

BRIEF REPORT

Antiprogramed cell death-1 therapy with microspheres for metastatic liver tumors

Hiroteru Kamimura, ¹ Nobutaka Takeda, Takashi Owaki, Takeshi Mizusawa, Takahiro Iwasawa, Satoshi Ikarashi, Satoru Hashimoto, Masaaki Takamura and Shuji Terai

Division of Gastroenterology and Hepatology, Niigata University Graduate School of Medical and Dental Sciences, Niigata, Japan

Key words

anti-programmed cell death-1 therapy, melanoma, metastatic liver tumor.

Accepted for publication 12 May 2019.

Correspondence

Hiroteru Kamimura, Division of Gastroenterology and Hepatology, Niigata University Graduate School of Medical and Dental Sciences, 757 Asahimachi-dori 1, Chuo-ku, Niigata 951-8510, Japan. Email: hiroteruk@med.niigata-u.ac.jp

Declaration of conflict of interest: None.

Immunotherapy is an effective treatment for many cancers that require surgery, radiotherapy, and chemotherapy.¹ The use of antibodies that specifically block programed cell death-1 (PD-1) was approved for the treatment of melanoma in 2014.² Combining immunotherapy with other therapies, including locoregional therapy (e.g. transarterial embolization), is an innovative field of clinical investigation for cancer treatment. With the advent of immunotherapy may be best combined with locoregional therapies is increasing.³ Advanced melanoma presents a significant therapeutic challenge to clinicians. Many therapies for metastatic melanoma are limited by low response rates, severe toxicities, and/or relatively short response durations.⁴ Here, we report a case of partial remission and tumor regression after antiprogramed cell death-1 (anti-PD-1) therapy with microspheres.

A 67-year-old man presented to our division for additional treatment of multiple liver metastases. Six years ago, he was diagnosed with left uveal melanoma, and his eyeball was enucleated. Four years ago, liver metastasis was detected in S6, and segmental resection was performed. Pathological examination confirmed the diagnosis of delayed metastasis from the uveal melanoma. Two years ago, the liver tumor had grown multifocally (Fig. 1a); hence, anti-PD-1 therapy was initiated. However, the metastatic liver tumor was progressive; thus, we performed additional therapy with transarterial embolization using microspheres, sized 100–300 μ m, in the left lobe, selectively. After a 3-month follow-up, computed tomography demonstrated a necrotic cyst and partial remission of the tumor in the left lobe (Fig. 1b). The treatment effects on the tumors persisted even after 1 year. Anti-PD-1 therapy was continued basically but was stopped for 2 weeks during the embolization therapy. No major complications occurred. Selective hepatic artery embolization with microspheres is a safe treatment for patients with metastatic melanoma.

References

- 1 Okazaki T, Chikuma S, Iwai Y, Fagarasan S, Honjo T. A rheostat for immune responses: the unique properties of PD-1 and their advantages for clinical application. *Nat. Immunol.* 2013; 14: 1212–18.
- 2 Philips GK, Atkins M. Therapeutic uses of anti-PD-1 and anti-PD-L1 antibodies. *Int. Immunol.* 2015; **27**: 39–46.
- 3 Greten TF, Mauda-Havakuk M, Heinrich B, Korangy F, Wood BJ. Combined locoregional-immunotherapy for liver cancer. J. Hepatol. 2019; **70**: 999–1007.
- 4 Mamalis A, Garcha M, Jagdeo J. Targeting the PD-1 pathway: a promising future for the treatment of melanoma. *Arch. Dermatol. Res.* 2014; 306: 511–19.

JGH Open: An open access journal of gastroenterology and hepatology 3 (2019) 542–543

© 2019 The Authors. JGH Open: An open access journal of gastroenterology and hepatology published by Journal of Gastroenterology and Hepatology Foundation and John Wiley & Sons Australia, Ltd.

This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

542



Figure 1 (a) Computed tomographic angiogram showing a high-density area in the left lobe in computed tomography during left hepatic arteriography (CTLHA) in the second phase. (b) The second-phase CTLHA demonstrated a necrotic cyst and tumor regression of the tumor in the left lobe after a 3-month follow-up treatment.