

沙利度胺对晚期肺腺癌并恶液质综合征的疗效研究

《现代肿瘤医学》[ISSN:1672-4992/CN:61-1415/R] 期数: 2019年24期 页码: 4359-4362 栏目: 论著(胸部肿瘤) 出版日期: 2019-11-08

Title: The study of thalidomide in the treatment of advanced lung adenocarcinoma with CACS

作者: 王希方¹; 秦思达²; 任宏²; 白俊¹; 何莉¹; 王海鹏¹; 刘屹¹; 王维佳¹; 陆王锋³

1.陕西省人民医院肿瘤内科, 陕西 西安 710068;2.西安交通大学第一附属医院胸外科, 陕西 西安 710061;3.商洛市中心医院胃肠外科, 陕西 商洛 726000

Author(s): Wang Xifang¹; Qin Sida²; Ren Hong²; Bai Jun¹; He Li¹; Wang Haipeng¹; Liu Yi¹; Wang Weijia¹; Lu Wangfeng³

1.Department of Medical Oncology,Shaanxi Provincial People's Hospital,Shaanxi Xi'an 710068,China;2.Department of Thoracic Surgery,the First Affiliated Hospital,Xi'an Jiaotong University,Shaanxi Xi'an 710061,China;3.Department of Gastrointestinal Surgery,Shangluo Central Hospital,Shaanxi Shangluo 726000,China.

关键词: 晚期肺腺癌; 恶液质; 沙利度胺; 安全性; 疗效

Keywords: advanced lung adenocarcinoma; CACS; thalidomide; safe; effective

分类号: R734.2

DOI: 10.3969/j.issn.1672-4992.2019.24.008

文献标识码: A

摘要: 目的: 评价沙利度胺对晚期肺腺癌并CACS治疗的有效性和安全性。方法: 选取210例体重减轻10%以上的,确诊晚期肺腺癌并发恶液质的患者, 随机分为实验组和对照组。治疗组每晚服用沙利度胺100 mg, 维持治疗8周, 同时两组均给予营养支持, 主要评价标准是体重和营养状况的变化。结果: 用药8周后, 204例(对照组104例, 实验组100例)用于评估, 治疗组在治疗前后平均体重、食欲、ECOGPS评分、乏力的改善均具有统计学意义。便秘、过度镇静在两组间有统计学差异(P=0.002 和P=0.004), 其他症状均无统计学差异。结论: 沙利度胺对缓解晚期肺腺癌并发恶液质患者治疗有效, 且耐受良好。

Abstract: Objective: To evaluate the efficacy and safety of thalidomide in treatment of CACS of advanced lung adenocarcinoma. Methods: 210 patients whose weight loss over 10%, with advanced lung adenocarcinoma and CACS, were randomly divided into trail group and control group. The trail group was given a daily dose of thalidomide 100 mg for 8 weeks, while patients of two groups were given nutritional support. The main criteria were changes in weight and nutritional status. Results: 204 cases (control group 104 cases, the trail group 100 cases) used to evaluate after 8 weeks. The average body weight, appetite, ECOGPS score and the improvement of fatigue had statistical significance in the trail group after treatment. Constipation and somnolence had difference statistically between the two groups (P=0.002 and P=0.004 respectively), and other symptoms had no statistical difference. Conclusion: Thalidomide is effective and well tolerated in the treatment of patients with advanced lung adenocarcinoma complicated with CACS.

参考文献/REFERENCES

- [1] Koichi Takayama, Shinji Atagi, Fumio Imamura, et al. Quality of life and survival survey of cancer cachexia in advanced non-small cell lung cancer patients-Japan nutrition and QOL survey in patients with advanced non-small cell lung cancer study [J]. Support Care Cancer, 2016 (24) :3473-3480.
- [2] Damrauer JS, Stadler ME, Acharyya S, et al. Chemotherapy-induced muscle wasting: Association with NF- κ B and cancer cachexia [J]. European Journal of Translational Myology, 2018, 28(2): 7590.
- [3] Mattox, Todd W. Cancer cachexia. Cause, diagnosis, and treatment [J]. Nutrition in Clinical Practice, 2017, 32(5): 599-606.
- [4] Miyamoto Y, Hanna DL, Zhang W, et al. Molecular pathways: Cachexia signaling-A targeted approach to cancer treatment [J]. Clin Cancer Res, 2016 (22) :3999-4004.
- [5] Chukwuemeka Charles Ezeoke, John E Morley. Pathophysiology of anorexia in the cancer cachexia syndrome [J]. J Cachexia, Sarcopenia Muscle, 2015, 6(4): 287-302.

