Azathioprine-induced pellagra in a child with autoimmune hepatitis: A case report and literature review

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Abstract
Pellagra is a clinical syndrome resulting from niacin deficiency with variety of manifestations. Azathioprine is among drugs that can lead to such condition. Physicians should be aware as proper management can lead to full resolution.

KEYWORDS
autoimmune hepatitis, azathioprine, drug eruptions, pellagra, vitamin B deficiency

1 | CASE PRESENTATION

Pellagra, a potentially fatal but easily treated disorder, is characterized by symmetrical photo-distributed skin lesions due to niacin deficiency. There are many known etiologies for pellagra including drug-induced cases. We, hereby, present a new case of azathioprine-induced pellagra and briefly review reported cases of azathioprine-induced pellagra in the literature. A 14-year-old girl was admitted with a 3-day history of abdominal pain and diarrhea. She had a recent diagnosis of autoimmune hepatitis 8 months ago and azathioprine (AZA) was initiated 4 weeks before. She was underweight (body mass index of 17.7 kg/m²). Further history revealed poor nutritional intake due to decreased appetite, but she denied any specific dietary restrictions. The patient also complained of a 1-day history of new onset painful skin lesions. She had no personal or family history of skin diseases. Well-demarcated violaceous to brown skin lesions with scaly desquamation and hyperpigmentation were present on the neck, dorsum of the hands, and feet. The patient reported that the rash developed shortly after having sun exposure in the preceding days (Figures 1 and 2).

Based on the photo distribution of dermatitis and associated scaling and hyperpigmentation in the pattern of casal’s necklace in association with the recent initiation of AZA, a clinical diagnosis of AZA-induced pellagra was made. Our patient’s total Naranjo Scale score was 6 (possible adverse drug reaction).

Histopathological examination showed confluent parakeratosis, mild acanthosis, and regular elongation of rete ridges. There was focal subepidermal blistering with dermal capillary proliferation and a mild perivascular infiltrate (Figure 3).

Discontinuation of azathioprine and administration of oral nicotinamide 50 mg twice daily resulted in significant improvement of the rash within 2 days. The dosage of nicotinamide was gradually increased over the following 4 weeks. After 2 months of treatment, only mild postinflammatory hyperpigmentation remained.
2 | DISCUSSION

Pellagra is a clinical syndrome resulting from niacin deficiency. It is an underdiagnosed but still existing disease. Pellagra is clinically characterized by the classic triad of 3 D's: (1) dermatitis (photo-distributed symmetrical skin lesions), (2) diarrhea, and (3) dementia (neurologic and psychologic disturbances) which can ultimately lead to death if left undiagnosed or untreated.⁴

Histological features of pellagra are perivascular lymphocytic infiltrate in the upper dermis and edema in the papillary dermis which can be observed in the acute stages. Hyperkeratosis, parakeratosis, and epidermal atrophy are mostly seen in late stages. Although these features are unspecific, they can support the clinical diagnosis.²

The main cause of pellagra is niacin or tryptophan (niacin precursor) deficiency. Pellagra has been reported to be associated with some medications including isoniazid, 6-mercaptopurine (6-MP), 5-fluorouracil, and also azathioprine (AZA).¹

AZA is metabolized to 6-MP. 6-MP decreases the synthesis of nicotinamide adenine dinucleotide and nicotinamide adenine dinucleotide phosphate, which are key coenzymes in niacin metabolism and other metabolic pathways. Consequently, lack of these coenzymes will result in major dysfunction in tissues with high energy demands such as brain, gut, and skin.³⁵

Thus, it is likely that AZA can lead to secondary niacin deficiency and development of pellagra.

In our patient, underlying poor nutrition as a result of her chronic disease may have caused a relative niacin deficiency that was exacerbated by addition of azathioprine, culminating in the clinical presentation.

AZA-induced pellagra has been rarely reported, predominantly in adults.²³⁵ Herein, we briefly review previous AZA-induced pellagra cases reported in the literature. (Table 1).

