригинальной статьи приводится библиографический список. В список литературы включаются все материалы, на которые имеются ссылки в тексте. Список составляется в алфавитном порядке и нумеруется. Литературный источник приводится на языке оригинала. В списке литературы сначала приводятся работы, написанные знаками грузинского алфавита, затем кириллицей и латиницей. Ссылки на цитируемые работы в тексте статьи даются в квадратных скобках в виде номера, соответствующего номеру данной работы в списке литературы. Большинство цитированных источников должны быть за последние 5-7 лет.
- 9. Для получения права на публикацию статья должна иметь от руководителя работы или учреждения визу и сопроводительное отношение, написанные или напечатанные на бланке и заверенные подписью и печатью.
- 10. В конце статьи должны быть подписи всех авторов, полностью приведены их фамилии, имена и отчества, указаны служебный и домашний номера телефонов и адреса или иные координаты. Количество авторов (соавторов) не должно превышать пяти человек.
- 11. Редакция оставляет за собой право сокращать и исправлять статьи. Корректура авторам не высылается, вся работа и сверка проводится по авторскому оригиналу.
- 12. Недопустимо направление в редакцию работ, представленных к печати в иных издательствах или опубликованных в других изданиях.

При нарушении указанных правил статьи не рассматриваются.

REQUIREMENTS

Please note, materials submitted to the Editorial Office Staff are supposed to meet the following requirements:

- 1. Articles must be provided with a double copy, in English or Russian languages and typed or computer-printed on a single side of standard typing paper, with the left margin of 3 centimeters width, and 1.5 spacing between the lines, typeface Times New Roman (Cyrillic), print size 12 (referring to Georgian and Russian materials). With computer-printed texts please enclose a CD carrying the same file titled with Latin symbols.
- 2. Size of the article, including index and resume in English, Russian and Georgian languages must be at least 10 pages and not exceed the limit of 20 pages of typed or computer-printed text.
- 3. Submitted material must include a coverage of a topical subject, research methods, results, and review.

Authors of the scientific-research works must indicate the number of experimental biological species drawn in, list the employed methods of anesthetization and soporific means used during acute tests.

- 4. Articles must have a short (half page) abstract in English, Russian and Georgian (including the following sections: aim of study, material and methods, results and conclusions) and a list of key words.
- 5. Tables must be presented in an original typed or computer-printed form, instead of a photocopied version. Numbers, totals, percentile data on the tables must coincide with those in the texts of the articles. Tables and graphs must be headed.
- 6. Photographs are required to be contrasted and must be submitted with doubles. Please number each photograph with a pencil on its back, indicate author's name, title of the article (short version), and mark out its top and bottom parts. Drawings must be accurate, drafts and diagrams drawn in Indian ink (or black ink). Photocopies of the X-ray photographs must be presented in a positive image in **tiff format**.

Accurately numbered subtitles for each illustration must be listed on a separate sheet of paper. In the subtitles for the microphotographs please indicate the ocular and objective lens magnification power, method of coloring or impregnation of the microscopic sections (preparations).

- 7. Please indicate last names, first and middle initials of the native authors, present names and initials of the foreign authors in the transcription of the original language, enclose in parenthesis corresponding number under which the author is listed in the reference materials.
- 8. Please follow guidance offered to authors by The International Committee of Medical Journal Editors guidance in its Uniform Requirements for Manuscripts Submitted to Biomedical Journals publication available online at: http://www.nlm.nih.gov/bsd/uniform_requirements.html http://www.icmje.org/urm_full.pdf
- In GMN style for each work cited in the text, a bibliographic reference is given, and this is located at the end of the article under the title "References". All references cited in the text must be listed. The list of references should be arranged alphabetically and then numbered. References are numbered in the text [numbers in square brackets] and in the reference list and numbers are repeated throughout the text as needed. The bibliographic description is given in the language of publication (citations in Georgian script are followed by Cyrillic and Latin).
- 9. To obtain the rights of publication articles must be accompanied by a visa from the project instructor or the establishment, where the work has been performed, and a reference letter, both written or typed on a special signed form, certified by a stamp or a seal.
- 10. Articles must be signed by all of the authors at the end, and they must be provided with a list of full names, office and home phone numbers and addresses or other non-office locations where the authors could be reached. The number of the authors (co-authors) must not exceed the limit of 5 people.
- 11. Editorial Staff reserves the rights to cut down in size and correct the articles. Proof-sheets are not sent out to the authors. The entire editorial and collation work is performed according to the author's original text.
- 12. Sending in the works that have already been assigned to the press by other Editorial Staffs or have been printed by other publishers is not permissible.

Articles that Fail to Meet the Aforementioned Requirements are not Assigned to be Reviewed.

ᲐᲕᲢᲝᲠᲗᲐ ᲡᲐᲧᲣᲠᲐᲓᲦᲔᲑᲝᲓ!

რედაქციაში სტატიის წარმოდგენისას საჭიროა დავიცვათ შემდეგი წესები:

