Throwing light on sweet syndrome and its homeopathic perspective

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Abstract
Sweet syndrome aka acute febrile neutrophilic dermatosis is one of the serious dermatological condition which goes undiagnosed due to several reasons. Unawareness of this condition may be one of the primary factors for missing the diagnosis which may lead to inaccurate treatment which may only lead to worsening the condition in a sick individual. Very few studies have been published in this area, hence the present investigator felt the need of reviewing few literatures and creating a material which will aid in early and accurate diagnosis of the condition. About 20 literature was reviewed based on the history, presentation, diagnosis and its probable homoeopathic treatment accordingly. All the information required for accurate diagnosis and its homeopathic approach was compiled into the present paper.

Keywords: Homoeopathy, sweet syndrome, acute febrile neutrophilic dermatosis, miasm, dermatology

1. Introduction
Homoeopathy has proved its efficacy time and again in various medical conditions, even when many systems of medicine have resorted to be of less advantageous in such conditions. The pathophysiology of the disease becomes the most important estimator for therapy in most other Medical systems but Homoeopathy is exclusively dependent on the symptoms so if a disease presents with symptoms then it can be managed with Homoeopathy.

There are certain problems existing in the medical science about certain diseases. Few of the diseases we do not know the exact aetiolo, few we are not yet sure about the exact pathophysiology, few diseases do not have the sufficient tools to establish the diagnosis. But Homoeopathy do not suffer with such ambiguity as the mere presence of signs and symptoms will guide them to the therapeutics. This is the biggest advantage of the second largest medical system in the world, Homoeopathy. The Homoeopathic pathology i.e. the miasmatic evaluation would help us to identify a position for the current disease in the Hahnemann’s classification of disease, which guides the physician in the treatment of that particular disease.

Homoeopathy has fashioned its scope in various dermatological conditions especially in all those diseases where the symptomatic presentation is clear. The gentle, harmless, permanent cure of these local manifestations according to the set of guidelines laid down by our Founder Dr. Samuel Hahnemann \(^\text{[1]}\) and other Stalwarts, where the malady can be removed without the use of external applications, thus making it the preferable mode of treatment for all. This review paper gives an insight about one of such rare dermatological conditions i.e. Sweet syndrome aka acute febrile neutrophilic dermatosis and its scope in homoeopathy.

2. History
In 1964, an English dermatologist named Dr. Robert D Sweet, described the disease \(^\text{[2]}\) that was observed in eight of his female patients. He described it as an illness with four distinct features

1. Fever
2. Acute, tender, red plaques
3. Leucocytosis
4. Papillary dermal infiltrate of neutrophils

This set of symptoms led to the nomenclature of the disease as ‘Acute febrile neutrophilic dermatosis’. Dr. Sweet expected the descriptive terms to be the regular nomenclature
of the condition. But ‘Sweet syndrome’[^3] ended up being the commonly used term in the fraternity. It is often confused with the illness to be related to disease caused by consumption sweet food items or Diabetes Mellitus due to its name.

In 1986, Dr. WP Su and Dr. HN Liu proposed a list of two major and four minor criteria for the diagnosis of Sweet Syndrome, but the modified and updated criteria given by Von Den Dreish P in 1994 is a generally accepted criteria among dermatologists.

3. Incidence

Sweet syndrome has shown a female predominance with the male female ratio 4:1[^4]. This female predominance is seen only in the classical varieties of SS and not seen in Malignancy associated SS and Drug induced Sweet Syndrome. Typically onset is seen between 30 to 60 years of age, but also seen in paediatric and geriatric ages also. There has been no observed racial predilection[^5].

In a study conducted by Hospach T, Von den Driesch P, Dannecker GE in 2009[^6], has shown that less than 70 children all over the world have reported with the diagnosis of Sweet syndrome out of which, only 42% had classical acute febrile neutrophilic dermatosis and the rest were associated with malignancies and drug induced cases. A need of further studies to identify the current prevalence/incidence of sweet syndrome is highly necessary for further studies on the condition.

4. Etiopathogenesis

Genetic predisposition, especially seen in Japanese population, showed a predominance of the gene HLA-Bw54 to be a risk factor in patients with Sweet syndrome[^7]. Although the exact aetiology is not identified till date, the various hypothesis gives a narrower explanation on what causes the disease. There are currently two well explained theories.

