

Case report

Radiation-induced intracranial vasculopathy after radiotherapy (laser surgery) for craniopharyngioma: a study case report

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Abstract

Brain irradiation is commonly used for many primary brain malignancies. After radiotherapy, the potential for the development of radiation-induced vasculopathy exists and should be kept in mind. Craniopharyngioma is one rare slowly developing, usually benign tumor. In the majority of cases, such tumors grow above the anterior upper lip of pituitary gland and are located mainly on Sella Turcica or on side sella areas. Macroscopically, they seem like big globe tumors with the white or red-blooded surface. Histologically, have ascertained two types, adamantinomatous and papillary type (though they have ascertained and mixed types). However, independently of their main type, they usually have cystic form. The size of craniopharyngioma is between 2 to 4 cm. Strike 0,12-2/100.000 people a year, 30 to 50% of all cases presenting during childhood and adolescence and they constitute the 2-5% of all brain tumors. The peak incidence rates have been shown in children of ages 5 to 14 years and adults of ages 50 to 74 years. The study below presents one case of post-radiation vasculopathy of the brain arteries of the patient who received radiation therapy to the brain as part of his craniopharyngioma management. It will be underscored the significance of this patient condition and also will be described possible routes to prevent the occurrence of vasculopathy.

Keywords: Cerebral vasculopathy, Radiation damage, Craniopharyngioma, Radiation therapy

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Introduction

Craniopharyngioma is a rare benign brain tumor arising along the craniopharyngeal duct and usually diagnosed during childhood or adult life with low histological grade (Hermann et al. 2014, Karavitaki et al. 2006). A plethora of patients are treated with both surgery and/or radiotherapy [Lo et al. 2016]. Radiotherapy is one of the primary treatments of brain tumors (Kralik et al. 2017). Although survival rates after removal of craniopharyngioma is 80-90% at 10 years by radioactive radiation (Habrand et al. 1999, Lo et al. 2014, Hetelekidis et al. 1993), patients with craniopharyngioma suffer potential adverse effects including hypopituitarism, visual problems, seizures, neurocognitive impairment, mood disorders (Varlotta et al. 2002, Laffond et al. 2012, Boop et al. 2012, Shiminski-Maher et al. 1990, Sterkenburg et al. 2015) and also radiation necrosis, atrophy, gliosis, telangiectasia, cavernous malformations (Wang et al. 2014). Cerebrovascular vasculopathy is also a potential adverse effect of cranial radiotherapy (Liu et al. 2009, Ayuk et al. 2012, Mueller et al. 2013, Ullrich et al. 2005). In the study of Brant-Zawadzki et al. was reported, when radiation-induced cerebrovascular vasculopathy occurs, histologically accompanying by the presence of subintimal foam cells and myointimal proliferation, fibrous thickening and hyalinization of the subintima, thickening of the elastica, fibrous thickening of the adventitia, and loose connective tissue thickening occluding the lumen (Brant-Zawadzki et al. 1980). It should also be noted that patients with a predisposition to insulin resistance, glucose intolerance, dyslipidemia, and hypertension have an increased risk of vasculopathy after

radiotherapy of craniopharyngioma (Srinivasan et al. 2004). In the proposed study we report a case of a vasculopathy caused by radiotherapy of craniopharyngioma to continue for 3 years because the area of craniopharyngioma's location is not surgically removable area. The craniopharyngioma detected for the first time by magnetic resonance imaging (MRI) due to often and intense headaches. The result of angiopathy became the automatic rupture of the vessels with the subsequent formation of subarachnoid bleeding and hematoma which together with the existing craniopharyngioma due to the pressure exerted on the basic brain structures, caused death because of cardiopulmonary failure. Thus, although radiotherapy carries a large number of degenerative changes in the surrounding highly functional areas, this method of treatment remains the best when it is not possible to perform a surgical intervention (Caldarelli et al. 2005).

