Gomaya churneeeyam of Bhela Indriya Sthana - An explorative study

Abstract

‘Maharshi Bhela’ is one among the six disciples of ‘Acharya Punarvasu Atreya’ and he has composed a treatise known as ‘Bhela samhita’. Bhela samhita is having 8 sections and 120 chapters. Indriya sthana is one among the 8 sections of ‘Bhela samhita’ deals with the prognostic aspects. Among 12 chapters of ‘Bhela indriya sthana’, ‘Gomaya churneeeyam’ is the 9th chapter which comprises of 21 verses dealing with various emergency conditions having poor prognosis. The contents of ‘gomaya churneeeyam’ are unique and further in-depth exploration is required. Previous works conducted on ‘Charaka indriya sthana’ and ‘Bhela indriya sthana’ have explored various hidden concepts having both clinical and prognostic significance. Studies on ‘gomaya churneeeyam’ of ‘Bhela indriya sthana’ have been lacking and the present study is aimed to explore the contents of this chapter in terms of its prognostic significance. Various conditions like seborrheic dermatitis in an immunocompromised patients, extra pulmonary tuberculosis, nasopatilne duct cyst, age related macular degeneration, oral malignant melanoma, trigeminal autonomic cephalgia, lateral medullary syndrome, periodontitis, autonomic dysreflexia, systemic lupus erythematosus, limbic encephalitis, temporal lobe epilepsy, congenital erythropoietic porphyria, white spot lesions, sub conjunctival haemorrhage, chronic kidney disease, end-of-life dreams and visions, fever of unknown origin and chronic widespread pain associated with mortality have been documented in ‘Gomaya churneeeyam’ of ‘Bhela indriya sthana’.

Further works are required to establish the facts documented in this chapter.

Keywords: bhela indriya sthana, bhela samhita, charaka indriya sthana, end of life stages, indriya sthana, prognosis

Abbreviations: SD, seborrheic dermatitis; AIDS, acquired immunodeficiency syndrome; HIV, human immunodeficiency virus; PD, parkinson’s disease; AD, alzheimer’s disease; TB, tuberculosis; EPTF, extra pulmonary tuberculosis; DIC, disseminated intravascular coagulopathy; NPDC, nasopatilne duct cyst; ABC, aneuryysmal bone cyst; AMD, age related macular degeneration; LHS, laugier-Hunziker syndrome; PJS, Peutz-Jeghers syndrome; OMM, oral malignant melanoma; ACTH, adrenocorticotropic hormone; SUNCT, short-lasting unilateral neuralgiform headache attacks with conjunctival injection and tearing; TAC, trigeminal autonomic cephalgia; MS, multiple sclerosis; SPS, spenopatilne ganglion; NMOSD, neuromyelitis optica spectrum disorder; SUNA, short-lasting unilateral neuralgiform headache attacks with cranial autonomic symptoms; LMI, lateral medullary infarction; PICA, posterior inferior cerebellar artery; LMS, lateral medullary syndrome; ICA, Internal carotid artery; LE, lupus erythematous; SLE, systemic lupus erythematous; DLE, Discoid lupus erythematous; AD, autonomic dysreflexia; SCI, spinal cord injury; ANS, autonomic nervous system; IP, Ictal piloerection; TLE, temporal lobe epilepsy; LE, limbic encephalitis; CEP, congenital erythropoietic porphyria; WSL, white spot lesions; SCH, sub conjunctival haemorrhage; CCFs, carotid cavernous fistulas; GFR, glomerular filtration rate; CKD, chronic kidney disease; EPO, erythropoietin; Hb, haemoglobin; CNS, central nervous system; PRES, posterior reversible encephalopathy syndrome; NMS, neuroleptic malignant syndrome; GCS, glasgow coma scale; ICU, intensive care unit; ELD, EOL dreams and visions; EBV, epstein-barr virus; CMV, cytomegalovirus; FLO, fever of unknown origin; OM, otitis media; TAO, thrombangaits obliterans; PAD, peripheral artery disease; NBT, non-beneficial treatments; EOL, End-of-life stages;

Introduction

Maharshi Bhela and his five colleagues Maharshi Agnivesha, Jatukarma, Parashara, Harita and Ksharapani were the disciples of ‘Acharya Punarvasu Atreya’. ‘Maharshi Bhela’ has composed an Ayurvedic treatise known as ‘Bhela samhita’ (1000 - 2000 BC) and it was quoted by a number of authors and commentators. Planning and arrangement of the contents in the ‘Bhela samhita’ is similar to that of ‘Charaka samhita’. Bhela samhita has a number of unique ideas which were hidden and unexplored till date.1 Bhela samhita consists of 120 chapters and 8 sections similar to that of ‘Charaka samhita’. ‘Indriya sthana’ is one among the eight sections of ‘Bhela samhita’ and it is comprised of 12 chapters. ‘Indriya sthana’ deals with various ‘Arishta lakshanas’ (fatal signs and symptoms) which denotes poor prognosis and an imminent death.2

‘Gomaya churneeeyam’ is the 9th chapter of ‘Bhela indriya sthana’ which consists of 21 verses. This chapter deals with some arisha lakshanas which denotes an imminent death.3,4 Previous works conducted on ‘Charaka indriya sthana’5-8 and ‘Bhela indriya sthana’9 have established their clinical and prognostic significance. The contents of ‘gomaya churneeeyam’ are unique and needs further in-depth exploration. The present is aimed to evaluate the contents of ‘gomaya churneeeyam adhyaya’ in scientific terms. Studies on ‘gomaya churneeeyam adhyaya’ of ‘Bhela indriya sthana’ have been lacking and the present study is planned to explore the contents of this chapter in terms of its prognostic significance.

