

Beau's Lines Resulting from Taxane Chemotherapy

Kumpol Aiempanakit, M.D.

Division of Dermatology, Department of Internal Medicine, Faculty of Medicine, Prince of Songkla University, Hat Yai, Songkhla 90110, Thailand.

Received 13 June 2018 • Revised 16 August 2018 • Accepted 27 August 2018 • Published online 10 October 2018

Abstract:

Nail abnormalities are frequently found in oncologic patients who have undergone chemotherapy. Although these changes do not require treatment, they could influence the treatment plan and the patient's quality of life. Some nail disorders lead to severe complications. Herein, the author reports on a patient with advanced breast cancer who received multiple kinds and cycles of chemotherapy. She developed multiple, parallel, transverse grooves, compatible with Beau's lines, on the nail plate of all her fingernails and toenails. This report aims to further the knowledge of medical students, physicians, and healthcare providers regarding the nails of patients who receive chemotherapy.

Keywords: adverse drug reaction, Beau's lines, chemotherapy, nails, taxane

Contact: Kumpol Aiempanakit, M.D.
Division of Dermatology, Department of Internal Medicine, Prince of Songkla
University, Hat Yai, Songkhla 90110, Thailand.
Email: akumpol@medicine.psu.ac.th

J Health Sci Med Res 2018;36(4):307-310
DOI: <http://dx.doi.org/10.31584/jhsmr.201826>
www.jhsmr.org

Introduction

Nails, a special skin appendage, consist of complex structures, including the nail matrix, the nail bed, the proximal nail fold, the hyponychium, and a hard keratin product, the nail plate. Nails have multiple functions, for example, supporting the ability to pick up small objects, protecting the distal fingers and toes, providing information about personal health and specific exposures, and influencing mental status.¹ Fingernail growth rate is 3–3.5 mm/month in adults, and grow two times faster than toenails, approximately 1–1.5 mm/month.^{1,2} The growth rate can be altered by some physiological or pathological factors and medications.^{2,3}

Nail abnormalities can be classified by the primary site of the structural damage, including the nail matrix (Beau's line, onychomadesis, pitting, true leukonychia, koilonychia), nail bed (onycholysis, apparent leukonychia, splinter hemorrhages), and nail fold (paronychia, pyogenic granuloma).¹ Chemotherapy, particularly the taxane regimen, can temporarily arrest nail matrix proliferation, causing nail plate abnormalities. It can also induce inflammation and pigmentation of the nail apparatus. This article reports on Beau's lines, an interesting nail abnormality resulting from taxanes chemotherapy, which correlates with the cycles of chemotherapy and the nail growth rate.

Case report

A 42-year-old woman presented with nail changes for a year. Fifteen months earlier, she was diagnosed with breast cancer, which metastasized to the liver, lungs, and pleura. The tumors were positive for estrogen and progesterone receptor, but negative for human epidermal growth factor receptor 2. She had received 4 courses of a combination of doxorubicin and cyclophosphamide, and then switched to paclitaxel 175 mg/mm² for 7 sessions, but

her disease sustained only a partial response. Her most recent chemotherapy regimen was 100 mg/mm² of docetaxel every 4 weeks/cycle for 5 cycles. She noticed the first nail changes occurring a month after receiving the first cycle of docetaxel.

Physical examination showed multiple, parallel, constant transverse grooves on the nail plate of all her fingernails (Figure 1) and toenails (Figure 2), compatible with Beau's lines, which correlated with the timing of her taxane chemotherapy cycles. She had no history of onychomadesis. Her nails also showed melanonychia and distal onycholysis.



Figure 1 Multiple parallel Beau's lines, melanonychia, and distal onycholysis of all fingernails



Figure 2 Multiple parallel Beau's lines, melanonychia, and distal onycholysis of all toenails

Discussion

Beau's lines, first described by French physician Dr. Joseph Honoré Simon Beau in 1846, are transverse depressions of the nail plate resulting from a temporary arrest of mitotic activity of the proximal nail matrix.⁴ This transverse line is a distal migration following the nail growth rate. The multiple parallel lines, as shown in this case, indicate repeated normal damage from the constant course of chemotherapy. The other conditions causing multiple Beau's lines are systemic, including viral infection, especially hand-foot-and-mouth disease, high fever, and localized Beau's lines caused by trauma, infections, or inflammation of the nails, e.g., manicures, chronic paronychia, and eczema.^{1,5,6} Severe damage can produce a complete interruption of proximal and distal nail matrix activity, which causes a complete detachment of the distal nail plate from the proximal nail fold, and is called onychomadesis or *defluvium unguium*.⁵

Taxanes (e.g., paclitaxel, docetaxel) are potent cytotoxic chemotherapy agents used in the treatment of many kinds of solid neoplasms, including breast, lung,

gastric, ovary, prostate, and head and neck cancers.⁷ The all-grade incidence of nail changes with paclitaxel treatment is reported to be approximately 43.7%.⁸ These nail adverse reactions can be hyperpigmentation, melanonychia, splinter hemorrhages, onycholysis, photo-onycholysis, Meuhrcke's nails, acute paronychia, pyogenic granuloma, Beau's lines, and onychomadesis.^{1,7,8} The patient in this report had melanonychia, distal onycholysis, and Beau's lines after treatment with paclitaxel and docetaxel.

