Childhood Occurrence of Pemphigus

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ABSTRACT

Pemphigus is a chronic mucocutaneous disease that initially manifests in the form of intraoral blisters which spread to other mucous membrane and skin. This study describes an unusual case of chronic generalized childhood pemphigus disease in an 11-year-old girl, who presented with multiple vesicles all over her body. Such a condition is seen more often in older people rather than children. It is crucial for dental professionals to be familiar with the diagnosis of bullous skin diseases in children and adolescents, especially in its initial stages in order to prevent the serious consequences and morbidity. The article highlights clinical presentation, histopathology, and successful management strategies useful for pediatric dental practice.

Keywords: Acantholysis, Autoimmune, Blistering disease, Corticosteroids, Pemphigus.

CASE REPORT

An 11-year-old girl presented to the department of pediatric dentistry, with a complaint of multiple eruptions and blisters all over the mouth, which increased in size gradually over a period of 2 to 3 months and ruptured to form a crusty erosive surfaces with watery discharge (Fig. 1). Later, similar sores appeared on limbs, trunk, and the genital area which were painful and led to considerable discomfort (Figs 2 and 3).

Entire oral mucosa including the tongue was eroded and erythematous, causing extreme discomfort and pain during eating. There was no history of any drug intake during the past 6 months nor any systemic condition identified. The child presented with such a condition for the first time and there was no such disorder noted in the family. Nikolsky’s perilesional sign was positive.

The girl was hospitalized in the medical unit and comprehensively managed with the help of a dermatologist (Tables 1 and 2). Direct immunofluorescence was positive and perilesion biopsy containing intact lesion, revealed Tzanck cells, intraepidermal blister and suprabasilar acantholysis (Fig. 4). The connective tissue stroma showed dense mononuclear infiltration. A significant improvement in the condition was observed after 3 to 4 weeks following the standardized steroid treatment regime (Figs 5 to 7).
DISCUSSION

Unusual childhood occurrence, though quick response to treatment, however potentially life-threatening nature with substantial morbidity, justifies its consideration in routine dental practice. These chronic recurrent and painful lesions interfere with the daily activities of life, such as eating, drinking, talking, and personal relationships. Pediatric dentists have the unique opportunity since initial lesions occur in the oral cavity and complete remission is possible only with early diagnosis.

Prompt diagnosis and early initiation of aggressive therapy can combat the malignant course of disease in children. The treatment strategies should be based on the understanding of underlying pathogenic processes and
recurrence\textsuperscript{3,11-14} (Tables 3 and 4). Systemic corticosteroids and immunosuppressive therapy are the mainstay treatments for PV. Apart from steroids, adjuvant therapies include azathioprine, mycophenolate mofetil, dapsone, and rituximab in refractory cases.\textsuperscript{4,7,8} These modern therapies can effectively reduce the circulating antibodies, allowing patients to lead a normal life. Adverse effects associated with long-term use of steroids, such

\begin{table}[h]
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\begin{tabular}{|l|c|l|}
\hline
\textbf{Drugs} & \textbf{Dose, route, and duration*} & \textbf{Action} \\
\hline
Dexamethasone & 0.5 mL Inj IM (50-100 mg) 3 to 4 weeks & Modification of immune response (immunosupresion) \\
Roxithromycin & 150 mg Tab BID – 2 to 3 week & Antibacterial for secondary infection \\
Prednisolone & 10-20 mg Tab tapering to 5 mg BID – 2 to 3 months & Anti-inflammatory and modification of immune response \\
Hematopoietics & Oral capsule OD – 1 month & Nutritional supplement \\
NaCl saline & IV fluid & Electrolytic balance \\
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\caption{Systemic treatment regime}
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\begin{tabular}{|l|c|l|}
\hline
\textbf{Drugs} & \textbf{Dose, route, and duration} & \textbf{Action} \\
\hline
Triamcinolone & Local application for more than 3 weeks & Potent anti-inflammatory and alters immune response \\
Silver sulfadiazine and chlorhexidine & Local application for more than 2 weeks & Broad spectrum antimicrobial \\
Gentamycin with propyl salicylic acid & Local application for more than 2 weeks & Prevents secondary infections \\
Saline compresses over erosive lesions & Local application for more than 2 weeks & For soothing effect and control of edema \\
Chlorhexidine & Oral gargle for more than 3 weeks & Oral antimicrobial \\
\hline
\end{tabular}
\caption{Topical treatment regime}
\end{table}

Fig. 4: Acantholysis and suprabasilar separation

Fig. 5: Posttreatment view

Figs 6A and B: Lesions disappear following standard treatment regime
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Figs 7A to C: Healing of lesions all over the body

Table 3: Protocols for preventing recurrence

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<tr>
<th>Protocol</th>
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<tbody>
<tr>
<td>Maintaining healthy diet and weight</td>
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<tr>
<td>Avoiding sunlight and friction of body folds</td>
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<tr>
<td>Keeping flexural areas clean and dry</td>
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<tr>
<td>Wearing cool garments with absorbent pads</td>
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<tr>
<td>Regular evaluation of secondary infections</td>
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<tr>
<td>Systemic antibiotics, such as tetracycline and erythromycin</td>
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<tr>
<td>Topical use of antibacterial creams, such as benzyl peroxide</td>
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<tr>
<td>Long-term low-dose steroid maintenance therapy</td>
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<td>Controlling side effects of long-term steroids</td>
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Table 4: The bullous management portfolio

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<th>Protocol</th>
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<tr>
<td>Gold line mainstay of therapy – Steroids (Systemic prednisone 1 mg/kg/day and topical triamcinolone)</td>
</tr>
<tr>
<td>Broad-spectrum antibiotics for control of secondary infections</td>
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<tr>
<td>Improving the general health and hygiene of the patient (Fluid replacement, electrolytic balance, and multiple vitamins/minerals)</td>
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<tr>
<td>Symptomatic relief of pain, discomfort, burning, and itching (Paracetamol, astringents, and aluminium acetate)</td>
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<tr>
<td>Steroid sparing immunosuppressant and adjuvants (Mycophenolate mofetil, tracolimus, azathioprine, dapsone, retinoids methotrexate, cyclophosphamide, gold, cyclosporine, and chlorambucil)</td>
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<tr>
<td>Newer vistas - Plasmapheresis, intravenous immunoglobulins, anti-B cell monoclonal antibodies, CO2 laser vaporization, dermabrasion, proteinase inhibitors, chimeric molecules, cholinergic agonists, etc.</td>
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as weight gain, menstrual irregularities, growth retardation, osteoporosis, and hormonal disturbances in adolescence have always led to the search for newer steroid sparing and novel avenues for eradication of blisters at the molecular level. As we probe deeper into molecular aspects of the disease, our understanding of the pathogenesis begins to gain focus, offering new novel, and improved methods of therapy or even an opportunity to achieve a cure, which should mark the end of an era of blistering diseases.

REFERENCES

6. Amagai M, Koch PJ, Nishikawa T, Stanley JR. Pemphigus vulgaris antigen (Desmoglein 3) is localized in the lower