

## 罕见白癜风并发环状扁平苔藓1例报告

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**【摘要】** 患者女, 36岁, 因“腰部白斑10<sup>+</sup>年, 其上新发环状斑块6<sup>+</sup>月”来院就诊。入院后完善伍德灯检查示: 阳性。右腰部环状隆起边缘处皮肤组织病理示: 表皮角化过度, 颗粒层楔形增厚, 基底细胞液化变性, 真皮浅层淋巴细胞呈带状浸润。诊断: 白癜风并发环状扁平苔藓。予以卤米松乳膏治疗, 2次/d, 治疗1个月后紫红色环状斑块基本消退, 随访2个月未见复发。环状扁平苔藓是扁平苔藓的一种少见亚型, 迄今国内外尚未见到此病合并白癜风的病例报道。

**【关键词】** 环状扁平苔藓 扁平苔藓 白癜风

**Vitiligo Combined With Annular Lichen Planus: A Rare Case Report** ZHOU Hai-yan<sup>1</sup>, LUO Dan<sup>2</sup>, PU Xiao-lan<sup>1Δ</sup>.

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**【Abstract】** A 36-year-old female patient came to our hospital with the chief complaint of having white patches on her waist for 10-plus years and having new annular plaques appearing on the white patches for 6-plus months. Wood's lamp examination done in the hospital showed a positive result. Histopathology of skin tissue from the edge of the annular swelling on the right waist revealed epidermal hyperkeratosis, wedge-shaped thickening of the granular layer, liquefactive degeneration of the basal cells, and a band-like infiltration of lymphocytes in the superficial dermis. The patient was diagnosed with vitiligo combined with annular lichen planus (ALP). The patient was treated with topical halometasone cream administered twice a day. The purplish-red annular plaques subsided and disappeared almost completely one month after the treatment was started and no signs of recurrence were observed during the 2-month follow-up. ALP is a rare variant of lichen planus. There has been no reported case of vitiligo combined with ALP so far.

**【Key words】** Annular lichen planus Lichen planus Vitiligo

### 1 病例资料

患者, 女性, 36岁, 因“腰部白斑10<sup>+</sup>年, 其上新发环状斑块6<sup>+</sup>月”来院就诊。患者入院前10<sup>+</sup>年无明显诱因于双侧腰部出现直径约4~6 cm白色斑片, 边界清晰、不规则, 无鳞屑, 不伴疼痛、瘙痒, 因白斑进展缓慢, 患者未重视, 未治疗。6<sup>+</sup>月前患者无明显诱因于双侧腰部白斑上及内踝部出现直径约1~3 cm紫红色环状斑块, 环状边缘堤状隆起, 其上可见少许白色鳞屑, 偶有轻微瘙痒, 无水疱、糜烂、渗出等。既往予以“萘替芬酮康唑乳膏”“他克莫司软膏”治疗, 无明显好转。为求进一步诊治, 遂至我院就诊。内科查体无特殊。皮肤科查体: 双侧腰部可见直径约4~6 cm白色斑片, 其上及内踝部可见直径约1~3 cm紫红色环状斑块, 环状边缘堤状隆起, 其上可见少许白色鳞屑(图1)。口腔黏膜、指甲未查见明显异常。既往史及家族史: 饮食无特殊, 近期无用药史、疫苗注射史、化学品接触史。实验室检查: 血常规、肝肾功能、抗核抗体、抗可提取性核抗原(extractable nuclear antigen, ENA)抗体谱结果未见异常。皮肤真菌荧光镜检: 阴性。伍德灯

检查阳性: 左侧腰部亮白色斑片, 边界清晰, 部分白斑可见毛囊复色(图2)。组织病理(右腰部环状隆起边缘): 表皮角化过度, 颗粒层楔形增厚, 基底细胞液化变性, 真皮浅层淋巴细胞呈带状浸润。结合患者典型的皮损特征、皮肤真菌荧光镜检、伍德灯检查及组织病理学结果, 诊断为白癜风合并环状扁平苔藓(annular lichen planus, ALP)。予以卤米松乳膏治疗, 2次/d, 治疗1个月后紫红色斑块基本消退, 随访2个月未见复发。但白斑尚未见明显好转, 目前患者仍在随访中。

### 2 讨论

ALP和白癜风是两个独立的疾病, 目前病因不明。ALP是扁平苔藓的一种少见亚型, 约占扁平苔藓的10%<sup>[1-2]</sup>。目前研究表明, ALP可能与细胞介导的免疫反应、遗传因素、感染因素、神经精神因素、药物因素、细胞内某些蛋白的改变及合并其他疾病都有着密不可分的关系<sup>[3]</sup>。临床上环状皮损主要表现为两种类型: ①多个小丘疹排列呈环状, 多为单环, 少数可呈多环; ②单个丘疹或斑块离心性向外扩展, 中央消退, 可伴有中央萎缩或色素沉着<sup>[4]</sup>。ALP好发于唇、龟头及阴茎, 腋窝、腹股沟和四肢也可发

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图 1 临床照片

Fig 1 Clinical pictures

A: left lumbar region; B: right lumbar region; C: ankle.