- 1. სტატია უნდა წარმოადგინოთ 2 ცალად, რუსულ ან ინგლისურ ენებზე,დაბეჭდილი სტანდარტული ფურცლის 1 გვერდზე, 3 სმ სიგანის მარცხენა ველისა და სტრიქონებს შორის 1,5 ინტერვალის დაცვით. გამოყენებული კომპიუტერული შრიფტი რუსულ და ინგლისურენოვან ტექსტებში Times New Roman (Кириллица), ხოლო ქართულენოვან ტექსტში საჭიროა გამოვიყენოთ AcadNusx. შრიფტის ზომა 12. სტატიას თან უნდა ახლდეს CD სტატიით.
- 2. სტატიის მოცულობა არ უნდა შეადგენდეს 10 გვერდზე ნაკლებს და 20 გვერდზე მეტს ლიტერატურის სიის და რეზიუმეების (ინგლისურ,რუსულ და ქართულ ენებზე) ჩათვლით.
- 3. სტატიაში საჭიროა გაშუქდეს: საკითხის აქტუალობა; კვლევის მიზანი; საკვლევი მასალა და გამოყენებული მეთოდები; მიღებული შედეგები და მათი განსჯა. ექსპერიმენტული ხასიათის სტატიების წარმოდგენისას ავტორებმა უნდა მიუთითონ საექსპერიმენტო ცხოველების სახეობა და რაოდენობა; გაუტკივარებისა და დაძინების მეთოდები (მწვავე ცდების პირობებში).
- 4. სტატიას თან უნდა ახლდეს რეზიუმე ინგლისურ, რუსულ და ქართულ ენებზე არანაკლებ ნახევარი გვერდის მოცულობისა (სათაურის, ავტორების, დაწესებულების მითითებით და უნდა შეიცავდეს შემდეგ განყოფილებებს: მიზანი, მასალა და მეთოდები, შედეგები და დასკვნები; ტექსტუალური ნაწილი არ უნდა იყოს 15 სტრიქონზე ნაკლები) და საკვანძო სიტყვების ჩამონათვალი (key words).
- 5. ცხრილები საჭიროა წარმოადგინოთ ნაბეჭდი სახით. ყველა ციფრული, შემაჯამებელი და პროცენტული მონაცემები უნდა შეესაბამებოდეს ტექსტში მოყვანილს.
- 6. ფოტოსურათები უნდა იყოს კონტრასტული; სურათები, ნახაზები, დიაგრამები დასათაურებული, დანომრილი და სათანადო ადგილას ჩასმული. რენტგენოგრამების ფოტოასლები წარმოადგინეთ პოზიტიური გამოსახულებით tiff ფორმატში. მიკროფოტო-სურათების წარწერებში საჭიროა მიუთითოთ ოკულარის ან ობიექტივის საშუალებით გადიდების ხარისხი, ანათალების შეღებვის ან იმპრეგნაციის მეთოდი და აღნიშნოთ სუ-რათის ზედა და ქვედა ნაწილები.
- 7. სამამულო ავტორების გვარები სტატიაში აღინიშნება ინიციალების თანდართვით, უცხოურისა უცხოური ტრანსკრიპციით.
- 8. სტატიას თან უნდა ახლდეს ავტორის მიერ გამოყენებული სამამულო და უცხოური შრომების ბიბლიოგრაფიული სია (ბოლო 5-8 წლის სიღრმით). ანბანური წყობით წარმოდგენილ ბიბლიოგრაფიულ სიაში მიუთითეთ ჯერ სამამულო, შემდეგ უცხოელი ავტორები (გვარი, ინიციალები, სტატიის სათაური, ჟურნალის დასახელება, გამოცემის ადგილი, წელი, ჟურნალის №, პირველი და ბოლო გვერდები). მონოგრაფიის შემთხვევაში მიუთითეთ გამოცემის წელი, ადგილი და გვერდების საერთო რაოდენობა. ტექსტში კვადრატულ ფჩხილებში უნდა მიუთითოთ ავტორის შესაბამისი N ლიტერატურის სიის მიხედვით. მიზანშეწონილია, რომ ციტირებული წყაროების უმეტესი ნაწილი იყოს 5-6 წლის სიღრმის.
- 9. სტატიას თან უნდა ახლდეს: ა) დაწესებულების ან სამეცნიერო ხელმძღვანელის წარდგინება, დამოწმებული ხელმოწერითა და ბეჭდით; ბ) დარგის სპეციალისტის დამოწმებული რეცენზია, რომელშიც მითითებული იქნება საკითხის აქტუალობა, მასალის საკმაობა, მეთოდის სანდოობა, შედეგების სამეცნიერო-პრაქტიკული მნიშვნელობა.
- 10. სტატიის ბოლოს საჭიროა ყველა ავტორის ხელმოწერა, რომელთა რაოდენობა არ უნდა აღემატებოდეს 5-ს.
- 11. რედაქცია იტოვებს უფლებას შეასწოროს სტატია. ტექსტზე მუშაობა და შეჯერება ხდება საავტორო ორიგინალის მიხედვით.
- 12. დაუშვებელია რედაქციაში ისეთი სტატიის წარდგენა, რომელიც დასაბეჭდად წარდგენილი იყო სხვა რედაქციაში ან გამოქვეყნებული იყო სხვა გამოცემებში.

აღნიშნული წესების დარღვევის შემთხვევაში სტატიები არ განიხილება.

GEORGIAN MEDICAL NEWS NO 7-8 (364-365) 2025

Содержание:

Babry I. Oren, Marina I. Devdariani, Gela V. Beselia, Nino N. Sikharulidze, Manana G. Dashniani, Maia A. Burjanadze, Ia R. Kvachakidze, Marina I. Nebieridze, Lena Sh. Davlianidze, Lali M. Gumberidze, Nodar P. Mitagvaria. ROLE OF ANTIOXIDANT FOLIUM EXPOSURE ON OXIDATIVE SRESS IN A VALPROIC ACID-INDUCED ANIMAL MODEL OF AUTISM
Hajdi Gorica, Pavllo Djamandi, Gentian Vyshka. DELAYED ONSET OF MYASTHENIA GRAVIS FOLLOWING COLECTOMY FOR ULCERATIVE COLITIS: A CASE STUDY16-17
Zhadyra Yersariyeva, Bagdad Suleyeva, Botagoz Turdaliyeva, Yeldos Tussipbayev. HEMOSTASIS GENE POLYMORPHISM IN RETINAL VASCULAR OCCLUSION: A SYSTEMATIC REVIEW
Ilia Nakashidze, Nameera Parveen Shaikh, Shota Nakashidze, Aleena Parveen Shaikh, Sarfraz Ahmad, Irina Nakashidze. EVALUATION OF TNF-A LEVELS IN MALE PATIENTS WITH STROKE: PROGNOSTIC IMPLICATIONS
Yerbolat Iztleuov, Marat Iztleuov, Altynbek Dushmanov, Gulmira Iztleuova. PREVENTION IN THE PARENTAL GENERATION OF EXPOSED RATS: CONSEQUENCES OF TOXIC EXPOSURE TO CHROMIUM AND GAMMA IRRADIATION IN AN EXPERIMENTAL MODEL
Rashid Nassar, Nadine Khayyat, Michele Halasa, Fahad Hussain. TRAUMATIC ANTERIOR SHOULDER INSTABILITY (TUBS): A NARRATIVE REVIEW OF CURRENT LITERATURE46-50
Albadawi Abdelbagi Talha, Mawaheip A. Abdo Jeweser, Abubakr Ali Elamin Mohamed Ahmed, Abdelrahman Eldaw Mohammed, Elhadi Abdalla Ahmed, GadAllah Modawe, Sanaa Elfatih Hussein. THE HBV AND HCV SEROPREVALENCE AMONG BLOOD DONORS IN Al-DAMAZIN STATE, SUDAN: A THREE-YEAR RETROSPECTIVESTUDY
Hiba Salah Hasan, Teeb Ali, Kadhim Adnan Ali, Al hassan Ali, Hany A. Al-hussaniy. MODELING DRUG-ORGAN INTERACTIONS AND OPTIMIZING IMMUNOTHERAPY: A QUANTITATIVE SYSTEMS PHARMACOLOGY AND ODRONEXTAMAB DYNAMICS
Zilola Mavlyanova, Davron Ravshanov, Malika Ibragimova, Lola Irbutaeva, Khalimova Fariza, May K. Ismail, Shawgi A. Elsiddig, Marwan Ismail, Salma E R Mohamed, Sara Mohammed Ali. PROGNOSTIC SIGNIFICANCE OF PROLIFERATION (KI-67) AND ANGIOGENESIS (CD34) MARKERS IN MENINGIOMAS FOR THE DEVELOPMENT OF REHABILITATION STRATEGIES
A.R. Abzaliyeva, K.K. Kurakbayev, A.R. Ryskulova, Z.R. Abzaliyev, E. Tasmagambet, D.Zh. Saussanova. TURNOVER INTENTIONS AMONG PHYSICIANS AND NURSES IN KAZAKHSTAN DURING THE COVID-19 PANDEMIC: A CROSS- SECTIONAL STUDY OF PSYCHOLOGICAL AND PROFESSIONAL CHALLENGES
A.A. Mammadov, A.N. Mustafayev, A.H. Aliyev. RADIOLOGICAL IMAGING METHODS FOR ACCURATE DIAGNOSIS OF ABDOMINAL POSTOPERATIVE COMPLICATIONS73-76
I.A. Lebedev, E.V. Zakharchuk, Yu.V. Boldyreva, I.A. Aptekar, E.I. Malinina. OSSIFICATION OF THE POSTERIOR LONGITUDINAL LIGAMENT: A CASE REPORT AND LITERATURE REVIEW
Zhanar Balmukhamedova, Gulmira Derbissalina, Aliya Dzholdasbekova, Dariga Blyalova, Luiza Murzakhalova. SPECKLE-TRACKING ECHOCARDIOGRAPHY FOR EARLY DETECTION OF SUBCLINICAL SYSTOLIC DYSFUNCTION IN PERIMENOPAUSAL WOMEN WITHOUT APPARENT DIASTOLIC DYSFUNCTION
Arkam Thabit Al Neama, Musab Mohammed Khalaf, Ahmed A.J. Mahmood. PATTERNS OF ACETYLCHOLINESTERASE AND BUTYRYLCHOLINESTERASE ACTIVITY IN COMMON CARDIOVASCULAR PHENOTYPES
Argjira Veseli, Shefqet Mrasori, Ivana Čuković-Bagić, Lul Raka, Kaltrina Veseli, Enis Veseli. PARENTAL QUALITY OF LIFE WHEN RAISING CHILDREN WITH AUTISM SPECTRUM DISORDER: A NARRATIVE REVIEW
Anas Ali Alhur, Daliya T. Sendi, Miad M. AlZahrani, Layla T. Abusharha, Rahaf Y. Abudaak, Rahmah Alsinan, Rama R. Alharbi, Lamia Almadhi, Laila M. Alotaibi, Mona A. Hadadi, Shaima H. Alattas, Fatimah Almisbah, Fathi Almisbah, Abdulrahman Alrashed, Kawkab Alharbi. EVALUATING THE TRUSTWORTHINESS OF CHATGPT-GENERATED HEALTH INFORMATION AMONG FUTURE HEALTH CARE PROFESSIONALS
Ting-Ting Wang, Yan Wang. HUMANISTIC CARE NURSING FOR PATIENTS IN THE OPERATING ROOM DURING THE PERIOPERATIVE PERIOD: FULL-CYCLE CARE FROM PHYSIOLOGY TO PSYCHOLOGY
Zauresh Barmanasheva, Mariya Laktionova, Anna Onglas, AyauIym Kossetova, Ivan Melnikov. PREVALENCE AND RISK FACTORS OF UTERINE FIBROIDS IN WOMEN OF REPRODUCTIVE AGE: A FACILITY-BASED STUDY IN A MEGACITY
Bolat Ashirov, Assel Kassymova, Jamilya Mansurova, Andrey Orekhov, Meiramgul Tokbulatova, Mirgul Kapakova, Zhanar Toktarova, Aisulu Zhunuspekova. PROGNOSTIC MARKERS OF ISCHEMIC AND HEMORRHAGIC COMPLICATIONS IN PATIENTS WITH ATRIAL FIBRILLATION