Review methodology

A literature search has been undertaken. Both ‘Google’ and ‘Google scholar’ databases have been used for search. Full text
articles with open access and abstracts published in ‘English’ were only considered. Articles published until November 2020, were only considered irrespective of their publication year or date of appearance. Ayurvedic literature (books and articles) published on ‘Indriya sthana’, ‘Charaka indriya sthana’, ‘Bhela indriya sthana’, and ‘Arishta lakshanas’ was considered. Articles related to emergency medicine and having prognostic importance have been searched from electronic databases by using relevant key words. No filters were applied during search. Authentic textbooks and pertinent websites were also referred.

**Exploration of the contents of gomaya churuneyam**

‘Purnam shirasi yasyaiva — maasaaddeham jahati sa’ (Verse 1).4,6

Scalp appears as covered with a dried powder of cow dung (gomaya churnam) which is unctuous (snehino) indicates an imminent death within a month (maasaaddeham jahati sa). The word ‘gomaya churnam’ denotes cow dung like powder (dandrufr) and ‘snehino’ denotes unctuousness or excessive sebum. Similar verse is documented in ‘Charaka indriya sthana’, 12th chapter (gomaya churuneyam indriyam adhyaya) and it was correlated as ‘SD in an immunocompromised patient’.6,8 SD is a chronic inflammatory skin disorder commonly seen in immunocompromised patients characterized by periods of remissions and exacerbations. The incidence of SD markedly increases in patients with AIDS. Dandruff (pityriasis sicca) is a fine scalp scaling considered as part of the adult SD spectrum. The scalp scaling associated with SD and dandruff is troublesome due to continuous shedding of these flakes (gomaya churnam) from the scalp. SD is associated with increased sebum secretion (senhino). Malassezia species have been found in immunocompromised patients. SD is seen in persons with an increased sebaceous gland activity (snehino), patients suffering with immunodeficiency (lymphoma & HIV-AIDS), neurological conditions (PD, AD, stroke and autonomic dysfunction), psychiatric conditions (major depression) and low ambient temperatures.20 The present verse denotes a condition of SD or Malassezia infection in an immunocompromised patient which may indicates death within a month.

‘Krushasaya kapha rogena — dvau maasau naativartate’ (Verse 2).4,6

The emaciated person (krushasya) having a kapha disease (respiratory diseases with cough and sputum production), cachexia (parikshaya) and aural bleeding (karnau raktam) with hemoptysis (raktau mukham) will die within two months (dvau maasau naativartate). The present verse denotes conditions like TB or EPTB or lung carcinoma metastasis to middle ear or temporal bone. Respiratory tract bleeding (epistaxis and hemoptysis) (raktau mukham) may be a marker of cancer (dvau maasau naativartate?) at several sites. Cancer-associated bleeding (karnau raktam & raktau mukham) may be caused by tumour compression or invasion of blood vessels, or by DIC, thrombocytopenia, or platelet abnormalities. Haemoptysis (raktau mukham) is also frequently seen in cancers and it varies from blood-streaked sputum to coughing up large amounts of pure blood (raktau mukham). Causes of hemothysis include malignancy, pulmonary embolism and other vascular causes, infection, trauma, inflammation, and bronchiectasis.20 Coughing up bright red or blood clots (raktau mukham) can be seen in carcinoma of the lung, tuberculosis (slesha parikshaya?) and pulmonary embolism. Blood-streaked, purulent sputum (raktau mukham) can be seen in bronchitis, bronchiectasis, or pneumonia and blood-tinged, white, frothy sputum is seen in congestive heart failure. Foul-smelling and bloody sputum (raktau mukham) is found in an anerobic lung process. Hemothysis in lung cancer (dvau maasa naativartate?) or tuberculosis (slesha parikshaya?) usually is a late symptom preceded by weight loss (krushasaya), change in cough (kapha rogena), fatigue, and other chronic symptoms.21 Bleeding from the ear (karnau raktam) is presumably the most significant symptom and it may occur early in cancer (middle ear and mastoid) (dvau maasa naativartate?).22 The clinical picture demonstrated in the present verse denotes TB or EPTB or lung carcinoma metastasis to middle ear or temporal bone associated with cachexia.

