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# A CLINICAL ASSESSMENT OF *AYURVEDIC* TREATMENT OUTCOMES IN A PATIENT WITH STAGE II CKD AND HYPERTENSION

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# **ABSTRACT**

Chronic Kidney Disease (CKD) Stage II often coexists with hypertension, forming a complex bidirectional relationship that complicates treatment and worsens outcomes. This case study explores the integrative management of CKD Stage II with hypertension in a 36-year-old male patient treated at Jeena Sikho Lifecare Limited Hospital, Derabassi, India. The patient presented with complaints of generalized weakness and bilateral leg swelling. A personalized *Ayurvedic* treatment plan was administered over four months. Clinical outcomes showed significant improvement. Blood pressure normalized from a peak of 180/110 mmHg to 120/90 mmHg, and pulse rate stabilized. Global GFR improved from 65.8 to 68.1 ml/min, while serum creatinine decreased from 2.11 to 1.40 mg/dL. Hemoglobin levels rose to 11.7 gm/dL, and urea and BUN levels declined, indicating better renal and systemic function. The findings suggest that *Ayurvedic* management may offer a safe and effective approach for stabilizing CKD and associated hypertension. Although this case demonstrated encouraging outcomes, it is based on a single patient observation. To substantiate the efficacy, safety, and reproducibility of integrated *Ayurvedic* interventions in CKD with hypertension, larger-scale randomized controlled trials are essential.

**KEYWORDS:** Ayurveda, Chronic Kidney Disease (CKD), Daruharidra, Guduchi, Hypertension, Punarnava, Shilajit, Uchcha Raktachap, Vrikk vikar.

# INTRODUCTION

Chronic Kidney Disease (CKD) stage II is frequently associated with hypertension, which not only contributes to kidney damage but also complicates disease progression and outcomes.<sup>[1]</sup> The relationship between CKD and hypertension is bidirectional, hypertension can both cause and result from declining kidney function.<sup>[2]</sup> Hypertension is common among CKD patients, with studies showing that approximately 46% of those with CKD stage II are affected.<sup>[3]</sup> In a cohort study, 32.35% of stage II CKD patients had systolic hypertension.<sup>[4]</sup> Uncontrolled hypertension in these patients is linked to a

higher risk of mortality, with systolic blood pressure (SBP) levels above 120 mmHg associated with increased death rates. [5] Pulmonary hypertension is another concern, reported in 51.35% of CKD patients, further complicating treatment and elevating mortality risks. [4] Management strategies typically involve the use of reninangiotensin system (RAS) inhibitors and diuretics, both effective in controlling blood pressure and reducing proteinuria. [6] However, it is also noted that not all patients with hypertension experience the same progression in kidney function decline, highlighting the

need for personalized management strategies based on individual patient factors.

In Ayurveda, the Samprapti (pathogenesis) of CKD stage II with hypertension is understood through the lens of Dosha, Dhatu, and Srotas imbalance. The disease primarily involves vitiation of Vata and Kapha doshas, with secondary involvement of Pitta in some cases. The affected Srotas are Mutravaha (urinary channels), Raktavaha (circulatory channels), and Medovaha (fat tissue channels). Long-standing Agnimandya (digestive fire suppression) and accumulation of Ama (metabolic toxins) initiate the pathology, leading to obstruction (Srotorodh) in the Mutravaha Srotas. This results in Mutrakshaya (decreased urine output) and Mutradah (burning micturition) in early stages, progressing to Mutraghaat (obstructive uropathy) in advanced stages. [10]

The progressive tissue damage corresponds to *Dhatu Kshaya*, particularly of *Rakta*, *Meda*, and *Majja Dhatus*, leading to systemic weakness and renal failure features. Hypertension in this context is seen as a manifestation of *Vyana Vata* and *Rakta Dushti*, often due to *Avarana* (encapsulation or occlusion) of *Vata* by *Kapha* or *Meda*. Ayurvedic diagnosis is based on *Rog* and *Rogi Pariksha* (examination of disease and patient), involving *Dashavidh Pariksha* (ten-fold examination), *Ashta vidh Pariksha* (eight-fold examination), and pulse diagnosis (*Nadi Pariksha*), which help in assessing *doshic* predominance, tissue damage, and prognosis. This comprehensive diagnostic approach guides the individualized treatment plan aimed at restoring *Dosha-Dhatu-Srotas* equilibrium. The *Samprapti Ghatak* hateleft.

Table 1: The Samprapti Ghatak of this case.