Khalilov Sh. Dzh. ELECTROCARDIOGRAPHY CHARACTERISTICS OF THE PATIENTS WITH NON-ST-ELEVATION MYOCARDIAL INFARCTION (NS TEMI)
Salome Kordzaia, Elene Dolmazashvili, Khatuna Tsiklauri, Lasha Khmaladze, Nana Chikhladze. FROM INFUSION REACTION TO IMMUNE CASCADE: A CASE OF SEQUENTIAL TAXANE AND CAPECITABINE TOXICITIES IN TRIPLE-NEGATIVE BREAST CANCER
Yu Zhu, Fandong Zeng, Weiwei Chang, Liying Wen, Lijun Zhu, Yuelong Jin. AN EMPIRICAL STUDY ON THE ASSOCIATION BETWEEN ASPIRATION INDEX AND ACADEMIC PERFORMANCE AMONG PREVENTIVE MEDICINE STUDENTS
Alaa O Ahmed, Mubarak S Karsany, Mohamed Elfatih Abdelwadoud, Mutaz Ali, Osama Mohamed, Amged Gaffer Mostafa, Hussam Ali Osman, Elryah I Ali, Elyasa Elfaki, Tagwa Yousif Elsayed Yousif, Ayman H. Alfeel, Mohammed Ibrahim Saeed. MOLECULAR DETECTION OF HIGH RISK HUMAN PAPILLOMA VIRUS SUBTYPES IN CERVICAL SMEARS AMONG SUDANESE WOMEN
Tchernev G, Tchernev KG Jr, Krastev DS, Krastev NS, Kordeva S. DERMATOLOGIC SURGERY ROUNDS: RECONSTRUCTIVE SURGERY EMPLOYING THE SHARK ISLAND FLAP FOR BASAL CELL CARCINOMA AFFECTING THE NASAL ALA
Saltanat Imanalieva, Bayan Sagindykova, Rabiga Anarbayeva, Murat Omirali, Gulnara Ospanova, Murat Ashirov. CURRENT STATUS AND PROSPECTS FOR THE DEVELOPMENT OF PEDIATRIC DOSAGE FORMS BY THE EXAMPLE OF COMBINED MELOXICAM AND VITAMIN B12 TABLETS
Ahmed Miri Saadoon. INCIDENCE OF PRESSURE SORE IN THE INTENSIVE CARE UNIT AT AL-DIWANYIA TEACHING HOSPITAL
Isoyan A.S, Danielyan M.H, Antonyan I.V, Azizyan N.H, Mkrtchyan A.A, Karapetyan K.V, Nebogova K.A. MORPHOHISTOCHEMICAL ANALYSIS OF CORTICAL STRUCTURES IN AN EXPERIMENTAL MODEL OF PROLONGED COMPRESSION SYNDROME OF THE HIND LIMB IN RATS
Abdulaziz Alroshodi, Faisal A. Al-Harbi, Rasil Sulaiman Alayed, Fahad M. Alharbi, Khalid A Alkhalifah, Mayadah Assaf Alawaji, Ibrahim S. Alsabhawi. FACTORS IMPACTING HEMODIALYSIS TREATMENT ADHERENCE IN END-STAGE RENAL DISEASE PATIENTS RECEIVING INCENTER HEMODIALYSIS IN QASSIM REGION
Gulshat Alimkhanova, Marat Syzdykbayev, Rinat Ashzhanov, Kulsara Rustemova, Maksut Kazymov, Rustem Kazangapova, Saule Imangazinova, Yernar Kairkhanov, Bazar Tuleuov, Sanzhar Khalelov, Roman Khripunov, Samatbek Abdrakhmanov, Abay Mijatov. THE TRANSVERSUS ABDOMINIS PLANE BLOCK AS A METHOD OF MULTIMODAL OPIOID-SPARING POSTOPERATIVE ANALGESIA: A NARRATIVE REVIEW
Zhengmei Fang, Xiaoling Ran, Lijun Zhu, Yingshui Yao, Yuelong Jin. THE IMPACT OF BMAL1 GENE POLYMORPHISM ON SLEEP QUALITY IN HEALTHY CHINESE YOUTH: A GENDER-SPECIFIC ANALYSIS
Muwafaq H. Zaya, Ahmed A. J. Mahmood, Musab M. Khalaf. CROSS SECTIONAL EVIDENCE FOR OPPOSING EFFECTS OF HYPERGLYCAEMIA AND HYPERLIPIDAEMIA ON CHOLINESTERASEACTIVITIES
Erleta Muçaj, Erëza Durmishi, Serbeze Kabashi Muçaj, Leart Kuçi, Elza Muçaj, Gerta Durmishi. CHALLENGES IN RADIOLOGICAL DIAGNOSIS: CRANIOPHARYNGIOMA VS ASTROCYTOMA
Uday Mahajan, Imran Khan, Ria Gupta, Meraj Akhtar, Vibhore Gupta, Edward Spurrier, Mohamed Kabary, Adnan Asif, Salman Shoukat Ali Parpia.
NAMING CONVENTIONS FOR UNIDENTIFIED PATIENTS IN EMERGENCY AND TRAUMA SETTINGS: A NARRATIVE REVIEW
Xuexue Li, Wenjie Wen, Dandan Ren. MOLECULAR MECHANISMS OF DIABETIC PERIODONTITIS: IDENTIFICATION OF KEY OXIDATIVE STRESS-RELATED GENES AND POTENTIAL THERAPEUTIC ROLE OF METFORMIN THROUGH MMP14 AND PXDN
Davron Ravshanov, Zilola Mavlyanova, Kholmirzayev Bakhtiyor, Malika Tursunovna, Khalimova Fariza. HISTOPATHOLOGICAL PREDICTORS AND FUNCTIONAL RECOVERY IN PATIENTS WITH INTRACRANIAL MENINGIOMAS
Aymuhambetov Y, Khismetova Z A, Iskakova N, Akhmetova K, Serikova-Esengeldina D, Shalgumbayeva G.M. ASSESSMENT OF QUALITY OF LIFE IN BREAST CANCER PATIENTS BY USING EORTC QLQ-C30 QUESTIONNAIRE IN EAST KAZAKHSTANREGION
Yujing Tao, Long Hua, Liu Zhang, Ying Feng, Liying Wen, Weiwei Chang. THE CORRELATION BETWEEN STRESS, ACADEMIC PERFORMANCE, AND SLEEP DISTURBANCES AMONG HIGH SCHOOL STUDENTS IN ANHUI PROVINCE: A CROSS-SECTIONAL STUDY
Fahad AlAmr, Muhannad Essa S. Alghamdi, Ahmed Saeed A. Alghamdi, Osama Khamis A. Alghamdi, Hassan Mahfouz B. Alghamdi, Osama Mesfer S. Alghamdi, Abdullah Ali A. Almimoni, Abdulmalik Ahmed S. Al-Zahrani. PREVALENCE AND ASSOCIATED RISK FACTORS OF NOCTURNAL ENURESIS AMONG CHILDREN AGED 5-18 YEARS IN ALBAHA REGION, SAUDI ARABIA