4.2 Sweet syndrome is a dermatological representation of classic immune complex vasculitis[^8]

Dr. Robert D Sweet, on careful histo-pathological analysis of cases with acute febrile neutrophilic dermatosis, consider the lack of vasculitis to be characteristic. But on more careful analysis of the histological sections of the classical SS cases, it showed that majority of cases have leukocytoclastic vasculitis. Although, in contrast, the presence of immunoglobulin and complement factors in direct immunofluorescence of the skin lesion, which is the hallmark of classic vasculitis, is absent in SS. Hence, even though the cases of classical SS showed leukocytoclastic vasculitis, there is no firm evidence to directly correlate Sweet Syndrome to be a representation of classical vasculitis.

4.3 A cascade of immunological events that is carried by T cells[^8]

**Fig 1:** A cascade of immunological events that is carried by T cells
Excessive infiltration of neutrophils into the skin (characteristic of Sweet Syndrome) will lead to dermal oedema. The continuous infiltration of these certain factors like (IL-1), IFN-γ, IL-8, G-CSF and TNF explains the histopathological changes seen in SS. Presence of all these factors, puts SS to close proximity to Pustular Psoriasis. T’cell research on SS will give a better understanding of pathogenic process occurring in the cells of SS.

5. Causes/risk factors [9]
Sweet Syndrome is a relative phenomenon than an individual entity itself. Presence of classical presentation of Sweet Syndrome can be used as an indicative factor for the underlying disease. Some of the diseases which may have a role in pathogenesis of Sweet Syndrome are

5.1 Infections
- Upper respiratory infection, commonly Streptococcal infection (Important risk factor of SS)
- GIT infection – Yersinia
- Chlamydia
- Cyтомegalovirus
- HIV
- Leprosy
- Tuberculosis

5.2 Malignancies [20% of Sweet Syndrome]
- Acute Myeloid Leukaemia
- Solid tumours of Gastro Intestinal tract, Genitourinary Tract, Breasts
- Patients with older age, anaemia or thrombocytopenia, and without arthralgia are more likely to have malignancy-associated SS. Studies have recommended that patients with SS without clear aetiology should be followed up for at least 16 months to exclude a possible underlying haematological malignancy [10].

5.3 Drug induced
a. G-CSF (Granulocyte Colony Stimulating factor)-Hematopoetic drug substance used to treat AML)
  b. Cotrimaxazole- the combination of trimethoprim (TMP) and sulfamethoxazole (SMX), used for the treatment of critically ill patients with infections caused by sensitive pathogens
  c. Minocycline- effective against gram-positive and-negative infections.
  d. Carbamazepine- anticonvulsants
  e. Oral contraceptives

5.4 Autoimmune disorders
a. Systemic lupus erythematosus
b. Rheumatoid arthritis
c. Dermatomyositis
d. Hashimoto’s thyroiditis

5.5 Inflammatory bowel diseases
a. Crohns disease
b. Ulcerative colitis

5.6 Others
a. Bechets disease
b. Erythema nodosum

6. Classification [11]
Hence Sweet Syndrome is broadly classified based on the clinical setting in which Sweet Syndrome occur
- Classical or idiopathic Sweet Syndrome
- Malignancy associated Sweet Syndrome
- Drug induced Sweet Syndrome

6.1 Histiocytoid Sweet syndrome [12]
This entity was first described by Requena et al. in 2005. It is considered as a variant of classic SS. It is histopathologically characterized by papillary dermal oedema with infiltration of histiocyte-like cells into the upper dermis. In this type the immature myeloid cells take the place of inflammatory cells which is usually seen in SS.

7. Clinical features

7.1 Cutaneous manifestations
- Abrupt/acute on set of
  - Well demarcatced lesions (Referred to as ‘Relief of mountain range’)
  - Tender lesions
  - Erythematous plaques, papules, nodes
  - Pseudo vesiculosis
  - Occasionally blisters that leads to annular slash aciform pattern
- Dermal oedema (Illusion of vesiculation) - Some lesions with dermal oedema gives a visual appearance as that of a vesicular lesion which may confuse the dermatologist to diagnose the condition clinically. Seen in the Histiocytoid variety of Sweet Syndrome
- Papules and plaques may combine to form large plaques up to 20 cm diameter
- In haemorrhagic malignancies, if local infection occurs, then atypical haemorrhagic bullae and ulceration may be formed.
- Ahn Et al- described variant of Xanthoma Like Lesions in Sweet Syndrome [12].