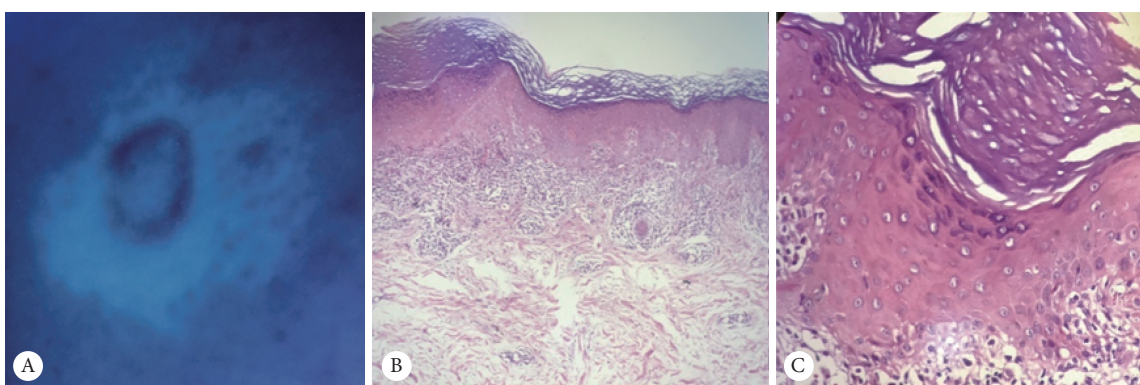


图 2 伍德灯检查阳性 (A) 和病理HE染色 (B: ×40; C: ×200)

Fig 2 Positive result from Wood's lamp examination (A) and pathological HE staining (B, ×40; C, ×200)

生,多不累及头皮、黏膜及甲<sup>[1-2]</sup>。患者通常没有自觉症状,但偶有瘙痒或灼热感。本例患者的皮损基本符合上述特点。白癜风是一种获得性、特发性色素脱失性皮肤病,发病机制主要涉及遗传、免疫、氧化应激、黑素细胞自毁等<sup>[5]</sup>。最新对T细胞介导的免疫应答机制的研究表明,T淋巴细胞在扁平苔藓和白癜风的免疫发病机制中均具有重要作用,辅助性T淋巴细胞(Th)1、Th9、Th17、Th22和细胞毒性T淋巴细胞是二者免疫发病机制的共同组成部分<sup>[6]</sup>。

通过文献检索,迄今国内外尚未见到ALP合并白癜风的病例报道。目前已有研究报道环状扁平苔藓可合并银屑病关节炎、红斑狼疮、生殖器黑子(生殖器黏膜黑斑)、Sneddon综合征、克罗恩病、人类免疫缺陷病毒感染、新型冠状病毒(SARS-CoV-2)感染等<sup>[7-13]</sup>。此外,也有一些文献报道了白癜风合并扁平苔藓的情况,二者皮损可以重叠,也可分别发生,但报道中扁平苔藓皮损主要表现为圆形、椭圆形或多角形紫红色扁平丘疹、斑块<sup>[14-20]</sup>。本例患者ALP皮损多发,位于双侧腰部及内踝部,初起均呈环状,且绝大部分ALP皮损与白癜风皮损为重叠发生,本例患者在原有的白癜风皮损基础上出现ALP皮损,进

一步证实了ALP和白癜风可能存在共同的发病机制。

本例患者的ALP皮损对于局部外用糖皮质激素治疗敏感,疗效显著,但白癜风色素脱失尚未见明显恢复,仍在进一步随访中。ALP有时与环状肉芽肿、环状银屑病、持久性色素异常性红斑、皮肤癣菌感染、蕈样肉芽肿和二期梅毒等难以区分,但本例患者通过实验室检查、组织病理学检查及对糖皮质激素治疗反应可明确诊断。

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**利益冲突** 所有作者均声明不存在利益冲突

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