Aya Saad Aldewachi, Mohammed I Aladul. APPETITIVE TRAITS AND QUALITY OF LIFE IN WOMEN WITH OBESITY USING GLUCAGON-LIKE PEPTIDE-1 RECEPTOR AGONISTS: INSIGHTS FROM A PCOS-ENRICHED SAMPLE
George Shaburishvili, Nikoloz Shaburishvili, Georg Becker, Solomon Zeikidze, Bacho Tsiklauri. INCIDENCE OF ADVERSE EVENTS RESULTING FROM BETA-BLOCKER TITRATION IN PATIENTS WITH HEART FAILURE
Blushinova A.N, Orazalina A.S, Shalgumbayeva G.M. INDUCED ABORTION IN KAZAKHSTAN: WOMEN'S PERCEPTIONS AND EXPERIENCES BASED ON CROSS-SECTIONAL STU DY
Qunru Hu, Liying Wen, Jingqi Zhang, Weiwei Chang, Yuelong Jin, Anshi Wang, Lijun Zhu. IS CORE SELF-EVALUATION A PROTECTIVE FACTOR FOR COLLEGE STUDENTS'MARITAL ATTITUDES? THE MODERATING ROLE OF PSYCHOLOGICAL STATUS
Gulfariza Gani, Ubaidilla Datkhayev, Kairat Zhakipbekov, Serzhan Mombekov, Murat Ashirov, Nurgali Rakhymbayev, Zhanerke Seitova. STUDY OF THE CHEMICAL COMPOSITION AND ANTIMICROBIAL ACTIVITY OF SUBCRITICAL CO ₂ EXTRACT FROM <i>EUPHORBIA HUMIFUSA</i> WILLD
Maysoon Mohammed Hassan, Mohammed Abdulwahab Ati Al-askeri, Naseer Kadhim Jawad. PROGNOSTIC IMPACT OF EGFR2 AND KI-67 OVEREXPRESSION WITH DOWNREGULATION OF <i>miR-17</i> AND <i>miR-1307</i> IN FEMALE BREAST CANCER PATIENTS
Imzharov Talgat Abatovich, Zhakiev Bazylbek Sagidollievich, Sarkulov Marat Nukinovich, Pavlov Valentin Nikolaevich, Kurmangaliev Oleg Maratovich.
THE EFFECTIVENESS OF METAPHYLAXIS OF NEPHROLITHIASIS DURING PERCUTANEOUS NEPHROLITHOTRIPSY: A SYSTEMATIC REVIEW AND META-ANALYSIS
Yan Wang, Ting-Ting Wang, Chang-Sheng He. PROGRESS IN T-CELL IMMUNE RESEARCH ON HYPERLIPIDEMIC PANCREATITIS
Marwan I Abdullah. MINING THE CELLMINER DATABASE TO IDENTIFY SHARED BIOMARKERS OF 5-FU AND OXALIPLATIN RESPONSE327-341
Shyngys Adilgazyuly, Tolkyn Bulegenov, Akmaral Mussakhanova, Tasbolat Adylkhanov, Kanat Abdilov, Zhannur Altybayeva, Gulmira Bazarova, Malike Kudaibergenova, Makpal Alchimbayeva, Aigul Utegenova, Gulnara Otepova. ASSESSING THE INFLUENCE OF MEDICAL EDUCATION REFORMS ON ONCOLOGIST WORKFORCE AND LUNG CANCER MORTALITY IN KAZAKH-STAN: AN INTERRUPTED TIME SERIES ANALYSIS WITH PREDICTIVE MOD-ELING OF NATIONWIDE DATA FROM 1998 TO 2023
Wen-Wen Liu, Zhi-Juan Xu, Fang Xu. NEW INSIGHTS INTO THE PATHOGENESIS AND TREATMENT ADVANCES OF AGE - RELATED MACULAR DEGENERATION
Zhamilya Zholdybay, Zhanar Zhakenova, Madina Gabdullina, Yevgeniya Filippenko, Suria Yessentayeva, Galymzhan Alisherov, Aigerim Mustapaeva, Jandos Amankulov, Ildar Fakhradiyev. 68GA-FAPI PET/CT IN DIAGNOSIS OF THE BREAST CANCER DEPENDING ON THE MOLECULAR SUBTYPES AND EXPRESSION STATUS OF HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2/NEU)
A.I. Rybin, V.E. Maksymovskyi, O.V. Kuznetsova, V.V. Osyk, A.S. Bohdan. THE RESULTS OF LIFE QUALITY ASSESSMENT IN PATIENTS WITH PRIMARY OVARIAN CANCER DURING TREATMENT: EFFECT OF DIFFERENT TACTICS AND HIPEC
Miranda Sejdiu Abazi, Arbër Prokshaj, Shpëtim Prokshaj, Fitim Alidema, Nora Leci, Linda Abazi Morina. ASSESSMENT OF PRACTICAL PERFORMANCE IN ORTHODONTIC CLASP FABRICATION AMONG DENTAL TECHNICIAN STUDENTS AT UBT: A REAL-TIME ANALYSIS OF WORKING TIME AND PERCEIVED STRESS
Abylay Baimakhanov, Ainash Oshibayeva, Temirkhan Kozhakhmetov, Nazarbek Omarov, Dinara Akhmetzhanova, Berikuly Duman. RESULTS OF MEDICAL CARE FOR PERSONS WITH POLYTRAUMA IN ALMATY AND CORRECTION OF THE ORGANIZATIONAL APPROACH
Khatia Mikeladze, Nino Chikadze, Nino Gachechiladze, Marina Tediashvili, Irina Datikashvili-David, Peter Lydyard, Nina Porakishvili. SERUM IL-6, IL-12, AND IL-10 LEVELS IN EARLY-STAGE, UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA PATIENTS: INSIGHTSFROMGEORGIA
Musayeva H.H. FREQUENCY OF COMPLICATIONS IN PATIENTS WITH ADENTIA (BASED ON ARCHIVAL DATA)
Hong-Xia Wang, Xiao-Xia Hou, Jie Xu. NURSING RESEARCH ON EMERGENCY GASTROSCOPIC TREATMENT OF UPPER GASTROINTESTINAL FOREIGN BODIES
Tolegenova Z.Zh, Tokanova Sh.E, Baibussinova A.Zh, Kalikhanova K, Iskakova A.M, Shalgumbayeva G.M. ASSESSMENT OF INFECTIOUS DISEASE RISK FACTORS, INCLUDING COVID-19, AMONG HEALTHCARE WORKERS IN EAST KAZAKHSTAN REGION

Bassam A. Al- jabery, Majid R. Al-bahrani.	
ENVIRONMENTALLY SAFE CsPbBr3/MXene/MWCNTs HYBRID NANOCOMPOSITES: OPTOELECTRONIC AND STRUCTURAL	
CHARACTERISTICS FOR POSSIBLE BIOMEDICAL AND HEALTH APPLICATIONS	5-414
Hasan AlAidarous.	
PIGMENTED VILLONODULAR SYNOVITIS IN THE ANKLE OF A PEDIATRIC PATIENT: A CASE REPORT415	5-419
Kuat Zhussupov, Nazarbek Omarov, Sagit Imangazinov, Saule Imangazinova, Yernar Kairkhanov, Olga Tashtemirova, Rustem Kazangapov,	
Aldiyar Masalov, Darkhan Otkenov.	
ENDOSCOPIC INJECTION HEMOSTASIS AND LOCAL TREATMENT OF GASTRODUODENAL BLEEDING. LITERATURE REVIE	$\exists \mathbf{W}$
AND OWN DEVELOPMENTS420)-424

PIGMENTED VILLONODULAR SYNOVITIS IN THE ANKLE OF A PEDIATRIC PATIENT: A CASE REPORT

Hasan AlAidarous*.

College of Medicine, AlBaha University, Saudi Arabia

Abstract.

Background: Pigmented villonodular synovitis (PVNS), also known as diffuse tenosynovial giant cell tumor, is a rare, benign but aggressive synovial proliferative disorder. It most commonly affects the knee; involvement of the foot and ankle is uncommon (incidence ~1.8 per million), and pediatric cases are especially rare. This rarity often leads to delayed diagnosis, as early symptoms can mimic benign causes (e.g. sprains or "growing pains") and initial radiographs may be unremarkable.

Case Presentation: We report a case of a 12-year-old girl with a four-year history of chronic ankle pain and swelling. She had been misdiagnosed with growth-related pain or recurrent ankle sprains after undergoing multiple X-ray examinations that showed no abnormalities. Over time, a progressive ankle joint swelling and pain with restricted motion developed. On examination, she had tenderness over the ankle (medially and laterally) with limited range of motion; neurovascular status was intact. Laboratory work-up showed a normal white blood cell count and an elevated erythrocyte sedimentation rate (ESR of 30 mm/hour, normal <20), consistent with mild inflammation. Plain radiographs remained normal, but MRI revealed diffuse synovial thickening with confluent nodular masses in the anterior and posterior ankle joint recesses, with heterogeneous signals and minor bone erosion, suggestive of PVNS. Arthroscopic biopsy confirmed tenosynovial giant cell tumor (PVNS). The patient underwent an open synovectomy via combined anterior and posterolateral incisions (double incision) to achieve maximal synovial resection. Approximately six weeks post-operatively, a single-session adjuvant radiosynoviorthesis (RSO) was performed - an intra-articular injection of Yttrium-90 colloid (approximately 5 mCi) - to irradiate any remaining synovial tissue and reduce recurrence risk. Postoperatively, her pain improved with no immediate complications. At 12-month follow-up, the patient had no evidence of recurrence and demonstrated marked improvement: her ankle pain resolved (VAS pain score improved from 6/10 preoperatively to 0-1/10 post-treatment) and range of motion normalized (dorsiflexion 20° and plantarflexion 45°, restored from a preoperative ~50% limitation). She returned to full, painfree weight-bearing ambulation without assistive devices.

