

Case Report

Aseptic necrosis of the femoral and humeral heads in a patient who has received a course of cell therapy.

ABSTRACT)

Aims: The article is devoted to the problem of complications of pulse therapy and long-term use of corticosteroids in patients with multiple sclerosis.

Presentation of case: There was described a case of aseptic necrosis of the femoral and humeral heads in a patient suffering from multiple sclerosis after a course of cell therapy.

Discussion & Conclusion: This case is interesting not only because the patient got aseptic necrosis of the femoral as well as humeral heads but also due to possible role of stem cell therapy applied after intensive course of corticosteroids.

Keywords: aseptic necrosis, multiple sclerosis, femoral bone, humeral bone, stem cell therapy

1. INTRODUCTION

High-dose corticosteroids are widely used to treat acute relapses in patients suffering from multiple sclerosis [1-3]. Attempts to use pulse-therapy with methylprednisolone in the treatment of a secondary progressive multiple sclerosis are known [4]. Intravenous methylprednisolone pulse therapy (IVMP) inhibits the

inflammatory cytokine cascade, dampens T cell activation, facilitates apoptosis of activated immune cells, reduces extravasation of immune cells to the CNS, and decreases expression of class II histocompatibility antigens on antigen-presenting cells [1].

Pulse therapy is the administration of ultra-high doses of glucocorticoids for a short period. Methylprednisolone is the most commonly used medication, which in the form of sodium succinate is administered in a dose of 1-2 g intravenously in 30-60 minutes once a day for 3-5 days. The maximum concentration of the drug in the blood develops after 1 hour, followed by a decrease in 6-7 hours. Methylprednisolone accumulates in various tissues, and more in the inflammatory than normal (including in the brain), as well as in red blood cells. Given the characteristics of the distribution, minimal mineralocorticoid action, a weaker, compared with prednisone, effects on the gastrointestinal tract and the central nervous system, methylprednisolone is considered the drug of choice during pulse therapy [4].

Osteonecrosis (aseptic necrosis) of the tubular bone heads is one of the most severe complications of multiple sclerosis (MS) pulse therapy. It is known that every third case of non-traumatic osteonecrosis is associated with prolonged use of corticosteroids,

and, in turn, from 3-20% of patients receiving high doses of glucocorticoids are at risk of developing aseptic necrosis [5-7].

Currently, there is no clear understanding of what doses and what duration of therapy lead to osteonecrosis, however, compared with other nosoforms, the incidence of osteonecrosis of the heads of the tubular bones in MS remains low [5, 7, 8-12]. So, **Sahraian M.A. et al. (2012)** for 5 years of observation at the University Hospital of Tehran revealed only 5 cases of osteonecrosis after pulse therapy with methylprednisolone in a dose of 5 to 15 g per course [8]. Another study showed that in patients with MS, the frequency of osteonecrosis after pulse therapy is 15.5% [9]. Italian researchers consider the occurrence of osteonecrosis as a result of the influence of several factors: increased blood clotting, impaired lipid metabolism and fatty embolism of small-caliber vessels, increased peripheral vascular resistance, and activation of osteocytes apoptosis.

To date, Ukraine has no statistics on the prevalence of osteonecrosis in patients with multiple sclerosis. At the same time, the number of patients with multiple sclerosis has increased in recent years **[13]**, which may, under the conditions of limited use of disease modifying therapy lead to an increase in the number of cases of osteonecrosis after pulse therapy.

2. PRESENTATION OF CASE

This publication is devoted to the clinical case of aseptic necrosis of the heads of the humeral and femoral bones in patient B., born in 1982, who received repeated courses of pulse therapy with methylprednisolone in preparation for cell therapy in one of Moscow's clinics (Russia).