Nail changes can commonly occur due to various medications other than taxanes (Table 1). Medications frequently reported with melanonychia are hydroxyurea, zidovudine, and psoralen.^{1,9,10} For onycholysis, other medications are tetracyclines, psoralen, and nonsteroidal anti-inflammatory drugs.¹ Retinoids, indinavir, methotrexate, capecitabine, sirolimus, and epidermal growth factor receptor inhibitors reported paronychia and pyogenic granuloma.^{1,11} However, another cause of nail changes, including infections and cutaneous metastasis, should be considered in patients who have advanced cancer with immunocompromised status.¹²

Table 1 Drug-induced nail abnormalities^{1,7-11}

Nail changes	Affected anatomy	Common medications
Beau's lines, onychomadesis	Nail matrix	Taxanes
Melanonychia	Nail matrix	Taxanes, hydroxyurea, fluorouracil, zidovudine, psoralen
True leukonychia (e.g. Mee's lines)	Nail matrix	Doxorubicin, vincristine, fluorouracil, methotrexate, cyclophosphamide
Apparent leukonychia (e.g. Muehrcke's lines)	Nail bed	Taxanes
Onycholysis	Nail bed	Taxanes, hydroxyurea, capecitabine, cyclophosphamide, doxorubicin, fluorouracil, tetracyclines, psoralen, nonsteroidal anti-inflammatory drugs
Paronychia and periungual pyogenic granuloma	Nail fold	Taxanes, methotrexate, capecitabine, sirolimus, epidermal growth factor receptor inhibitors, retinoids, indinavir

There is a lack of guidelines and effective treatment interventions for nail abnormalities associated with chemotherapy.⁸ Most nail changes are asymptomatic and do not require treatment. Local wound care and topical antibiotics and corticosteroids should be adjusted in patients with symptoms. In severe reactions, a reduced dose of chemotherapy might help prevent events afterward.

Conclusion

Nail changes resulting from chemotherapeutic agents are common. Patients should get information before treatment plans, and physicians should also evaluate adverse reactions. When patients experience nail abnormalities, they should receive an appropriate intervention.

Acknowledgment

The author is grateful to the International Affairs Office, Faculty of Medicine, Prince of Songkla University for language editing services. This case report was approved by the Research Ethics Committee, Faculty of Medicine, Prince of Songkla University.

References

1. Tosti A, Piraccini BM. Nail disorders. In: Bologna JL, Schaffer JV, Cerroni L, editors. *Dermatology*. 4 ed. China: Elsevier; 2018; p.1203–19.
2. Aiempanakit K, Geater A, Limtong P, Nicoletti K. The use of topical minoxidil to accelerate nail growth: a pilot study. *Int J Dermatol* 2017;56:788–91.
3. Geyer AS, Onumah N, Uyttendaele H, Scher RK. Modulation of linear nail growth to treat diseases of the nail. *J Am Acad Dermatol* 2004;50:229–34.
4. Weismann K. J.H.S. Beau and his descriptions of transverse depressions on nails. *Br J Dermatol* 1977;97:571–2.
5. Braswell MA, Daniel CR 3rd, Brodell RT. Beau lines, onychomadesis, and retronychia: a unifying hypothesis. *J Am Acad Dermatol* 2015;73:849–55.
6. Jacobsen L, Zimmerman S, Lohr J. Nail findings in hand–foot–and–mouth disease. *Pediatr Infect Dis J* 2015;34:449–50.
7. Sibaud V, Leboeuf NR, Roche H, Belum VR, Gladieff L, Deslandres M, et al. Dermatological adverse events with taxane chemotherapy. *Eur J Dermatol* 2016;26:427–43.
8. Capriotti K, Capriotti JA, Lessin S, Wu S, Goldfarb S, Belum VR, et al. The risk of nail changes with taxane chemotherapy: a systematic review of the literature and meta-analysis. *Br J Dermatol* 2015;173:842–5.
9. Kaewdech A, Aiempanakit K, Apinatriyo B. Acral hyperpigmentation resulting from hydroxyurea therapy in primary myelofibrosis. *Indian J Hematol Blood Transfus* 2018;34:551–2.
10. Aiempanakit K. Cutaneous hyperpigmentation in general practice. *Songkla Med J* 2015;33:165–75.
11. Fox LP. Nail toxicity associated with epidermal growth factor receptor inhibitor therapy. *J Am Acad Dermatol* 2007;56:460–5.
12. Aiempanakit K. Digital metastasis of tongue squamous cell carcinoma. *JAAD Case Rep* 2018;4:200–2.