Location of Lesions [8]
- Head
- Neck
- Legs (lower limb lesions similar to erythema nodosum- morphologically and histologically)
- Arms (Particularly back of hands and fingers)
- Trunk is rarely involved

7.2 Neutrophilic dermatosis of hands [14]
It is a localized variant of Sweet syndrome seen on hands. It has a pustular appearance clinically and features of leukocytoclastic vasculitis histologically. Hence it has been derived that neutrophilic dermatosis of hands are a variant of localised classical Sweet syndrome [13].

After resolution of cutaneous manifestations
- Persistent erythema may be present
- Scarring is rare
- Some reports of focal elastolysis at sites of previous inflammation was seen in children.

7.3 Non cutaneous manifestations

Fever conditions
Fever, as mentioned in the nomenclature of the disease, is
seen in more than 50% of patients with SS. This is usually due to the underlying infection or chronic inflammatory disease manifestation. High fever maybe present before the onset of rash and may persist throughout the duration of illness [9].

7.4 Constitutional symptoms
- Arthritis or Arthralgia is commonly seen among the individuals affected
- Myalgia maybe seen during or after fever
- Headache is also a recurring phenomenon in Sweet syndrome.

7.5 Systemic complications
- Occular-Conjunctivitis (38%), Iritis, episcleritis, Limbal nodules
- Sudden visual loss due to occlusive venous vasculitis [15]
- Panuveitis with optic nerve involvement that resembles bechets disease [16]
- Liver- Transient liver enzymes elevation
- Oral- Aphthous ulcers
- Lung- Aseptic pneumonia
- Children- Tibial pain
- Renal- Proteinuria followed by hematuria, reduced creatitine clearance, Acute renal failure
- CVS complications- Most dangerous complication in childhood (<3yrs of age). Occurs in 14% of those affected. Mortality rate of those with CVS complications is 40%.
- CNS complications- Neuro-Sweet syndrome, a neurological complication of sweet syndrome, is seen mostly in Asians. Wide range of symptoms and signs can occur, depending on which part of the nervous system is affected. Usually presented with the SS typical skin lesions followed by neurological involvement or may even occur simultaneously. The most representative neurologic symptom that is seen is altered state of consciousness, followed by headache and memory disorders [17].

7.6 Histopathological features
- Dense dermal infiltrate of neutrophils with leucocytoclasia, prominent papillary dermal oedema
- Focal neutrophil micro abscess formation maybe seen.

8 Dreisch’s criteria for diagnosis [18]

Major criteria
- Abrupt onset of tender or painful erythematous plaques or nodules occasionally with vesicles, pustules or bullae.
- Predominantly neutrophilic infiltration of the dermis without obvious leukocytoclastic vasculitis.

Minor criteria
- Preceded by a nonspecific respiratory or gastrointestinal tract infection or vaccination or associated with inflammatory diseases, haemoproliferative disorders, solid malignant tumours, pregnancy.
- Accompanied by periods of general malaise and fever.
- Laboratory values during onset (3/4 necessary):
  - ESR >20 mm
  - C-reactive protein elevated
  - Segmented nuclear neutrophils and stabs >70% in peripheral blood smear
- Leucocytosis >8000.
- Excellent response to treatment with systemic corticosteroids or potassium iodide.

Two major and two minor criteria have to be present in order to diagnose the case as acute febrile neutrophilic dermatosis.