- [6] Fukawa T, Yanjiang BC, Minwen JC, et al. Excessive fatty acid oxidation induces muscle atrophy in cancer cachexia [J]. *Nature Medicine*, 2016, 22(6):666.
- [7] Sundaresan S, Puthanveetil P. Is FoxO1 the culprit, partner in crime, or a protector in systemic inflammation [J]? *American Journal of Physiology-Cell Physiology*, 2017:ajpcell.00194.2016.
- [8] Bye A, Wesseltoft-Rao N, Iversen PO, et al. Alterations in inflammatory biomarkers and energy intake in cancer cachexia: A prospective study in patients with inoperable pancreatic cancer [J]. *Med Oncol*, 2016 (33) :54.
- [9] Sulaieva O, Chereshneva Y, Kartashkina N, et al. Secretory function of white adipose tissue and adipokines: Biological effects and clinical significance [J]. *Georgian Medical News*, 2018(274):116.
- [10] Tara C Mueller, Jeannine Bachmann, Olga Prokopchuk, et al. Molecular pathways leading to loss of skeletal muscle mass in cancer cachexia can findings from animal models be translated to humans [J]. *BMC Cancer*, 2016 (16) :75.
- [11] Dong M, Lin J, Lim W, et al. Role of brown adipose tissue in metabolic syndrome, aging, and cancer cachexia [J]. *Frontiers of Medicine*, 2017.
- [12] GoKcen D, Serkan D, Erhan U, et al. Effects of serum leptin and resistin levels on cancer cachexia in patients with advanced-stage non-small cell lung cancer [J]. *Clinical Medicine Insights: Oncology*, 2017(11):117955491769014.
- [13] Fangyuan Z, Aomei S, Yinghui J, et al. The management strategies of cancer-associated anorexia: A critical appraisal of systematic reviews [J]. *BMC Complementary and Alternative Medicine*, 2018, 18(1):236.
- [14] Chen JF, Xiao LF, Zhang R, et al. Bortezomib, thalidomide, and dexamethasone (VTD) induction results in better overall survival than adriamycin, thalidomide, and dexamethasone (ATD) induction in previously untreated myeloma patients eligible for transplants [J]. *Acta Haematologica*, 2017, 137(4):207.
- [15] Han J, Meng Q, Lei S, et al. Interleukin-6 induces fat loss in cancer cachexia by promoting white adipose tissue lipolysis and browning [J]. *Lipids in Health & Disease*, 2018, 17(1):14.
- [16] Alejandro Schcolnik-Cabrera, Alma Chávez-Blanco, Guadalupe Domínguez-Gómez, et al. Understanding tumor anabolism and patient catabolism in cancer-associated cachexia [J]. *Am J Cancer Res*, 2017, 7(5):1107-1135.
- [17] Sadeghi M, Keshavarz-Fathi M, Baracos V, et al. Cancer cachexia: Diagnosis, assessment, and treatment [J]. *Critical Reviews in Oncology/hematology*, 2018, 127(8):91-104.
- [18] Brigo J, Elorza AA, Riedel CA, et al. Role of oxidative stress as key regulator of muscle wasting during cachexia [J]. *Oxidative Medicine and Cellular Longevity*, 2018, 2018(8):2063179.
- [19] Ou YC, Li JR, Wang JD, et al. Aspirin restores ABT-737-mediated apoptosis in human renal carcinoma cells [J]. *Science of the Total Environment*, 2018, 502(2):187-193.

备注/Memo: National Natural Science Foundation of China(No.81402506);国家自然科学基金资助项目 (编号: 81402506)

更新日期/Last Update: 1900-01-01