‘Vasya lohitakaabhaasa -- sa maasam naativartate’ (Verse 3).4,6

The emaciated or cachexic person having a reddish cyst like swelling at palate will die within a month. The word ‘lohitakaabhaasa’ denotes reddish lesion and ‘ambu’ denotes cystic swelling. The present verse denotes various conditions such as NPDC, ABC, radicular cysts, hemangiomas, mucoepidermoid carcinoma, pleomorphic adenoma, squamous cell carcinoma, non-Hodgkin’s lymphoma, and metastasis in the oral cavity and maxillofacial region.23 NPDC is located on the palatal aspect (taaluni drushyate) in midline of maxilla and it is the most common developmental, non-neoplastic, non-odontogenic cysts of the oral cavity. Odontogenic cysts (e.g., lateral radicular cyst, lateral periodontal cyst, odontogenic keratocyst), odontogenic tumors (e.g., ameloblastoma, odontogenic myxoma) and non-odontogenic tumors (e.g., central giant tumor, brown tumor of hyperparathyroidism, central hemangioma) etc conditions should be considered in the differential diagnosis of NPDC. NPDC may give rise to squamous cell carcinoma (sa maasam naativartate?) in the anterior zone of the upper maxilla.24 ABC lesions are nonneoplastic, locally aggressive and have high recurrence rates. ABC is an expansive osteolytic lesion consisting of blood-filled spaces (lohitakaabhaasa) and channels divided by connective tissue septa that can contain osteoid tissue and osteoclast-like giant cells. Hemorrhage (lohitakaabhaasa) can occur, especially if the alveolar bone is involved. Aspiration of a dark red or brownish hemorrhagic fluid (lohitakaabhaasa ambu) is suggestive of ABC.25

Radicular cysts are inflammatory jaw cysts and they develop as a consequence of advanced carious lesions. Increased vascularity with numerous dilated endothelial lined blood capillaries and thickened vessels filled with RBCs are evident along with areas of haemorrhage (lohitakaabhaasm ambu). Radicular cysts may become large enough to extensively erode adjacent bony structures (sa maasam naativartate?).26 Hemangiomas are blood vessel tumours occasionally seen on the palatal mucosa (taaluni drushyate), can occur as a capillary or cavernous type. The lesions may be bright-red, erythematous (lohitakaabhaasam ambu), and bilobulated with well-defined margins. Capillary hemangioma is associated with gingival vascular features and complications (sa maasam naativartate?) in the form of impaired nutrition and oral hygiene, increased accumulation of plaque and microorganisms, and increased susceptibility to oral infections, which can impair the systemic health of the affected individual. Chances of profuse bleeding (sa maasam naativartate?) are always a threat while dealing with hemangiomas.27

‘Arundhati na pashyet -- kshipram praan aur vimuchyate’ (Verse 4).4,6

The person unable to see (na pashyet) a star called ‘Arundhati’ in the sky (constellation of saptarshi) will die (praan aur vimuchyate) within 3 months (maasaaddeham jahati sa). The word ‘Arundhati’ denotes a star belonging to the sky (constellation of saptarshi) which will die (praan aur vimuchyate) within 3 months. The verse ‘Purnam shirasi yasyaiva -- maasaaddeham jahati sa’ (Verse 1) denotes conditions like TB or EPTB or lung carcinoma metastasis to middle ear or temporal bone associated with cachexia.

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The person who is having small or minute (anukaabhi), black or brownish-black or bluish-black or grey coloured (krishnaabhi), confluent (anucchhinnam), hyperpigmented or melanotic lesions in the oral mucosa (aasya), tongue (jihwa) and palate (taalu) will die soon (na sa jeevati). The present verse denotes a condition of visual impairment (due to various ocular and systemic disorders or diseases) associated with mortality.

‘Anukaabhishecha -- na sa jeevati taadrusha’ (Verse 5) 3,4

The person who is having small or minute (anukaabhi), black or brownish-black or bluish-black or grey coloured (krishnaabhi), confluent (anucchhinnam), hyperpigmented or melanotic lesions in the oral mucosa (aasya), tongue (jihwa) and palate (taalu) will die soon (na sa jeevati). The present verse denotes various life threatening conditions associated with oral melanotic macules. Increase in melanin content (krishnaabhi) of the oral epithelium (aasya) occurs in several conditions. The oral melanotic macule is a small (anukaabhi), discrete, brownish-black (krishnaabhi), flat lesion of the oral mucosa (aasya). Multiple, small (anukaabhi), melanotic macules in oral mucosa (aasya) can be seen in PJS. The oral mucosal pigmentary changes may occasionally be the earliest sign of Addison’s disease (increased ACTH). 3,5 Labial and oral melanotic macules (krishnaabhi aasya) are commonly encountered in a broad range of conditions ranging from physiological pigmentation to a sign of an underlying life-threatening disease (na sa jeevati). The most common lesion sites in LHS are the lips, the oral cavity (aasya), the tongue (jihwa), gingiva, and palatal mucosa (taalu). Typically, the cutaneous or mucosal lesions manifest as gray, brown, blue-black, or black macules (krishnaabhi) with a flat, smooth surface and relatively well-defined or indistinct margin (anucchhinnam) having the size of 2 to 5mm in diameter (anukaabhi) and lenticular, oval, or irregular in shape. Hyper pigmented macules (krishnaabhi) are distributed in variable numbers. Single or multiple lesions are observed, and occasionally, the lesions are confluent (anucchhinnam). Labial and OMM can be seen in various other conditions such as melanotic macule, melanocytic nevus, melanocanthoma, oral melanoma, PJS, McCune-Albright syndrome, Neurofibromatosis type 1, Carney complex, LEOPARD syndrome and Cronkhite-Canada syndrome. 3,2 The lesions in oral melanotic macule may be elevated and a well-defined blackish-brown colour (krishnaabhi) discoloration present on right buccal (aasya) and lingual mucosa (jihwa). Various conditions such as oral melanotic macule, melanoplakia, pituitary-based Cushings syndrome, post-inflammatory pigmentation, melanocanthoma, amalgam tattoo, Addison's disease, PJS and melanocytic nevi should be considered in the differential diagnosis of OMM. Poor prognosis of melanoma (na sa jeevati) may be due to early invasion of deeper structures due to proximity of bone and muscles increasing likelihood of metastasis. 3,6