Drug-related Sweet syndrome: Diagnostic criteria [19]
A. Abrupt onset of painful erythematous plaques or nodules
B. Histopathological findings of dense neutrophilic infiltrate without evidence of leukocytoclastic vasculitis
C. Fever >38°C
D. Temporal relation between use of medication and clinical presentation or relapse with re-administration
E. Disappearance of lesions after drug discontinuation or treatment with systemic corticosteroids

9. Prognosis
- Depends on the severity of underlying or associated systemic disease
- Recurrences are not rare, and a chronic relapsing variant affects 15% of the patients
- In children, CVS complications have become associate with higher mortality rate [20]

10. Homeopathic perspective
Understanding the cause or the disease that is in the background becomes very necessary for homeopathic prescription for the condition. Hence in cases of Drug Induced or Malignancy Associated Sweet Syndrome, consideration of the background process should be definitely considered along with the scope of homeopathy in those conditions, before treating
In cases of classical SS where no background processes are in action, the patients may be treated symptomatically.

The first thing that needs to be considered when approaching any condition is the Hahnemannian classification of the disease and the treatment protocol suggested to deal with that phenomenon.
Sweet syndrome maybe classified according to the hahnemann classification of diseases under the following headings.
1. Classical Sweet syndrome (Individual Acute disease)
2. Drug induced sweet syndrome (Artificial chronic disease)
3. Malignancy assoc. sweet syndrome and Idiopathic sweet syndrome (One sided disease)

10.1 Sweet syndrome as an individual acute disease
In Aphorism 5 [1], Hahnemann says “Useful to the physician in assisting him to cure are the particulars of the most probable exciting cause of the acute disease....” About acute diseases. As we already mentioned in the above sections of this article, there are multiple exciting causes which may lead to the formation of this entity. Here the treatment of the cause becomes the primary focus which may remove the symptoms of Sweet syndrome correspondingly.

HA Roberts said “In acute manifestations, the vital functions are often restored to complete harmony by and through their own power”. Hence it is vital to understand that in mild cases, superficial treatment might be more
The acute manifestations most of the times, if not always, are the transient explosion of the latent psora. Hence it is advised to treat with homeopathic remedies based on acute totality and then complete the cure/avoid recurrence, if necessary, with an anti miasmatic remedy.

10.2 Sweet syndrome as an artificial chronic disease
This is the type of manifestation of Sweet syndrome which is due to the prolonged violent action of strong medication in its increasing doses. Because of the violent actions of these drugs, the vital force develops a reaction in the body to protect itself from the destructive primary action. According to Aphorism 75, these kind of diseases are the most incurable ones. This has to be remedied by the vital force itself. Identification of these factors and the corresponding action to avoid further progression becomes necessary before any therapeutic action is taken.

10.3 Sweet syndrome as a one sided disease
When we talk about the last variety, i.e. Malignancy associated Sweet Syndrome, it falls under the Hahnemannian classification of one sided diseases [1], where the diseases has only few developed symptoms. Here we need to follow the criteria of treatment of one sided diseases according to Hahnemann. The treatment of Malignancy Associated Sweet syndrome maybe done on the basis of symptom similarity and correspondence i.e. by administration of the remedy which covers the simillium of the most prominent symptoms of the presentation. This can be done using proved symptoms of homeopathic Materia Medica and with the aid of different repertories which guide us to the remedies of similar manifestation.

Of course, as Hahnemann suggested, we should focus on the accessory symptoms of the case available. If the remedy covers the uncommon peculiar symptoms, then cure may take place with the single remedy itself. If the selected remedy covers only the general symptoms and not the peculiar symptoms, then our totality should consist of both the accessory symptoms of medicine and disease. After the remedy administration, every time, a new totality will be formed based on the remaining symptoms and medicine may be prescribed. This process continues in succession until the cure is achieved.

The following are the remedies that is found in different repertories which may aid the homeopathic doctor is finding the remedy when dealing with cases of Sweet Syndrome.

### 10.3.1 Tender skin lesions

| Kents repertory [21] | BBCR [22] | Boericke repertory [23] |
|----------------------|-----------|-------------------------|
| Arnica               | Arnica    | Badiaga                 |
| Belladonna           | Belladonna| Cinchonha               |
| Nux-Vomica           | Phosphoric acid | Hepar              |
| Phosphoric Acid      | Asafoetida| Kali.C                  |
| Silicea              | Nux vomica| Lachesis                |
| Sulphur              | Sepia     | Oleander                 |
| Mezereum             | Vertrum Alb| Petroleum              |
|                      | Petroleum | Psorinum                |