Methods

The male patient was taken to the hospital with the following symptoms - often and intense headaches. Hereinafter was diagnosed with the craniopharyngioma in the pituitary area after the execution of the MRI. The maximum size of the craniopharyngioma was 2 cm. Subsequently, the patient was treated with laser for 3 years, because the craniopharyngioma was located in was not surgically removable area. The irradiation of the area caused postoperative metacentric vasculopathy and in combination that the patient suffered from hypertension, occurred the automatical rupture of the vessels and the formation of subarachnoid bleeding and hematoma.

Probably the bleeding started intraparenchymally, in the area where the tumor was previously located. Mild brain edema also was reported. Regarding the symptomatology, the patient was observed visual disorders and breathlessness, for around 3-4 days before his death. Eventually, the cause of death was a cardiopulmonary failure due to pressure on the basic brain structures.

Results

Notwithstanding craniopharyngioma was treated with radiation therapy, three years after it initiated, the patient developed vasculopathy, which led to the rupture of cerebral vessels with the subsequent bleeding. It is worth noting that vasculopathy rarely occurs as a consequence of radiation therapy. Only in two studies out of 7, on the basis of which a review of Bradley et al. was performed, vasculopathy was reported as a consequence of radiotherapy (Bradley et al. 2014). In the study of Luu et al. 16 patients who have treated craniopharyngioma with radiation therapy, only two developed vasculopathy which did not lead to death (Luu et al. 2006). Also in the study of Bishop et al. only two of 19 patients developed vasculopathy, which was not fatal (Bishop et al. 2013). Thus, death from vasculopathy was not reported in any case. Nevertheless, in the proposed study we report the vasculopathy is considered the primary cause of death, since subarachnoidal bleeding is a consequence of vasculopathy, and was caused by an additional pathology, hypertension, related to an increased risk of subarachnoidal bleeding (Feigin et al. 2005). Subarachnoidal bleeding, in turn, caused secondary consequences, such as pressure on the surrounding brain structures

responsible for the basic functions of the body, which was the ultimate cause of death.

Discussion

Craniopharyngioma is a tumor in the pituitary region with an incidence rate of 1.34 in all ages per 1 million (Nielsen et al. 2011). It represents approximately 2.5 – 3% of brain tumors in adults in the United States (Bunin et al. 1998; Karavitaki et al. 2006). Notwithstanding it is histologically benign, its infiltrative behavior affects the pituitary axis, the visual pathways, and other relevant vascular and neural structures and often has a long-term effects sequelae after surgical removal, that obstructs total removal of disease. Optimal therapeutic management of craniopharyngioma continues to be debated (Pietro Mortini et al. 2011). There are no commonly accepted guidelines or a clear consensus for best treatment's type of craniopharyngiomas in adults, although in some reports published have recommended radiation therapy after subtotal or partial removal of craniopharyngiomas, at least for adults (De Vile et al. 1996; Duff et al. 2000). Nonetheless, optimal treatment's type should always be determined considering, for example, the patient's symptoms, age, tumor localization, and extension.

Since the discovery of the effects of radiotherapy on cranial tumors, many investigations have been conducted with the patient's observation after treatment and possible outcomes, such as a vasculopathy. The syndrome of radiation-induced intracranial vasculopathy is accepted in the community of radio oncology. Vasculopathy is frequently not comprised of informed consents as a complication of radiotherapy to the brain for pediatric and young adult patients.

The first cases, published in the international bibliography, were reported in 1967 by Darmody et al. (Darmody et al. 1967) after brain irradiation for pituitary adenoma and Lee and Hodes (Lee and Hodes 1967) in consequence of irradiation for optic glioma. Ishibashi et al. reported a case of a 64-year-old woman, who had a chromophobe pituitary adenoma removed and followed by radiotherapy, six years after treatment she developed vasculopathy (Ishibashi et al. 1982). Turel et al., who evaluated retrospectively 42 adults' patients of which 34 had undergone radiation therapy, reported vasculopathy in one case with worsening of vision, as evidenced by MRI (Turel et al. 2016). Sawamura et al. reported vasculopathy of large intracranial arteries in 3 of 111 patients after irradiation over the parasellar region; two-thirds of these patients had a stroke, 2 and 14 years after irradiation, and one-third developed a huge dural arteriovenous malformation 11 years after (Sawamura et al. 1998).