‘Sheershaabhitaaapino -- naayamasteeti nirdishet’ (Verse 6) 3,4

The person who has been suffering with headache (sheershaabhiitaapino), respiratory symptoms (sleshma rogavata), hiccup (hikka) and rhinorrhea (vinishyandate) will die soon (naayamasteeti nirdishet). The present verse denote various conditions such as LMI, LMS, NMOSD, SENA, SUNCT, SPG, TAC with or without MS, pituitary apoplexy, pulmonary embolism, lower lobe pneumonia, acute or chronic upper or lower respiratory tract infections and their complications. LMS or Wallenberg syndrome is known to be associated with multitudes of peculiar symptoms such as hiccup (hikka), rhinorrhea (vinishyandate) and headache (sheershaabhiitaapino) in patients. Patients of LMS in the distribution of the PICA have shown intermittent rhinorrhea (vinishyandate). Rhinorrhea (vinishyandate) can be a part of the autonomic symptom of cluster headache (sheershaabhiitaapino) and SUNCT. ICA dissection can be associated with a clinical picture consistent with cluster headaches (sheershaabhiitaapino). The lateralized rhinorrhea (vinishyandate) suggests that the underlying cause is stroke and previous studies have reported the occurrence of contra-lateral rhinorrhea (vinishyandate) after a right caudate infarct.3,4 Middle level and dorsolateral lesion locations in LMI can induce hiccup (hikka) frequently. Hiccup (hikka) are infrequent result of LMI. Patients with LMI presenting with hiccup have also shown vertigo, dizziness, nausea (sleshma rogavata?), vomiting (sleshma rogavata?), and dysphagia.3,5

TAC is a primary headache syndrome (sheershaabhiitaapino) characterized by severe short-lasting unilateral headaches with ipsilateral cranial autonomic symptoms. Previous studies have described the association of TAC-like headache with intracranial lesions (e.g., pituitary adenomas, arteriovenous malformation of frontal and temporal lobes, brain tumour in posterior fossa, vertebral artery aneurysm, and orbitosphenoidal aspergillosis). Ipsilateral tearing, rhinorrhea (vinishyandate), eye redness and hiccup (hikka) were found in patients of MS presenting with TAC.3,6 SPG is the largest collection of neurons in the calvarium outside of the brain and it is also known as pitryogopatalline ganglion, nasal ganglion or Meckel’s ganglion. SPG has been postulated to be involved in facial pain and headaches (sheershaabhiitaapino) for over a century. TAC is characterized by parasympathetic (lacrimation, rhinorrhea, nasal congestion and edema) (sleshma rogavata?) activation and sympathetic dysfunction. Persistent hiccup (hikka) can also be seen in patients of SPG.3,7 Intractable hiccup (hikka), vomiting, lacrimation, and ipsilateral rhinorrhea (vinishyandate) can be seen in NMOSD patients. SENA / SUNCT are rare forms of TAC and can be found in patients of NMOSD.3,8

‘Yasya lomaani keshaaashche -- na sa jeevati taadrusha’ (Verse 7) 3,4

A person whose hairs on the body (lomaani) and the head (keshaa) appear as if being scorched or singed (plushyanteeva) or hairs standing up (horripilation or goose bumps or piloerection) (keshaani) appear as if being scorched or singed (plushyanteva) because this sign is so striking that it can suggest the diagnosis of SLE. LE is a chronic multi-organ autoimmune disease (na sa jeevati?) with a spectrum of clinical and serological presentations. The major target organs are the joints, skin, kidneys, lungs, and the nervous and serous systems (na sa jeevati?). In DLE patients, hair on the affected sites is usually brittle (plushyanteeva) and easily detachable.3,9 Piloerection
bone loss, halitosis and tooth loss. There is a positive association between periodontal disease and risk of lung, oral, and pancreatic cancers (na sa jeevati?).

‘Kshaarena vindhram gaatram – na cha sa jeevati taadrusha’
(Verse 9)\(^4\)

Body (gaatram) appears (drushyate) as washed with caustics (kshaarena vindhram) in both hot and cold seasons (samamushne cha sheete cha) denote an impending death (na sa jeevati). The present verse denotes skin colour changes associated with or without edema (kshaarena vindhram gaatram) and temperature changes at the EOL stages (na sa jeevati). Change in skin colour (skin becomes deadly pale or earth like colour, skin colour drains or turns white, skin turns pale or other colour changes) (kshaarena vindhram gaatram?), edema (of whole body or extremities or instep or underside of foot) (kshaarena vindhram gaatram?), body temperature decline (samamushne cha sheete cha?), vital power decline, blood pressure decline, bedsore/ wound deterioration (kshaarena vindhram gaatram?), and cyanosis (kshaarena vindhram gaatram?) are some among the signs and symptoms shown by the senile dementia patients at the end stages of life (na sa jeevati).\(^9\) The present verse explains the appearance of body or colour and temperature changes at the end stages of life.