The patient has suffered from MS since 2003, when, after a stressful situation in the conditions of the maritime transition, there was a dysfunction of the pelvic organs, **manifestations** of central prosoparesis, lower paraparesis. The flow is steadily progressive with temporary spontaneous disturbance. According to MRI of the brain without contrast, in 2003, MS was diagnosed **and** courses of nootropic therapy were conducted. In 2007–2008, there was a restriction of movements **s** due to weakness in the legs (EDSS 5.0–6.0); in 2010, decrease in motor activity **was noted** (EDSS 6.0–6.5). Further deterioration occurred in 2011, the patient spent a long time in bed, sitting wheelchair, walking for short distances (EDSS 7.0). In 2012, she was examined and consulted at the National Medical and Surgical Center **n.a.** NI Pirogov (Moscow, Russia). She also passed a course of high-dosage immunosuppressive therapy (pulse therapy with **solumedrol** - 6000 mg per course) with the

support of autologous hematopoietic cells, a course of robotic kinesitherapy. For a long time, she received consolidation therapy with mitoxantrone and ondasetron as a support therapy. In the spring of 2014, after a regular course of kinesitherapy, pain occurred in the shoulder and hip joints. In May 2015, the diagnosis of aseptic necrosis of the heads of the humerus and femur was diagnosed. The patient received vazoprostan, denozumab (prolia), calcium supplements, vitamin D3 but her condition was not improved.

In October 2018, the patient passed the re-examination. At the time of the survey she complained of pain in the hip and shoulder joints, had restrictions on walking, numbness of the left leg, reduced visual acuity. Blood pressure was 135/80 mm Hg on the right hand, 130/80 mm Hg on the left hand. HR - 82 beats per minute.

On examination, the palpebral fissures were equal, the pupils were equal, photoreactions were alive, ophthalmodynamics was in full range, the adjusting nystagmus was present when looking to the right, weakness of convergence from two sides, more to the left. Muscle strength was reduced, more to the left (4 points). Tendon and periosteal reflexes in hands were raised without a clear difference of the parties. Positive reflexes of Jacobson-Laske, Zhukovsky and Wenderovich were positive in both sides, more

111 pronounced on the left. Knee reflexes were reduced without a clear
112 difference of sides, the Achilles reflex on the right was missing, on
113 the left it was reduced.

114 Gait was severely impaired, she had paraparesis, more manifested
115 on the left. Active and passive movements in the shoulder and hip
116 joints were limited - flexion in the shoulder 60-70°, right abduction -
117 80°, left - 60°. Pathological reflexes of Babinsky, Pussep,
118 Rossolimo were positive on both sides.

119 Coordinator tests were performed uncertainly from 2 sides, with
120 intention tremor. Decrease in sensitivity on the left in the Th9-Th10
121 innervation zone was found. There were signs of constant
122 incontinence. Meningeal signs were negative, no fasciculations
123 were detected. The patient was emotionally labile, asthenized, the
124 phenomenon of acrohyperhidrosis was determined.

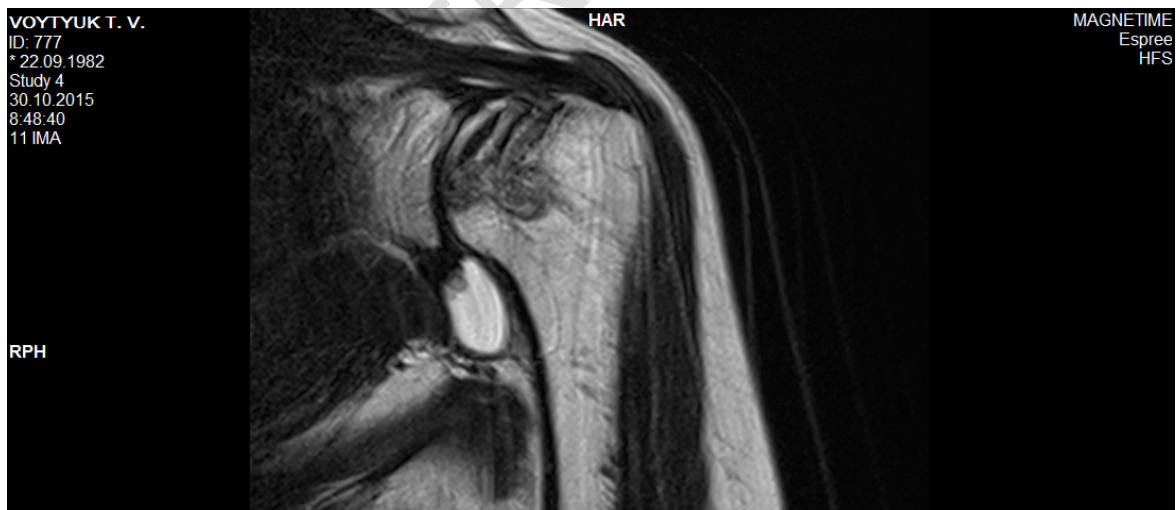
125 The MRI signs of pronounced avascular necrosis of the femoral
126 heads, the deformation of the left femoral head, an excess amount
127 of fluid in the joints, more to the left were determined (Fig. 1). In
128 addition, signs of avascular necrosis of the lateral femoral condyles
129 on both sides were identified. MRI signs of avascular necrosis of
130 the heads of both humerus bones and fluid in the joint cavity were
131 identified also.