Conclusion: PVNS of the ankle is a challenging diagnosis in children due to its rarity and nonspecific presentation. This case highlights that persistent pediatric ankle pain with swelling, even with normal initial X-rays, should raise suspicion for uncommon pathologies like PVNS. Early MRI is warranted for definitive diagnosis. Complete surgical synovectomy remains the primary treatment for diffuse PVNS; however, adjuvant intra-articular RSO (radioisotope synovectomy) can be safely utilized to minimize the high recurrence risk associated with diffuse disease. Our patient's excellent outcome at one year

- with restored function and no recurrence - underscores the effectiveness of an aggressive combined approach in managing diffuse PVNS of the ankle in a pediatric patient.

Key words. Pigmented villonodular synovitis, tenosynovial giant cell tumor, ankle, pediatric, synovectomy, radiosynoviorthesis, radiation synovectomy, case report.

Introduction.

PVNS of the ankle is a very rare, locally aggressive proliferative synovial condition of unknown etiology [1-7]. The knee is the most commonly affected joint, while foot and ankle involvement is uncommon (estimated ~1.8 cases per 1,000,000 population) [2,8]. PVNS can occur in localized or diffuse forms; the diffuse type involves the entire synovium and tends to be more aggressive and destructive than the localized nodular form [9,10]. First described in 1941 by Jaffe et al., PVNS is benign but can cause severe joint erosions and bone destruction if untreated [11,12]. It typically affects adults aged 30–40, and pediatric cases are exceedingly uncommon [13].

Clinically, PVNS often presents with chronic joint pain, swelling, and stiffness. In children and adolescents, such symptoms are easily mistaken for more common issues. Indeed, in pediatric ankle pain, diagnoses like ankle sprain/strain or "growth pain" are often presumed, and a rare entity like PVNS may be initially overlooked in the differential diagnosis. Additionally, early PVNS may not show abnormalities on plain X-rays, which further contributes to diagnostic delay. In a recent multicenter study, the median delay from symptom onset to PVNS diagnosis was 18 months [14,15]. In our patient's case, diffuse ankle PVNS went undiagnosed for four years due to such factors.

Magnetic resonance imaging (MRI) is the most sensitive tool for early detection, often showing characteristic synovial proliferation with low to intermediate T1 and heterogeneous T2 signals with hemosiderin deposition [16,17]. Definitive diagnosis is confirmed by histopathology.

The mainstay of treatment for diffuse PVNS is complete synovectomy, either via arthroscopic or open surgery [18]. An open surgical approach often achieves a more extensive synovectomy (especially for diffuse or extra-articular disease) and is associated with lower recurrence rates in diffuse PVNS, albeit with more surgical morbidity [19]. Arthroscopic synovectomy, while minimally invasive, can be effective for accessible localized lesions but may risk higher recurrence if any diseased tissue remains. Recurrence rates after surgery alone are substantial – reported between ~40% up to 60% or higher for diffuse disease in some series. Therefore, adjuvant therapies are often considered. External beam radiotherapy and intra-articular radiosynoviorthesis (RSO) with beta-emitting radioisotopes (often Yttrium-90) have been used as

© GMN 415

adjuncts to surgery to reduce recurrence [3]. Yttrium-90 (^90Y) radiosynovectomy is commonly employed, especially in the United States, for large joints like the knee [3]. It delivers high-energy β -radiation (mean penetration $\sim 3-5$ mm) to ablate remaining synovium. For smaller joints (ankle, subtalar), some protocols use lower-energy isotopes (e.g. Rhenium-186 or Erbium-169) to minimize soft tissue damage. We present this pediatric ankle PVNS case to emphasize the importance of early recognition and to describe our surgical and adjuvant treatment approach, including open synovectomy and postoperative RSO, which led to a successful outcome.

Case Presentation.

A 12-year-old girl presented with a four-year history of right ankle pain and intermittent swelling, exacerbated by walking. Initially, her symptoms were mild and attributed to benign causes; over the years she underwent multiple clinical evaluations and X-ray examinations, which were all unremarkable, leading to diagnoses of ankle sprain/strain or "growth pains." However, her ankle swelling gradually and persistently increased. By the time of referral to our clinic, she reported daily ankle pain (intensity ~6/10 on walking), noticeable swelling, and reduced range of motion. There was no history of significant trauma and no family history of rheumatologic disease or tumors.

On examination, there was a diffuse swelling around the ankle joint, with tenderness along both the lateral and medial aspects. Ankle motion was limited (dorsiflexion and plantarflexion were painful and approximately half of normal range). There were no signs of infection; the overlying skin was normal and the joint was stable. Neurovascular status of the foot was intact. Laboratory tests showed a normal white blood cell count. However, the erythrocyte sedimentation rate (ESR) was markedly elevated at 30 mm/hour (normal <20 mm/hour), consistent with moderate inflammation; this is in line with literature reports that about 45–50% of PVNS patients can have an elevated ESR [1]. C-reactive protein was within normal limits.

Plain radiographs of the ankle (Figure 1) remained essentially normal, with no osseous erosions, spurs, or periarticular soft tissue calcifications. Given the persistent symptoms, an MRI was obtained. The MRI of the ankle (Figure 2) revealed diffuse synovial thickening and proliferation throughout the tibiotalar and subtalar joints. There were multiple confluent nodular synovial masses in the anterior and posterior recesses of the ankle. The lesions exhibited low-to-intermediate signal intensity on T1-weighted images and heterogeneous high signal intensity on proton density and T2-weighted images, with some foci of low signal on all sequences (consistent with hemosiderin deposition). A subtle cortical erosion was noted at the anterior aspect of the talar body. The flexor hallucis longus tendon sheath was involved by ill-defined lobulated synovial tissue with similar signal characteristics. Mild joint effusion was present in the ankle and subtalar joints, and a small amount of fluid in the retrocalcaneal bursa suggested mild bursitis.

These imaging findings were highly suggestive of diffuse PVNS (tenosynovial giant cell tumor). An ultrasound-guided synovial biopsy was performed on January 28, 2023, yielding grayish-brown synovial tissue. Histopathological examination confirmed features of pigmented villonodular synovitis (diffuse-



Figure 1. Ankle X-ray (AP view) showing no bony erosions or notable soft tissue swelling.



Figure 2. MRI of the right ankle: (A) Proton density fat-sat image showing synovial hypertrophy with nodular lesions (white arrows) in the anterior ankle joint; (B) T2-weighted image showing posterior ankle joint with low-signal deposits (arrowheads) consistent with hemosiderin within hypertrophic synovium (white arrows indicate extent of synovial proliferation).

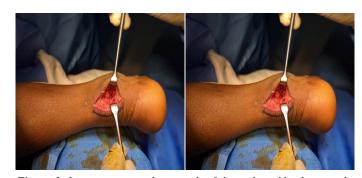


Figure 3. Intra-operative photograph of the right ankle showing the two incisions (anteromedial and posterolateral, marked by "*") used for open synovectomy in this diffuse PVNS case.

type tenosynovial giant cell tumor), with typical synovial villi, mononuclear cell proliferation, hemosiderin-laden macrophages, and multinucleated giant cells.

