Lesson of the month 2: An unusual adverse reaction associated with pramipexole

Authors: Yasmina Tashkent and Vinod Aiyappan

Dopamine agonists such as pramipexole are commonly used in the treatment of restless legs syndrome (RLS) as well as Parkinson’s disease. Pramipexole’s common side effects are well documented; however, adverse skin reactions are less well known. In this case, a 45-year-old male farmer presented with excessive daytime tiredness and reported a history suggestive of RLS. He was initiated on pramipexole but developed a maculopapular erythematous rash in sun-exposed areas 8 days after its commencement. The skin rash resolved following pramipexole’s cessation and it is thought the patient experienced a drug-induced photosensitivity reaction to pramipexole.

This case highlights the potential for photosensitivity reactions to pramipexole, which is especially significant in countries like Australia where UV solar radiation is especially high.

KEYWORDS: Pramipexole, adverse reaction, skin

Presentation

A 45-year-old male farmer presented with excessive daytime tiredness and reported a history suggestive of restless legs syndrome (RLS). He was otherwise fit and well with no significant past medical history, no history of allergies and was not on any medications. Further investigation revealed a normal biochemical profile and iron stores. A sleep study confirmed periodic limb movements in sleep associated with sleep fragmentation (Fig 1). He was initiated on pramipexole (Sifrol) 0.125 mg nightly and had significant symptomatic improvement immediately after initiation of therapy. On the 8th day after initiation of treatment, the patient developed a maculopapular erythematous rash (Fig 2) in sun-exposed areas, while working outdoors on a sunny day. There were no symptoms of angioedema nor cardiorespiratory compromise. The patient’s wife took photos of the rash on her mobile phone (Fig 2). His GP stopped the pramipexole and started him on a carbidopa-levodopa combination (Kinson 25/100 mg). The skin rash resolved and the patient did not develop an adverse reaction to the carbidopa-levodopa.

Discussion

Dopamine agonists, including pramipexole, a non-ergot, second-generation dopamine agonist, are recommended as first-line in the treatment of RLS. Pramipexole is also commonly used in the treatment of Parkinson’s disease. Common side effects include insomnia, nausea, constipation, hallucinations, asthenia, sedation, somnolence, orthostatic hypotension, urinary frequency, dyskinesia and extrapyramidal side effects. In this case, it is thought the patient experienced a drug-induced photosensitivity reaction to pramipexole.

Photosensitivity reactions associated with exogenous agents are divided into photoallergy, phototoxicity and the induction/exacerbation of systemic illness where photosensitivity is a major clinical finding (eg pellagra, lupus erythematosus and porphyria cutanea tarda). Phototoxic and phototoxic reactions can be difficult to distinguish but can be differentiated by certain features (Table 1). Phototoxicity is thought to be a special type of a cell-mediated hypersensitivity reaction in which the drug is converted into an active compound by UV radiation, creating a photoantigen, which then triggers an immune response. This is in contrast to phototoxicity which occurs when light of a specific wavelength penetrates the skin and is absorbed by the drug or active drug metabolite that has reached the skin’s cells, exciting electrons in the drug and producing unstable singlet/triplet...
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states. The damage to organelles and macromolecules and the inflammatory mediators produced by the energy transferred from the unstable molecules returning to ground state subsequently causes cell damage.

To date, only one other case of a phototoxic reaction from pramipexole has been reported in the literature. Although rare, photosensitivity reactions to pramipexole should be discussed with patients being initiated on this drug, especially in countries like Australia where solar UV radiation is 13% greater in the Southern Hemisphere compared to corresponding sites in the Northern Hemisphere.3 ☐

Consent

Consent was obtained from the patient to publish the clinical details and images in this article.

References


Address for correspondence: Dr Yasmina Tashkent, Flinders Medical Centre, Division of Medicine, Flinders Drive, Bedford Park, South Australia 5042, Australia.

Email: Yasmina.Tashkent@sa.gov.au

Table 1. The difference between photoallergy and phototoxicity.

<table>
<thead>
<tr>
<th></th>
<th>Photoallergy</th>
<th>Phototoxicity</th>
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<tbody>
<tr>
<td>Dosage of drug required</td>
<td>Small</td>
<td>Large</td>
</tr>
<tr>
<td>Onset</td>
<td>&gt;24 hours</td>
<td>Minutes – hours</td>
</tr>
<tr>
<td>Clinical features</td>
<td>Dermatitis (acute, subacute or chronic) on exposed skin but may also spread to unexposed areas</td>
<td>Exaggerated sunburn only on exposed skin</td>
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<tr>
<td>Persistent light reaction</td>
<td>Yes</td>
<td>No</td>
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<tr>
<td>Histopathology</td>
<td>Epidermal spongiosis</td>
<td>Degeneration of epidermal cells</td>
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<tr>
<td></td>
<td>Dermal mononuclear cell infiltrate</td>
<td>Dermal vasodilation and oedema</td>
</tr>
<tr>
<td></td>
<td>Exocytosis of mononuclear cells</td>
<td>Sparse mononuclear infiltrate</td>
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Adapted from Gould JW et al.4

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Fig 2. Maculopapular confluent erythematous eruption involving the anterior chest wall (a) and left auricle (b).