‘Gaatreasu swara varneshu – na cha sa jeevati taadrusha’
(Verse 10)\(^4\)

The word ‘vaarilava plava’ (which floats on the water) denotes lightness or feebleness or decline. The person who shows body weight reduction or weight loss (gaatreshu vaarilava plava?), feeble voice (vaarilava plava?), decline of skin colour/ complexion or skin turns pale or white (varneshu vaarilava plava?) and excessive unctuousness or oiliness or greasiness even without application of oil (anabhyakteshu vaarilava plava?) or hyperhidrosis (anabhyakteshu gaatreasu varneshu plava plava?) will leave or die soon (na cha sa jeevati) as per the present verse. Body weight reduction (gaatreshu vaarilava plava?) is commonly seen among the elderly with EOL dementia (na cha sa jeevati). Positive association between body weight reduction (gaatreshu vaarilava plava?) and the prognosis of elderly patients (na cha sa jeevati) has been established. Body weight reduction (gaatreshu vaarilava plava?) still occurs at the dementia EOL stage (na cha sa jeevati), even if the appropriate amounts of nutrition are given. Skin becomes deadly pale or earth like colour, draining of skin colour or colour turns white, skin turns pale or other colour changes of skin (varneshu vaarilava plava?) are some of the features among various signs and symptoms seen in EOL stages of senile dementia patients (na cha sa jeevati). Skin turns pale in colour and discoloration of skin can also be seen in advanced cases of cancer.\(^4\) Weakening or quietening of voice (swareshu vaarilava plava?) is one among many signs and symptoms to be anticipated in the last days of life (na cha sa jeevati).\(^3\) Hyperhidrosis (excessive sweating) (vaarilava plava) can be seen in systemic disorders such as diabetes mellitus, hyperthyroidism, PD and other neurologic disorders, and tumours such as pheochromocytoma and lymphoma (na cha sa jeevati).\(^5\) Seborrhoeic facies is a manifestation of PD and it presents as plagues with greasy scales (anabhyakteshu gaatreasu vaarilava plava plava?) in areas that have an increased density of sebaceous glands. Dyshidrosis, including hyperhidrosis (anabhyakteshu gaatreasu vaarilava plava plava?) and hypohidrosis, is a common feature of PD. SD in PD may be due to autonomic dysfunction. Dyshidrosis is considered as part of the spectrum of autonomic dysfunction in PD. Drenching sweats (vaarilava plava?) also have been described in PD patients.\(^5\)
Gomaya churneeyan of Bhela Indriya Sthana - An explorative study

Citation:

The person who has been suffering with restlessness (arati), anorexia or loss of appetite or indigestion (avipaaka), weight loss or cachexia or sarcopenia (kaarshya) and weakness or vital power decline (daurbalya) will die soon (na sa jeevati). The present verse denotes a condition of anorexia-cachexia syndrome associated with delirium. Cachexia (kaarshya) is a complex metabolic process associated with underlying terminal illnesses (na sa jeevati) and it is characterized by anorexia (avipaaka) and loss of fat and muscle mass (daurbalya? kaarshya?). Loss of body weight (kaarshya) and reduced energy levels (daurbalya) are also found in cachexia (kaarshya). Cachexia (kaarshya) is known to be associated with advanced dementia (na sa jeevati). Advanced stages of dementia are marked by substantial unintentional weight loss (kaarshya), malnutrition (avipaaka), sarcopenia (kaarshya? daurbalya?), anorexia (kaarshya), lethargy (daurbalya), altered immune function, and cachexia. The present verse denotes delirium due to an underlying terminal illness or delirium superimposed on advanced dementia patients with cachexia. Agitation (arati), restlessness (arati) and increased psychomotor activity (arati) are seen in delirium. During last days of life patients often experience progressive functional decline (daurbalya) and worsening symptom burden (na sa jeevati). Many symptoms such as anorexia-cachexia (avipaaka & kaarshya), weight loss (kaarshya / daurbalya), decreased quality of life and delirium can be found in last days of life (na sa jeevati).

‘Yastu deenam anaatho vaa -- yatha preta stathaiva sa’ (Verse 14)34

The person who has been suffering with depressed mood (deenam), loneliness or isolation or social withdrawal (anaatho), weakness or fatigue or lack of interest or anhedonia (balena pariheeyate) and hypochondriasis (bhidyam aarogam aapnoti) or feeling good while suffering with illness or disease (mithya arogyam aapnoti) should be considered as already dead (yatha preta stathaiva sa). The present verse denotes a wide variety of conditions like MDD, catatonia, negative symptoms of schizophrenia and mood disorders associated with poor prognosis. Depressed mood (deenam), anhedonia (loss of interest or pleasure in nearly all activities) (anaatho), significant weight loss (balena pariheeyate), psychomotor retardation (balena pariheeyate), fatigue or loss of energy (balena pariheeyate), feelings of worthlessness (anaatho) and interpersonal rejection sensitivity (anaatho) are the characteristic features of MDD.37 Catatonia of the retarded type is associated with signs reflecting a paucity of movement, including immobility, staring, mutism, rigidity, withdrawal (anaatho) and refusal to eat, along with more bizarre features such as posturing, negativism, grimacing, waxy flexibility, echolalia / echopraxia, verbigeration, stereotypy, and automatic obedience. Catatonia appears to be a risk factor for the development of neuroleptic malignant syndrome (which has a mortality rate of 10%) (yatha preta stathaiva sa). Negative symptoms are a core component of schizophrenia that account for a large part of the long-term disability and poor functional outcomes (yatha preta stathaiva sa) in patients with the disorder. The negative symptom domain consists of five key constructs: blunted affect (deenam), alogia (reduction in quantity of words spoken) (balena pariheeyate), avolition (reduced goal-directed activity due to decreased motivation) (balena pariheeyate), asociality (anaatho), and anhedonia (reduced experience of pleasure).38 Hypochondriasis is a unique disorder with the primary feature of persistent preoccupation (bhidyam aarogam aapnoti) with the possibility of having one or more
serious physical disorders. Hypochondriasis is also noted to have an association with anxiety and depression (deemaam).69