Given the diffuse involvement of the ankle joint, complete resection via a single incision was not feasible. On March 19, 2023, the patient underwent surgery consisting of an open synovectomy through two approaches (an anteromedial ankle incision and a posterolateral incision) to access and resect all affected synovial tissue. Intra-operatively, hypertrophic brownish synovium was observed in both anterior and posterior compartments; a meticulous open synovectomy (debridement) was performed, removing all visible diseased tissue (marginal resection). There were no immediate complications from surgery.

Post-operatively, the patient's ankle pain improved significantly. She was kept non-weight-bearing in a splint initially and started on gentle range-of-motion exercises in the first 2 weeks. At approximately 6 weeks after surgery, once the surgical wounds had healed, we administered adjuvant radiation synovectomy (RSO). Under fluoroscopic guidance, an intraarticular injection of a beta-emitting radioisotope was given to eradicate any residual microscopic synovium. Specifically, we utilized Yttrium-90 (^90Y) citrate colloid, which is commonly used for RSO in large joints [3]. A single dose of ~5 mCi (≈185 MBq) ^90Y was injected into the ankle joint (a dose on the lower end of the spectrum, adjusted for a smaller joint space). Triamcinolone acetonide (20 mg) was co-injected to mitigate post-injection inflammation. The patient's leg was immobilized for 48 hours post-RSO to allow the radionuclide to concentrate in the synovium. No immediate complications occurred; specifically, there were no signs of extra-articular leakage (confirmed by post-injection Bremsstrahlung imaging) and no skin or soft tissue reactions.

By the 6-week post-operative visit (just prior to RSO), the patient's ankle pain had already decreased (VAS 2–3/10 at that time) and her range of motion was improving with physical therapy. At final follow-up 12 months after surgery, she was doing well: the patient reported minimal to no pain in the ankle (VAS 0–1/10) and had regained full ankle motion compared to the contralateral side. She had returned to all normal activities and sports without limitations. Weight-bearing gait was normal, and no assistive devices were needed. Follow-up MRI at one year showed no evidence of PVNS recurrence.

Discussion.

Given an annual incidence of only ~1.8 per million in the United States [8], PVNS of the ankle is appropriately considered an orphan disease. Ankle involvement accounts for roughly 7-9% of PVNS cases [2]. This rarity, combined with the oftenindolent onset of symptoms, makes early clinical recognition challenging. Indeed, a multicentre review by Xie et al. noted an average ~18-month delay from symptom onset to diagnosis of PVNS [14]. In our patient, it took four years to arrive at the correct diagnosis. Initially, her complaints were attributed to benign causes (e.g. ankle sprain or growth pain), reflecting a common scenario where pediatric joint pain and swelling are presumed to be benign and self-limited. Only after symptoms persisted and progressed was advanced imaging obtained, revealing the true pathology. This case underscores the importance of considering rare diagnoses like PVNS in children with chronic monoarticular pain, especially when initial treatments for more common diagnoses fail. Furthermore, the lack of radiographic findings early in the disease (her initial X-rays were normal) likely contributed to the diagnostic delay, as PVNS-related bone erosions or calcifications typically appear only in later stages.

Clinically, diffuse PVNS of the ankle often presents with insidious ankle swelling, pain, and stiffness. Patients may or may not recall a precipitating injury. On exam, joint effusion or a boggy synovial mass may be palpable, and range of motion is often limited – consistent with our patient's presentation. Laboratory tests are usually non-specific; inflammatory markers can be normal or mildly elevated. Our patient's ESR was elevated (~30 mm/h), which is notable – a recent study found ESR elevation in 45.8% of PVNS cases, though typically only to a modest degree [1] (median ~13.5 mm/h in that series). Thus, while markedly high inflammatory markers are not the norm in PVNS, mild elevations of ESR/CRP can be present and should not exclude the diagnosis.

MRI is the imaging modality of choice. The pathognomonic MRI appearance of PVNS is due to hemosiderin deposition in the synovium: on T1-weighted images PVNS lesions often show low-to-intermediate signal, and on T2 or PD-fat sat sequences there is heterogeneous signal with blooming artifact from hemosiderin. Our case exhibited these classic features. Early in the disease, plain radiographs may appear normal, as in our patient. In more advanced cases, radiographs can show periarticular erosions with sclerotic margins, joint space narrowing, or soft tissue masses [12] – findings more often seen in weight-bearing joints like the hip, elbow, or ankle when PVNS has been long-standing.

The standard treatment for diffuse PVNS is complete synovectomy. Both arthroscopic and open techniques are employed. Open synovectomy generally allows more extensive clearance of diseased tissue (especially in hard-to-reach areas such as the ankle's posterior compartment or extra-articular tendon sheaths) and has been associated with lower recurrence rates in diffuse PVNS [19]. Arthroscopic synovectomy, on the other hand, offers quicker recovery and less stiffness, and is effective for localized disease or when lesions are accessible in a single compartment. According to the literature, recurrence rates after arthroscopic synovectomy for diffuse PVNS of large joints can range from about 40% up to 92%, whereas after open synovectomy they range roughly from 14% to 67%. (These wide ranges reflect varying follow-up durations and completeness of resection in different studies). In addition, the risk of osteoarthritis progression appears lower after thorough open synovectomy (0% in one series) compared to partial or arthroscopic approaches (reported 9-23%). In the ankle, complete excision is challenging due to the complex anatomy and confined joint space. A combined approach (arthroscopy for intra-articular lesions plus open surgery for extra-articular components) is sometimes recommended for extensive cases. In our patient, diffuse synovium extended throughout the ankle and into tendon sheaths, so we chose an open two-incision approach to maximally expose and resect all affected tissue. This "dual incision" open synovectomy allowed effective clearance of both anterior and posterior ankle compartments.

Even with optimal surgery, diffuse PVNS carries a substantial risk of local recurrence. Reported overall recurrence rates in the ankle without adjuvant therapy can be as high as ~30–50% [3]. Adjuvant therapies are therefore often considered, especially for diffuse disease. External beam radiotherapy has been used post-synovectomy in some cases and can reduce recurrence,

but concerns about radiation exposure and joint stiffness have limited its use. Chemical or radiosynovectomy is an alternative adjuvant aimed at selectively destroying residual synovium. RSO involves injecting a radioactive isotope (commonly ^90Y for large joints) into the joint space, where its β -radiation ablates remaining synovial cells with minimal systemic effect [3]. This technique has been used for decades in inflammatory arthritis and hemarthrosis, and has shown benefit in PVNS as well. Studies indicate that performing RSO after surgical synovectomy can significantly lower the recurrence rate of diffuse PVNS compared to surgery alone [5]. For example, without adjuvant RSO, diffuse PVNS recurrences may approach 40–50%, whereas combined surgery plus RSO has achieved recurrence rates on the order of ~0–20% in some reports [3]. (In other words, adjuvant RSO can roughly halve the recurrence risk, although results vary by study.) Our management aligns with this approach: we administered intra-articular ^90Y after surgery to eradicate residual disease and mitigate recurrence risk. It should be noted that while ^90Y is effective, its high radiation penetration can pose risks in smaller joints like the ankle [4]. Some authors recommend using a lower-energy isotope (such as Rhenium-186 or Erbium-169) for ankles to avoid complications like skin necrosis; indeed, cases of radionecrosis have been reported when high-dose ^90Y was used in ankle joints. In our case, we used a conservative ^90Y dose (~5 mCi) and encountered no adverse effects. The patient tolerated RSO well, and we observed no recurrence through one year of follow-up. Long-term vigilance is still required, as PVNS can recur even years later.