### 10.3.2 Red Papular eruptions

| Kents repertory [21] | BBCR [22] | Boericke repertory [23] |
|----------------------|-----------|-------------------------|
| Sulphur              | Mercurius | Causticum               |
| Mercurious           | Belladonna| Chamomilla              |
| Kali-carb            | Rhus-tox  | Graphitis               |
| Calcarea carb        | Phosphoric acid | Lycopodium          |
| Phosphorous          | Graphitis | Mercurious              |
| Kali-bich            | Aconitum. | Mercurious              |
| Ammonium carb        | Sepia     | Sepia                   |
| Sepia                | Dulcamara | Pulsatilla              |
| Petroleum            | Pulsatilla| Ant-crud                |
| Kali-iod             | Petroleum | Causticum              |
| Sul-acid             |           |                         |
| Mezereum             |           |                         |

### 10.3.3 Dermal oedema

| Kents repertory [21] | BBCR [22] | Boericke repertory [23] |
|----------------------|-----------|-------------------------|
| Mercurius            | Rhus-tox  | Acet.acid               |
| Rhus-tox             | Mercurius | Agaricus               |
| Sulphur              | Arsenic   | Anacardium              |
| Sepia                | Sulphur   | Aps Mellifica           |
| Sambucus             | Sambucus  | Arsenic                 |
| Belladonna           | Belladonna| Digitalis              |
| Pulsatilla           | Lycopodium| Helleborous             |
| Kali-carb            | Pulsatilla| Prunus sp               |
| Thuja                | Kali-carb | Rhus tox                |
|                      |           | Thyrodisium             |

### 10.3.4 Fever eruptions

| Kents repertory [21] | BBCR [22] | Boericke repertory [23] |
|----------------------|-----------|-------------------------|
| Ignititia            | Mercurius | Euphrasia               |
| Aps Mellifica        | Arsenic   | Gelsemium              |
| Rhus-tox             | Sulphur   | Kali bich              |
| Copaiva              | Arsenic   | Aconite                 |
| Sulphur              | Bryonia   | Pulsatilla              |
| Rhus-v.              | Hyoscamus | Sticta.                |
|                      | Stramonium|                         |
|                      | Phosphorous|                        |
|                      | Causticum |                         |
10.3.5 Neutrophilic dermatosis of hands (Based on Kent’s repertory)

![Fig 2: Neutrophilic dermatosis of hands (Based on Kent’s repertory)](image)

10.3.6 Specific remedies from herring guiding symptoms

[24, 28]

- Skin eruptions, nodular, painful - Natrum mur
- Skin eruptions, pustular painful - Tarentula
- Skin subcutaneous swelling - Kali-mur
- Skin, Eruption, Pustules, Malignant, Pyaemia, hectic fever - Crotonus horridus
- Skin, Inflammation (dermatitis), Pressure, tender on – Arnica

10.4 Miasmatic understanding of sweet syndrome

The Miasmatic diagnosis of any condition will depend on the stage of the disease and its underlying cause for most of its spectrum.

In Sweet syndrome, when considering the Malignancy Associated SS, it will be predominantly under the syphilis or syphillis miasm (proliferative/destructive nature) [29] with respect to the grading and progression of the underlying malignancy, and rarely seen under Psoric miasm.

Although, when considering the skin symptoms alone, in cases of Classical acute febrile neutrophilic dermatosis, the following miasms are involved [27]

- Rawness or soreness of any skin eruptions – Syphilitic
- Red and haemorrhagic skin lesions - Tubercular
- Eruptions after vaccinations (drug induced) - Sycotic
- Exhaustion and general symptoms with Skin eruptions - Tubercular
- Nodular swelling – Sycotic
- Papular eruptions- Psoric
- Painful boils- Psora
- Scars after tumours- Sycotic

11. Conclusion

The scope of homoeopathy in dermatological conditions is not to be questioned after many evidence based studies done over the years. But the preparedness of any homoeopath to treat rare conditions is always necessary, especially considering the scope of thousands of remedies at our disposal. Understanding the mental cause, the maintaining factors of mind and body, and also the thorough understanding of the miasmatic diagnosis and treatment will help us encounter any rare conditions that come to us.

Further specific case studies or research studies of Homoeopathy on such rare skin conditions will help us improve the scope and create a mark in treatment of dermatological conditions with Homoeopathy.

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