‘Anuvrutta yatha jantu -- yatha preta stathaiva sa’ (Verse 15)44

The person who has been suffering with loss of consciousness (caused by pitta) (pitrena parimurcchata) and making incomprehensible sounds or inappropriate words (samudhva vaakya) will die soon (yatha pretha stathaiva sa). The present verse denotes a condition of coma caused by various underlying fatal conditions. Impaired consciousness (parimurcchati) can be considered in terms of reduced alertness or ability to be aroused, awareness or both, with coma defined as ‘a completely unaware patient unresponsive to external stimuli (parimurcchati) with only eye opening to pain with no eye tracking or fixation, and limb withdrawal to a noxious stimulus at best’. The causative factors for non traumatic coma (parimurcchati) can be classified in to four main categories: neurological (ischemic stroke, intra-cerebral or subarachnoid haemorrhage, subdural hematoma, brain tumour, cerebral lymphoma, multiple brain metastases, CNS infections, cerebral abscess or oedema, hydrocephalus, axonic brain injury, PRES and trauma), metabolic (hypo / hyperglycaemia, hypo / hypernatraemia, hypercalcaemia, Addisonian crisis, hypothyroidism, uraemia, hypercapnia and septic / hepatic encephalopathy), diffuse physiological brain dysfunctions (seizures, alcohol intoxication, opioid toxicity, poisoning, hypothermia, NMS and serotonin syndrome) and psychiatric or functional (psychiatric coma and malingering). Scores allotted for ‘inappropriate words’ and ‘incomprehensible sounds’ (samudhva vaakya) in GCS are 3 and 2 respectively. Prognosis depends on a number of factors and the mortality rate varies from 25–87% in coma patients (yatha pretha stathaiva sa). Non traumatic coma patients presenting with a stroke have the highest mortality (yatha pretha stathaiva sa). Patients with a lower GCS at presentation 3 - 5 have a significantly higher mortality than those with a GCS of 7–10 (yatha pretha stathaiva sa).61

‘Yastvaasane atha shayane -- yashcha saashruni khaadati’ (Verse 16)44

The person who has been suffering with restlessness or agitation (ratim na labhate) and not getting comfort in sitting (aasane) or lying (shayane) will die soon (sheeghram kurute kaalam). The present verse denotes the condition of terminal restlessness or terminal agitation seen in delirium. Delirium in the last few days of life (often referred to as terminal restlessness or terminal agitation) (sheeghram kurute kaalam) is often ongoing and irreversible. Hyperactive or agitated delirium is characterised by increased motor activity with agitation (ratim na labhate), hallucinations and inappropriate behaviour (ratim na labhate). Delirium is prevalent at the EOL (sheeghram kurute kaalam), particularly during the final 24–48hours. All patients at the EOL (sheeghram kurute kaalam) can be considered at high risk of delirium.62

‘Aaruhya vaanaram yastu -- swapne tu kushalaar naraa’ (Verse 17)44

The person who sees dreams (swapne) such as riding on a monkey (aaruhya vaanaram) and roaming without any purpose or objective (sankalpa naavabudhyate) should be considered as already dead (tamaahu paralokaaya). The present verse denotes ELDVs or delirium. Since ancient times, people have recorded dreams and visions (swapne) experienced by individuals (naraa) at the end of their lives (tamaahu paralokaaya). These experiences may involve visual, auditory and kinesthetic experiences (aaruhya vaanaram). ELDV experiences can occur months, weeks, days or hours before death (aaruhya vaanaram). ELDVs including a theme of going or preparing to go somewhere (aaruhya vaanaram) were common. ELDVs may be prognostically significant based on changes in dream content (aaruhya vaanaram) and increased frequency as death nears (tamaahu paralokaaya). The main quality of pre-death dreams (tamaahu paralokaaya) is a sense of personal meaningfulness.63 Discharged ICU patients often recall experience vivid dreams, hallucinations or delusions (aaruhya vaanaram). These dreams may be persecutory in nature and are sometimes very frightening also. It is possible that these memories stem from times when the patient was experiencing delirium, a common syndrome in the critically ill (tamaahu paralokaaya). A longer ICU stay was significantly associated with the experience of ICU dreaming (aaruhya vaanaram).64 In present verse, ‘aaruhya vaanaram’ and ‘sankalpa naavabudhyate’ (riding on a monkey and roaming on it without any objective or purpose) denotes agitation or restlessness or increased psychomotor activity etc. especially seen in delirium patients. The dream object or symbol mentioned in the present verse is ‘monkey’ (which is the displacement for patient), dream content (riding on a monkey without having any purpose) denotes underlying ‘agitation’ or ‘restlessness’ or ‘increased psychomotor activity’, dream type is ‘absurd’, source of the dream is ‘internal organic pathology’ or ‘somatic stimuli’ and dream mechanisms involved are ‘displacement’, ‘representation’, ‘secondary revision’, ‘projection’, ‘inversion’ and ‘condensation’.65