Although there are many clinical cases of radiation-induced vasculopathy in the published literature, only a small part of them refers to the description of adult patients. The causes of vasculopathy can be implicated with various types of radiation such as Gamma Knife, colloidal gold intracystic instillation, fractionated external beam radiation therapy, and stereotactic radiosurgery. Moreover, it is obvious that any part of the brain may be exposed to the risk of developing vasculopathy. The time interval from the moment of radiation therapy until the onset of symptoms of vasculopathy ranges from 2 to 25 years. But in most cases, vasculopathy is manifested in the first 10 years (Aoki et al. 2002). Notwithstanding this disease has a long latency period,

the symptoms that manifest themselves are frequently acute. The course of the disease may have periodic improvements, but in general, it has a deteriorating pattern. Patients in many cases present stroke-like symptoms (Peñagaricano et al. 2004). Increased development of vasculopathy leads to systemic disorders that may involve motor impairment, sensory loss, ataxia, speech difficulty, or decreased cognitive ability. Due to the constantly recurring ischemic episodes caused by vasculopathy, there is a gradual deterioration in the functional status of patients.

In patients with vasculopathy, transdural collateral vascularization may be the main source of blood supply to the part of the brain where there is no damage to the brain vessels. So it is crucial to preserve this vascular anastomosis during the operation. Magnetic resonance imaging of patients suffering from vasculopathy, prior to the use of radiation therapy, can help in determining the sites of stenosis and obstruction in the blood vessels of the brain. The MRI pattern seen after the use of radiation therapy is similar to that expressed in vascular atherosclerosis. Thickening of the endothelium of the vascular walls of the brain can be observed on MRI and several years after the occurrence of occlusion or stenosis. In our case, the patient was treated only with radiation therapy. By dint of multimodal planning of radiotherapy with visual control, it is crucial to exclude large arteries of the brain that do not need treatment and can be injured. In contrast, MR studies and MR angiography can clearly define the boundaries of large arteries. Thus far, there are no proven methods that can reduce the possibility of vasculopathy after radiotherapy. The adjustment of time-dose-fractionation and volume

does not allow to avoid the occurrence of this complication. The administration of corticosteroids prior to and throughout the course of radiation therapy was proposed as a possible prevention option, but its effectiveness remains unproven. It has been reported that baby aspirin treatment is beneficial and may reverse some of the symptoms. Prevention by delaying radiation therapy in young patients for as long as possible remains the best method of treatment. If treatment still needs to be done, then to prevent radiation-induced vasculopathy, modern planning and delivery techniques with conformal coverage and conformal avoidance can be used as the preventive method.