To the best of our knowledge, there are only four cases of AZA-induced pellagra reported in the literature. All the reported cases were female patients, and only one case was in pediatrics. Duration of AZA use varied widely among patients from days to years. They all had the typical skin
| Number | Author/year | Patient age/sex | Preexisting disease | Duration of azathioprine use | Dermatologic manifestations | Other clinical manifestations | treatment | Prognosis |
|--------|-------------|----------------|---------------------|----------------------------|-----------------------------|-----------------------------|-----------|-----------|
| 1      | Jarrett et al. (1997)² | 42/female | Ulcerative colitis | 2 weeks | Browny-red lesions with peeling and distinct margin resembling sunburn on photo-exposed areas affecting both hands, the radial side of forearms, uncovered parts of feet, casal's necklace. | No mental confusion | Azathioprine was not discontinued, Nicotinamide 500 mg once daily. | Rash quickly improved and resolved |
| 2      | Jarrett et al. (1997)² | 17/female | Crohn's disease | 10 days | Marginated, reddish-brown, scaling rash extending down the radial aspects of forearms from just above elbows, casal's necklace, more erythematous on the dorsa of the hands, the front and backs of the legs, and the dorsa of the feet. | No mental confusion | Azathioprine discontinued, nicotinamide 50 mg three times daily. | No resolution until azathioprine discontinuation. Rapid improvement and full resolution after starting nicotinamide |
| 3      | Oliveira et al. (2011)⁵ | 47/female | Polymyositis | 15 years | Painful, well-defined, erythematous, arched plaque with exudative surface on anterior cervical region appearing after sun exposure. Multiple erosions on dorsum of the hand recovered by sero-hematic crusts. | Acute onset diarrhea, chelitis, glossitis | Azathioprine was discontinued, 300 mg niacin/day/oral began. | As skin lesions were improving, patient developed medullary aplasia leading to death three weeks later. |
| 4      | Zhao et al. (2018)⁴ | 50/female | Neuromyelitis optica | 3 months | Multiple painful erythematous, well-defined plaques on the dorsum of hands spreading to wrists, with scaling. | Multiple tongue and buccal mucus ulcersations, hyperpathia in thorax and abdomen, thoracic and abdominal pain, paroxysmal girdle-like tightening sensation leading to depression | Azathioprine was discontinued, nicotinamide 100 mg 3 times daily +vitamin B complex three times daily | Rapid improvement of skin lesions, within 1 month fully resolved skin lesions and no hyperpathia or zonesthesia |
| 5      | Present case | 14/female | Autoimmune hepatitis | 4 weeks | Well-demarcated violaceous to brown skin lesions with scaly desquamation and hyperpigmentation on the neck, dorsum of the hands, and feet which aggravated after sun exposure | Diarrhea, abdominal pain | Azathioprine was discontinued, oral nicotinamide 50 mg twice daily initiated. | Significant improvement after nicotinamide administration, full resolution after 2 months of treatment |
manifestations of pellagra, some experienced diarrhea, and none had neurologic disturbance. AZA was discon-
tinued in all the patients except one, and they were all initi-
ted on nicotinamide with different dosages from 150 to
500 mg/day. Significant improvement in skin lesions was
seen in all the cases. 2,3,5

Clinicians should consider AZA-induced pellagra in
any patient who develops a photo-distributed dermatosis
while undergoing treatment with AZA.

CONFLICT OF INTEREST
The authors declare that there is no conflict of interests
regarding the publication of this paper.

AUTHOR CONTRIBUTIONS
Bahareh Abtahi-Naeini had contributed to designing and
conducting the study. Parvin Rajabi had contributed to
pathological reports of the study. Shakiba Dehghani had
assisted in the preparation of the first draft of the manu-
script and manuscript revision. All authors have revised
the manuscript critically for important intellectual con-
tent, also have read and approved the content of the man-
uscript, and confirmed the accuracy or integrity of any
part of the work.

CONSENT
Written informed consents were obtained from the pa-
tient’s guardians for publication of this paper and any ac-
companying images.

DATA AVAILABILITY STATEMENT
The data that support the findings of this study are avail-
able from the corresponding author upon reasonable
request.

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