‘Parismavatsaraad yasya -- yatha preta stathaiva sa’ (Verse 18)44

The person who has been suffering with continuous or prolonged or persistent fever (jwara), which occurs both in hot and cold seasons (ushno va yadi va shheeto) and having more than a year duration (parisamvatsaraad) should be considered as already dead (yatha pretha stathaiva sa). A number of bacterial, viral, parasitic and fungal infections can cause prolonged fever (parisamvatsaraad jwara). Many infectious agents cause a chronic and low grade fever (parisamvatsaraad jwara). Chronic infections can be categorised according to the type of organism such as bacterial causes (oesomyelitis, occult abscess, bacterial endocarditis, Rickettsia, Coxiella, Bartonella spp, enteric fever, tuberculosis, brucellosis and yersiniosis), parasitic causes (leishmaniasis and malaria), infectious mononucleosis (caused by EBV, CMV and Toxoplasma cause a fever lasting more than three weeks), endemic fungi (deep seated mycoses), and other opportunistic pathogens that can present with FUO in immunosuppressed individuals (yatha pretha stathaiva sa), such as those with chronic renal impairment and HIV.66 Sepsis accounts for 74 % of fever in hospitalised patients and, of the remainder, malignancy, tissue ischaemia, and drug reactions account for the majority. Patients exposed to higher temperatures and for longer periods of time (parisamvatsaraad jwara) are more at risk of complications such as multi organ failure and death in extreme cases (yatha pretha stathaiva sa).67 Persistent fever (parisamvatsaraad jwara) suggests resistant bacteria, a viral cause, unsuitable antibiotic or a complication of OM. Unexplained persistent and recurrent fever (parisamvatsaraad jwara) for more than a month is considered as a moderate severe HIV (yatha pretha stathaiva sa’). Persistent or prolonged fever (parisamvatsaraad jwara) can also be seen in asecic or TB meningitis, pulmonary TB, disseminated TB, miliary TB, and in EPTB. Children with combined intrapulmonary and EPTB have a higher peak and a longer duration of fever (parisamvatsaraad jwara).67 The present verse denotes prolonged fever caused by a wide variety of conditions associated with poor prognosis.

‘Yasya jaata pramehasya -- yatha preta stathaiva sa’ (Verse 19)44

Manifestation of carbuncles (pitaka) or sinuses (shatapada) or multiple pus points (shatapada or pitaka paandura) other complications (upadraa) in congenital diabetes (jaata prameha) patients denotes an
impending death (yatha preta stathaiva sa). Skin carbuncle (pitaka) is a necrotizing infection of the skin and subcutaneous tissues composed of a cluster of furuncles with multiple draining sinuses (shatapada). Carbuncles usually open and drain through multiple channels (shatapada). They are commonly associated with diabetic patients (jaata prameha). Skin and soft tissue infections are common in diabetics (especially when uncontrolled) (jaata prameha). The bacteria penetrate the skin and the subcutaneous tissues to form a series of communicating abscesses (pitaka paandura?), which discharge by separate opening on the surface (sieve like appearance) (shatapada).46

‘Yasyordhwam kramate vaayu -- yatha preta stathaiva sa’ (Verse 20)47

The person (yasya), in whom the aggravated vaayu traverses (kramate) upper parts of the body (urdhwam) up to the ears (shrotram) and all over the body (adha pravartate & sarvaani cha) causing pain (rupturing type) (prabhihiyate) should be considered as dead (yatha preta stathaiva sa). An increased risk of death (yatha preta stathaiva sa) in people (yasya) with chronic pain (vaatajata), particularly from cancer has been established. Musculoskeletal pain (vaatajata) may also be associated with increased mortality (yatha preta stathaiva sa). Chronic pain (which lasts for three months or longer) (kramate) is experienced by 30% (approximately) of adults (yasya) and commonly occurs in multiple body sites (urdhwam, adha, shrotram and sarvaani). Chronic widespread pain (urdhwam, adha, shrotram and sarvaani vaayu) is a phenotype that captures people (yasya) with more severe pain (vaatajata) that has a greater impact on outcomes (yatha preta stathaiva sa). Few studies have established the positive association between chronic or widespread pain (urdhwam, adha, shrotram and sarvaani kramate vaayu) and an increased rate of all-cause mortality (yatha preta stathaiva sa). Significant associations between chronic or widespread pain (urdhwam, adha, shrotram and sarvaani kramate vaayu) and increased risk of death (yatha preta stathaiva sa) from cancer, cardiovascular disease, cerebrovascular disease, liver cirrhosis, influenza, pneumonia and septicemia were reported.48

The present verse denotes wide variety of conditions such as vascular (acute arterial occlusion, arterial embolus, aortic dissection, arterial thrombosis, TAO, PAD, haemorrhage, cardiovascular and cerebrovascular diseases etc.) or neurological (peripheral neuropathies, myelopathies, myelitis, neuroinflammatory, neurodegenerative, demyelinating conditions) or musculoskeletal (myopathies, infectious & non-infectious myositis, autoimmune inflammatory disease, connective tissue diseases, degenerative diseases etc.) or malignancy related or infectious etc. associated with poor prognosis.