a)



b)*



c)

Fig. 1 Manifestations of aseptic necrosis (a - heads of the femurs, b - lateral condyles of the femurs, c - heads of the humerus)

Multiple demyelination foci were defined in the brain, non-accumulating contrast, in the white matter of both hemispheres, in the legs of the brain, in the pons, in the medulla, in the corpus callosum, in the cerebellar hemispheres, in the cranial spinal cord, with a nominal diameter of 0.3 cm up to 2.8 cm, periventricular drain character. Cyst-like extensions of subarachnoid spaces in all areas of the brain, moderate expansion of the ventricular system, expansion of cerebellar sulci were determined. Cleavage of the posterior parts of the transparent septum was identified.

3. DISCUSSION

Considering the pronounced dysfunction and the absence of regress of symptoms, the patient was recommended surgery for prosthetic hip joints, however, in the current socio-economic conditions of Ukraine, this intervention could not be performed - waiting in line for a free prosthetic may take years. At the same time, the delay significantly worsens the prognosis and may lead to a further aggravation of the clinical picture.

This case is interesting not only because the patient got aseptic necrosis of the femoral as well as humeral heads but also due to possible role of stem cell therapy applied after intensive course of corticosteroids. There are no publications of such cases in the literature. However stem cell therapy was recommended for

164 treatment of avascular necrosis by some authors [11, 12]. Because
165 both high-dosage immunosuppressive therapy and other agents
166 influencing cellular immunity are used as a preparation to stem-
167 cells therapy than the consolidation therapy with mitoxantrone is
168 applied. However mitoxantrone itself could be a contributing factor
169 in medication-related osteonecrosis [14]. Also we still do not have
170 any data about the role of implanted stem cells in developing
171 avascular necrosis.

172 CONCLUSION

173 Avascular osteonecrosis remains rare complication of high dose
174 therapy with corticosteroids. Presented case shows outcomes of
175 several intensive courses of pulse-therapy provided for patient with
176 severe MS. It seems that we should avoid stem cell therapy with
177 preparatory high-dosage course of corticosteroids in the same year
178 with the previous relapse. Further investigations could help to
179 clarify if such approach is safe and efficient.

180 ACKNOWLEDGEMENTS

181
182 NONE TO DECLARE
183
184

COMPETING INTERESTS

Authors have declared that no competing interests exist

AUTHORS' CONTRIBUTIONS

All authors provide clinical guidance for the patient authors read and approved the final manuscript."

ETHICAL APPROVAL (WHERE EVER APPLICABLE)

The manuscript was approved by an Institutional Ethical Committee of the Center of Reconstructive & Renovative Medicine of Odessa National Medical University.