References

- Aoki S, Hayashi N, Abe O, Shirouzu I, Ishigame K, Okubo T, et al. (2002). Radiation-induced arteritis: thickened wall with prominent enhancement on cranial MR images report of five cases and comparison with 18 cases of Moyamoya disease, *Radiology*, vol. 223, no. 3, pp. 683-8.
- Ayuk J. (2012). Does pituitary radiotherapy increase the risk of stroke and, if so, what preventative actions should be taken?, *Clin Endocrinol (Oxf.)*, vol. 76, no. 3, pp. 328-31.
- Bishop AJ, Mahajan A, Okcu MF, Allen PK, Kahalley LS, McAleer MF, McGovern SL, Grosshans DR. (2013). Proton Therapy for the Treatment of Childhood Craniopharyngiomas: Cyst Dynamics and Initial Outcomes, *American Radium Society 95th Annual Meeting*, P054.
- Boop FA. (2012). Craniopharyngioma, *J Neurosurg Pediatr*, vol. 10, no. 4, pp. 291-2.
- Bradley JA, Indelicato DJ. (2014). Craniopharyngioma and Proton Therapy, *Int J Particle Ther.*, vol. 1, no. 2, pp. 386-98.
- Brant-Zawadzki M, Anderson M, DeArmond SJ, Conley FK, Jahnke RW. (1980). Radiation-induced large intracranial vessel occlusive vasculopathy, *AJR Am J Roentgenol*, vol. 134, no. 1, pp. 51-5.
- Bunin GR, Surawicz TS, Witman PA, Preston-Martin S, Davis F, Bruner JM. (1998). The descriptive epidemiology of craniopharyngioma, *J Neurosurgery*, vol. 89, no. 4, pp. 547-51.
- Caldarelli M, Massimi L, Tamburrini G, Cappa M, Di Rocco C. (2005). Long-term results of the surgical treatment of craniopharyngioma: the experience at the Policlinico Gemelli, Catholic University, Rome, *Childs Nerv Syst*, vol. 21, no. 8-9, pp. 747-57.
- Darmody WR, Thomas LM, Gurdjian ES. (1967). Postirradiation vascular insufficiency syndrome, *Case report Neurology*, vol. 17, no. 12, pp. 1190-2.
- De Vile CJ, Grant DB, Kendall BE, Neville BG, Stanhope R, Watkins KE, et al. (1996). Management of childhood craniopharyngioma: can the morbidity of radical surgery be predicted?, *Journal of Neurosurgery*, vol. 85, no. 1, pp. 73-81.
- Duff JM, Meyer FB, Ilstrup DM, Laws ER, Schleck CD, Scheithauer BW. (2000). Long-term Outcomes for Surgically Resected Craniopharyngiomas, *Neurosurgery*, vol. 46, no. 2, pp. 291-305.
- Feigin VL, Rinkel GJ, Lawes CM, Algra A, Bennett DA, van Gijn J, et al. (2005). Risk factors for subarachnoid hemorrhage: an updated systematic

review of epidemiological studies, *Stroke*, vol. 36, no. 12, pp. 2773-80.

Habrand JL, Ganry O, Couanet D, Rouxel V, Levy-Piedbois C, Pierre-Kahn A, et al. (1999). The role of radiation therapy in the management of craniopharyngioma: a 25-year experience and review of the literature, *Int J Radiat Oncol Biol Phys*, vol. 44, no. 2, pp. 255-63.

Hetelekidis S, Barnes PD, Tao ML, Fischer EG, Schneider L, Scott RM, et al. (1993). 20-year experience in childhood craniopharyngioma, *Int J Radiat Oncol Biol Phys*, vol. 27, no. 2, pp. 189-95.

Ishibashi Y, Okada H, Mineura K, Kodama N. (1982). A case of radiation necrosis with vascular changes on main cerebral arteries (author's transl), *No Shinkei Geka*, vol. 10, no. 3, pp. 337-41. [Article in Japanese]

Karavitaki N, Cudlip S, Adams CDT, Wass JAH. (2006). Craniopharyngiomas, *Endocrine Reviews*, vol. 27, no.4, pp.371–97.

Kralik SF, Watson GA, Shih CS, Ho CY, Finke W, Buchsbaum J. (2017). Radiation-Induced Large Vessel Cerebral Vasculopathy in Pediatric Patients With Brain Tumors Treated With Proton Radiation Therapy, *Int J Radiat Oncol Biol Phys*, vol.99, no. 4, pp. 817-24.

Laffond C, Dellatolas G, Alapetite C, Puget S, Grill J, Habrand JL, et al. (2012). Quality of life, mood and executive functioning after childhood craniopharyngioma treated with surgery and proton beam therapy, *Brain Injury*, vol. 26, no. 3, pp. 270-81.

Lee KF, Hodes PJ. (1967). Intracranial ischemic lesions, *Radiologic Clinics of North America*, vol. 5, no. 3, pp. 363.

Liu AK, Bagrosky B, Fenton LZ, Gaspar LE, Handler MH, McNatt SA, et al. (2009). Vascular abnormalities in pediatric craniopharyngioma patients treated with radiation therapy, *Pediatr Blood Cancer*, vol. 52, no. 2, pp. 227-30.

Lo AC, Howard AF, Nichol A, Hasan H, Martin M, Heran M, et al. (2016). A Cross-Sectional Cohort Study of Cerebrovascular Disease and Late Effects After Radiation Therapy for Craniopharyngioma, *Pediatr Blood Cancer*, vol.63, no. 5, pp. 786-93.