This case illustrates that diffuse PVNS of the ankle in a pediatric patient can be successfully managed with an aggressive combined-modality approach. Early recognition is crucial to prevent prolonged joint damage. For any child with chronic monoarticular joint swelling and normal radiographs, PVNS should be included in the differential diagnosis after ruling out infection or inflammatory arthritis. MRI should be employed early to identify synovial proliferative disorders. Once diagnosed, thorough synovectomy is paramount. Our preference is open excision for diffuse ankle PVNS to ensure complete resection. Adjuvant RSO, when available, is a valuable addition to reduce the likelihood of recurrence [5], provided it is used judiciously with appropriate isotope selection and dosing. The excellent functional recovery and absence of recurrence at 12 months in our young patient is encouraging, as PVNS in weight-bearing joints can often lead to significant morbidity. We plan to continue follow-up monitoring; if recurrence is detected, further intervention (repeat synovectomy and/or RSO) would be considered.

Conclusion.

Diffuse PVNS of the foot and ankle poses a complex clinical challenge, especially in pediatric patients where it can masquerade as benign conditions and lead to delayed diagnosis. This case underscores the importance of considering PVNS in the differential diagnosis of chronic ankle pain with swelling, even in children, and the utility of MRI in early detection. The preferred treatment is surgical synovectomy (open or combined open/arthroscopic for extensive disease) to achieve maximal resection. While arthroscopic excision minimizes surgical

morbidity, open synovectomy often yields more complete removal and thus lower recurrence in diffuse PVNS. Adjuvant intra-articular RSO can be safely and effectively employed after surgery to further reduce recurrence risk. In our patient, this multimodal approach led to resolution of symptoms, restoration of ankle function, and no recurrence to date. Long-term follow-up is essential, but the outcome suggests that even in pediatric diffuse PVNS of the ankle – a rare and difficult condition – a combination of timely diagnosis, aggressive surgical synovectomy, and adjunctive RSO can achieve an excellent result.

REFERENCES

- 1. Iakovou I, Symeonidis P, Kotrotsios D, et al. Radiosynoviorthesis after surgery in the treatment of patients with ankle pigmented villonodular synovitis: A case series. Journal of Clinical Medicine. 2020;9:597.
- 2. Mori H, Nabeshima Y, Mitani M, et al. Diffuse pigmented villonodular synovitis of the ankle with severe bony destruction: treatment of a case by surgical excision with limited arthrodesis. American Journal of Orthopedics. 2009;38:E187-E189.
- 3. Bisbinas I, De Silva U, Grimer R.J. Pigmented villonodular synovitis of the foot and ankle: A 12-year experience from a tertiary orthopedic oncology unit. Journal of Foot and Ankle Surgery. 2004;43:407-411.
- 4. Duncan N, Rajan R. Case report of pigmented villonodular synovitis arising from the calcaneocuboid joint in a 12-year-old male. The Foot. 2015;25:59-61.
- 5. Heller S.L, O'Loughlin P.F, Di Carlo G, et al. Pigmented villonodular synovitis about the ankle: two case reports. Foot & Ankle International. 2008;29:527-533.
- 6. Brien E.W, Sacoman D.M, Mirra J.M. Pigmented villonodular synovitis of the foot and ankle. Foot & Ankle International. 2004;25:908-913.
- 7. Korim M.T, Clarke D.R, Allen P.E, et al. Clinical and oncological outcomes after surgical excision of pigmented villonodular synovitis of the foot and ankle. Foot and Ankle Surgery. 2014;20:130-134.
- 8. Novikov D, Richardson M.W, Ho C, et al. A rare incidence of pigmented villonodular synovitis of the ankle in an adolescent. Journal of Foot and Ankle Surgery. 2018;57:1263-1266.
- 9. Xie G, Jiang N, Liang C, et al. Pigmented villonodular synovitis: a retrospective multicenter study of 237 cases. PLOS ONE. 2015;10:e0121451.
- 10. Schnirring-Judge M, Lin B. Pigmented villonodular synovitis of the ankle—radiation therapy as a primary treatment to reduce recurrence: a case report with 8-year follow-up. Journal of Foot and Ankle Surgery. 2011;50:108-116.
- 11. Cheng X.G, You Y.H, Liu W, et al. MRI features of pigmented villonodular synovitis (PVNS). Clinical Rheumatology. 2004;23:31-34.
- 12. Eckhardt B.P, Hernandez R.J. Pigmented villonodular synovitis: MR imaging in pediatric patients. Pediatric Radiology. 2004;34:943-947.
- 13. Karami M, Soleimani M, Shiari R. Pigmented villonodular synovitis in the pediatric population: literature review and a case report. Pediatric Rheumatology. 2018;16:14.

- 14. Bernthal N.M, Ishmael C.R, Burke Z.D. Management of pigmented villonodular synovitis (PVNS): an orthopedic surgeon's perspective. Current Oncology Reports. 2020;22:64.

 15. Gouin F, Noailles T. Localized and diffuse forms of tenosynovial giant cell tumor (formerly giant cell tumor of the tendon sheath and pigmented villonodular synovitis). Orthopaedics & Traumatology: Surgery & Research. 2017;103:S91-S97.
- 16. Lee M, Mahroof S, Pringle J, et al. Diffuse pigmented villonodular synovitis of the foot and ankle treated with surgery and radiotherapy. International Orthopaedics. 2005;29:403-405.
- 17. Dürr H.R, Capellen C.F, Klein A, et al. The effects of radiosynoviorthesis in pigmented villonodular synovitis of the knee. Archives of Orthopaedic and Trauma Surgery. 2019;139:623-627.
- 18. Quaresma M.B, Portela J, Soares do Brito J. Open versus arthroscopic surgery for diffuse tenosynovial giant cell tumors of the knee: a systematic review. EFORT Open Reviews. 2020;5:339-346.
- 19. Colman M.W, Ye J, Weiss K.R, et al. Does combined open and arthroscopic synovectomy for diffuse PVNS of the knee improve recurrence rates? Clinical Orthopaedics and Related Research. 2013;471:883-890.