REFERENCES

1. Yamasaki R, Matsushita T, Fukazawa T, Yokoyama K, Fujihara K, Ogino M, Yokota T, Miyamoto K, Niino M, Nomura K, Tomioka R, Tanaka M, Kawachi I, Ohashi T, Kaida K, Matsui M, Nakatsuji Y, Ochi H, Fukaura H, Kanda T, Nagaishi A, Togo K, Mizusawa H, Murai H, Kira J. Efficacy of intravenous methylprednisolone pulse therapy in patients with multiple sclerosis and neuromyelitis optica. *Mult Scler*. 2016 Sep;22(10):1337-48.
2. Smets I, Van Deun L, Bohyn C, van Pesch V, Vanopdenbosch L, Dive D, Bissay V, Dubois B; Belgian Study Group for Multiple Sclerosis. Corticosteroids in the management of acute multiple sclerosis exacerbations. *Acta Neurol Belg*. 2017 Sep;117(3):623-633
3. Ratzer R, Iversen P, Börnsen L, Dyrby TB, Romme Christensen J, Ammitzbøll C, Madsen CG, Garde E, Lyksborg M, Andersen B, Hyldstrup L, Sørensen PS, Siebner HR, Sellebjerg F. Monthly oral methylprednisolone pulse treatment in progressive multiple sclerosis. *Mult Scler*. 2016 Jun;22(7):926-34
4. Liu S, Liu X, Chen S, Xiao Y, Zhuang W. Oral versus intravenous methylprednisolone for the treatment of multiple sclerosis relapses: A meta-analysis of randomized controlled trials. *PLoS One*. 2017 Nov 27;12(11):e0188644
5. Carulli C, Nistri L, Bracco L, Giannini M, Amato MP. A steroid-induced bilateral avascular necrosis of the femoral head in an underage patient affected by multiple sclerosis. *Clin Cases Miner Bone Metab*. 2015 Sep-Dec;12(3):257-9.
6. Kale N, Agaoglu J, Tanik O. Correlation of cumulative corticosteroid treatment with magnetic resonance imaging assessment of avascular femoral head necrosis in patients with multiple sclerosis. *Neurol Int*. 2010 Nov 26;2(2):e17.
7. Ce P, Gedizlioglu M, Gelal F, Coban P, Ozbek G. Avascular necrosis of the bones: an overlooked complication of pulse steroid treatment of multiple sclerosis. *Eur J Neurol*. 2006 Aug;13(8):857-61
8. Sahraian MA, Yadegari S, Azarpajouh R, Foroughipour M. Avascular necrosis of the femoral head in multiple sclerosis: report of five patients. *Neurol Sci*. 2012 Dec;33(6):1443-6
9. Andriolo L, Merli G, Tobar C, Altamura SA, Kon E, Filardo G. Regenerative therapies increase survivorship of avascular necrosis of the femoral head: a systematic review and meta-analysis. *Int Orthop*. 2018 Jul;42(7):1689-1704.
10. Franceschi F, Franceschetti E, Paciotti M, Torre G, Samuelsson K, Papalia R, Karlsson J, Denaro V. Surgical management of osteonecrosis of the humeral head: a systematic review. *Knee Surg Sports Traumatol Arthrosc*. 2017 Oct;25(10):3270-3278
11. Vijapura A, Levine HB, Donato M, Hartzband MA, Baker M, Klein GR. Total Hip Arthroplasty in Patients With Avascular Necrosis After Hematopoietic Stem Cell Transplantation. *Orthopedics*. 2018 Mar 1;41(2):e257-e261.

- 229 12. Toguchida J, Aoyama T, Goto K, Kakinoki R, Kasai Y. [Cell therapy for aseptic necrosis of femoral head]. Nihon
230 Rinsho. 2011 Dec;69(12):2225-30
- 231 13. Nefyodov O, Mamchur V. [Pharmacotherapy for multiple sclerosis - modern standards.] Pharmacology and drug
232 toxicology, № 3 (44) / 2015 10-16 [Ukr]
- 233 14. Bagan JV, Bagan L, Poveda R, Scully C. Mitoxantrone as a contributing factor in medication-related
234 osteonecrosis of the jaws. Int J Oral Maxillofac Surg. 2016 Mar;45(3):377-9

UNDER PEER REVIEW