Lo AC, Howard AF, Nichol A, Sidhu K, Abdulsatar F, Hasan H, et al. (2014). Long-term outcomes and complications in patients with craniopharyngioma: the British Columbia Cancer Agency experience, *Int J Radiat Oncol Biol Phys*, vol. 88, no. 5, pp. 1011-8.

Luu QT, Loredano LN, Archambeau JO, Yonemoto LT, Slater JM, Slater JD. (2006). Fractionated proton radiation treatment for pediatric craniopharyngioma: preliminary report, *Cancer J.*, vol. 12, pp. 155–9.

Mortini P, Losa M, Pozzobon G, Barzaghi R, Riva M, Acerno S, et al. (2011). Neurosurgical treatment of craniopharyngioma in adults and children: early and long-term results in a large case series, *Journal of Neurosurgery JNS*, vol. 114, no. 5, pp. 1350-9.

Müller HL. (2013). Childhood craniopharyngioma, *Pituitary*, vol. 16, no. 1, pp. 56-67.

- Müller HL. (2014). Craniopharyngioma, *Endocrine Reviews*, vol. 35, no. 3, pp. 513–43.
- Nielsen EH, Feldt-Rasmussen U, Poulsen L, Kristensen LO, Astrup J, Jørgensen JO, et al. (2011). Incidence of craniopharyngioma in Denmark (n = 189) and estimated world incidence of craniopharyngioma in children and adults, *J Neurooncol*, vol. 104, no. 3, pp. 755-63.
- Peñagaricano JA, Linskey ME, Ratanatharathorn V. (2004). Accelerated cerebral vasculopathy after radiation therapy to the brain, *Neurol India*, vol. 52, no. 4, pp. 482-6.
- Sawamura Y, Ikeda J, Shirato H, Tada M, Abe H. (1998). Germ cell tumours of the central nervous system: treatment consideration based on 111 cases and their long-term clinical outcomes, *Eur J Cancer*, vol. 34, no. 1, pp. 104-10.
- Shiminski-Maher T, Rosenberg M. (1990). Late effects associated with treatment of craniopharyngiomas in childhood, *J Neurosci Nurs*, vol. 22, no. 4, pp. 220-6.
- Srinivasan S, Ogle GD, Garnett SP, Briody JN, Lee JW, Cowell CT. (2004). Features of the Metabolic Syndrome after Childhood Craniopharyngioma, *The Journal of Clinical Endocrinology & Metabolism* vol. 89, no. 1, pp. 81–6.
- Sterkenburg AS, Hoffmann A, Gebhardt U, Warmuth-Metz M, Daubenbüchel AMM, Müller HL. (2015). Survival, hypothalamic obesity, and neuropsychological/psychosocial status after childhood-onset craniopharyngioma: newly reported long-term outcome, *Neuro-Oncology*, vol. 17, no. 7, pp. 1029–38.
- Turel MK, Tsermoulas G, Gonen L, Klironomos G, Almeida J, Zadeh G, et al. (2016). Management and outcome of recurrent adult craniopharyngiomas: an analysis of 42 cases with long-term follow-up, *Neurosurgical Focus FOC*, vol. 41, no. 6, pp. 11.
- Ullrich NJ, Scott RM, Pomeroy SL. (2005). Craniopharyngioma therapy: long-term effects on hypothalamic function, *Neurologist*, vol. 11, no. 1, pp. 55-60.
- Varlotto JM, Flickinger JC, Kondziolka D, Lunsford LD, Deutsch M. (2002). External beam irradiation of craniopharyngiomas: long-term analysis of tumor control and morbidity, *Int J Radiat Oncol Biol Phys*, vol. 54, no. 2, pp. 492-9.
- Wang L, Ni M, Jia W, Jia G, Du J, Li G, et al. (2014). Primary adult infradiaphragmatic craniopharyngiomas: clinical features, management, and outcomes in one Chinese institution, *World Neurosurg*, vol. 81, no. 5-6, pp. 773-82.