‘Ityete lakshani yuktam -- rakshaan aatma yasha sputham’ (Verse 21)44

The brave (veero) and wise (dheero) physician (bhishak) should not attempt to treat (nopakrameta) the patient (evam maanavam), who displays the signs and symptoms mentioned in this chapter (ityete lakshani yuktam). Physician can protect his fame or reputation (rakshaan aatma yasha by avoiding such patients (who displays the signs and symptoms mentioned in this chapter). Treating incurable diseases leads to financial losses, defamation, loss of reputation, social harm, liability to legal punishments etc. to the concerned physician. Hence physician should avoid treating incurable diseases to protect his reputation and dignity.1 The conditions explained in the present chapter represent NBT especially at the EOL stages, which are futile and physician should avoid treating such patients. Attempt to treat conditions with high mortality rates or poor prognosis seen especially at EOL stages may generate false hope, inappropriate use of scarce healthcare resources or infrastructure and staff dissatisfaction with anticipated poor outcomes. Most of the ICU patients will die no matter how best the treatment is provided to them (just like the conditions explained in the current chapter). Medically inappropriate care aggravates pain, suffering, and discomfort to the patients.26 According to ‘Maharshi Bhela’, patients with EOL stages should not be treated (as they won’t receive any benefit to the treatments no matter how good they are).

### Table 1 Contents of ‘Gomaya churneeyam’ chapter

| Verse | Relevant condition |
|-------|--------------------|
| ‘puram shirasi yasyaiva -- maasaddeham jahaati sa’(B. I. 9/1) | Seborrheic dermatitis in immunocompromised patient or AIDS patients; Tuberculosis or extra pulmonary tuberculosis or lung carcinoma with metastasis to middle ear & temporal bone; |
| ‘krushaya kapha rogena -- dvau maasu naativrata(te)(B. I. 9/2) | Nasoplatine duct cyst; Aneurysmal bone cyst; radicular cysts, hemangiomas, squamous cell carcinoma, non-Hodgkin’s lymphoma, and metastasis in the oral cavity and maxillofacial region; |
| ‘yasya lohitakaabhaasa -- sa maaasaatnaativrata(te)(B. I. 9/3) | Visual impairment due to various ocular and systemic diseases associated with mortality;Age related macular degeneration; Cataract; |
| ‘arundhati na pashyat -- khipram praan aur vimuchyate(B. I. 9/4) | Oral malignant melanoma; Laugier-Hunziker syndrome; Peutz-Jeghers syndrome; |
| ‘anukabhishcha -- na sa jeevati taadrusha(B. I. 9/5) | Short-lasting unilateral neuralgiform headache attacks with conjunctival injection and tearing; Trigeminal autonomic cephalgia; Neuromyelitis optica spectrum disorder; |
| ‘sheershaabbitaapino -- naayamasteeti nirdishhet’(B. I. 9/6) | Systemic lupus erythematosus; Autonomic dysreflexia; Spinal cord injury; Lctal piloerection; Temporal lobe epilepsy; Limbic encephalitis; |
| ‘yasya lomaani keshhaaasha -- na sa jeevati taadrusha(B. I. 9/7) | Periodontitis; White spot lesions; Congenital erythropoietic porphyria; |
| ‘yasya kaalaantare dantaa -- na sa jeevati taadrusha(B. I. 9/8) | Changes of skin colour and temperature at end stages of life; Hyperhidrosis and seborrheoa in Parkinson’s disease; |
| ‘kshaarena vidhrutam gatram -- na cha sa jeevati taadrusha(B. I. 9/9) | |
| ‘gaatreshu swara varneshu -- na sa jeevati taadrusha(B. I. 9/10) | |
Conclusion

‘Gomaya churneeyam’ is the ninth chapter of ‘Bhela Indriya sthana’, which has 21 verses. Various signs and symptoms that are having prognostic significance have been explained in this chapter. Most of the verses are unique and untraceable in other classical Ayurvedic texts. Various conditions such as seborrhoeic dermatitis in an immunocompromised patients, extra pulmonary tuberculosis, nasalopatine duct cyst, age related macular degeneration, oral malignant melanoma, trigeminal autonomic cephalgia, lateral medullary syndrome, periodontitis, autonomic dysreflexia, systemic lupus erythematosus, limbic encephalitis, temporal lobe epilepsy, congenital erythropoeitic porphyria, white spot lesions, sub conjunctival haemorrhage, chronic kidney disease, end-of-life dreams and visions, fever in situations where fever is not a sign of infection, chronic widespread pain associated with mortality etc. are documented in this chapter by ‘Maharshi Bhela’. Physician should avoid treating the end-of-life stages mentioned in this chapter. Further studies are required to substantiate the findings claimed in this chapter.

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Conflicts of interest

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