Dosha (Bio-energetic forces) • Vata (especially Apana and Vyana Vata) - primary involvement • Kapha (structure and stability) and Pitta (metabolism) - secondary involvement Dushya (Affected body tissues) Ras (plasma/lymph), Rakta (blood), Meda (fat), Mamsa (muscle), Majja (bone marrow/nervous tissue) progressive tissue depletion (Dhatu Kshaya) Srotas (Body channels) Mutravah Srotas (urinary channels), Raktavah Srotas (blood circulation channels), Medovah Srotas (metabolic channels) Srotodushti Prakar (Type of channel vitiation) Sanga (obstruction), Siragranthi (vascular blockage), Vimarg gaman (misdirection of flow) Udabhav Sthan (Origin site) Pakvashaya (colon) Adhisthan / Waktasthan (Site of manifestation) • Basti (urinary system, kidneys), Hridaya (heart – due to Vyana Vata disturbance) Rog Marg (Pathway of disease) · Abhyantar Rog Marg (internal pathway) Vyadhi Swabhav (Nature of disease)

• Yapya (manageable with ongoing treatment), Chirakari (chronic), Krichhra Sadhya (difficult to cure)

The *Ayurvedic* approach to managing CKD Stage II with hypertension is rooted in restoring balance among the disturbed *doshas*, primarily *Vata* and *Kapha*, and supporting the proper function of the urinary and circulatory systems. <sup>[16]</sup> Treatment emphasizes correcting the underlying imbalances responsible for obstruction and degeneration in the body, particularly addressing the dysfunction of *Vyana Vata* and the weakness of tissue metabolism (*Dhatu Kshaya*). <sup>[17]</sup> Attention is given to improving digestion and assimilation to reduce the formation of toxins (*Ama*), which can obstruct channels (*Srotas*) and aggravate disease progression. <sup>[18]</sup> Lifestyle and dietary adjustments are also integral, aimed at reducing systemic stress and maintaining physiological harmony.

# **OBJECTIVE**

Examine the impact of *Ayurvedic* interventions in a 36-year-old male patient with CKD II and hypertension.

# MATERIALS AND METHODS

### I. Case Report

A 36-year-old male diagnosed with CKD stage II and hypertension presented at Jeena Sikho Lifecare Limited Hospital in Derabassi, India, on February 10, 2024. She reported experiencing general weakness and bilateral leg swelling. There was no significant family history. Findings from the initial *Ashta-vidh Pariksha* are provided in **Table 1**. The patient's vital signs recorded during the first visit are shown in **Table 2**, and the laboratory test results throughout the course of treatment are detailed in **Table 3**.

Table 1: The Ashta vidh Pariksha (examination) on the visits.

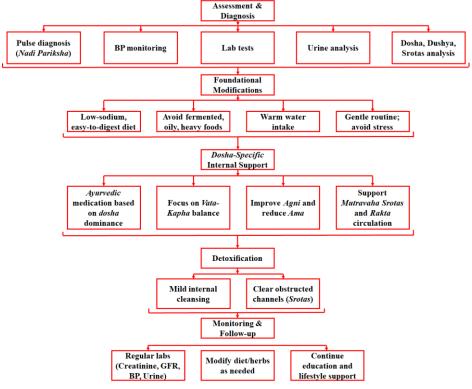
Parameter	06-01-2025		
Naadi (Pulse)	Vataj Pittaj		
Mala (Stool)	Malsang (Constipated)		
Mutra (Urine)	Avikrit (Normal)		
Jiwha (Tongue)	Saam (Coated)		
Shabda (Voice)	Spashta (Clear)		
Sparsha (Touch)	Anushna sheeta (Normal)		
Drika (Eye)	Avikrit (Normal)		
Akriti (Physique)	Madhyam		

Table 2: The vitals during the initial assessment.

Date	<b>Blood Pressure</b>	Pulse Rate/ min	Weight
06-01-2025	152/104 mmHg	102/min	60 Kg
10-02-2025	140/90 mmHg	72/min	58 Kg
18-04-2025	180/110 mmHg	64/min	62 Kg
17-05-2025	140/100 mmHg	80/min	61 Kg
06-06-2025	120/90 mmHg	61/min	61 Kg

Table 3: The laboratory investigation reports during the treatment (Fig 2).

Parameter	Haemoglobin	Urea	BUN	<b>Serum Creatinine</b>	Uric Acid
06-01-2025	11.7 gm/dL	41.50 mg/dL	19.37 mg/dL	1.50 mg/dL	7.42 mg/dL
13-01-2025	-	30.4 mg/dL↓	14.19 mg/dL↓	1.48 mg/dL↓	6.79 mg/dL↓
10-02-2025	11.6 gm/dL↓	50.36 mg/dL↑	23.50 mg/dL↑	2.11 mg/dL↑	8.46 mg/dL↑
18-02-2025	10.7 gm/dL↓	59.93 mg/dL↑	<b>27.97 mg/dL</b> ↑	1.52 mg/dL↓	5.29 mg/dL ↓
17-05-2025	10.7 gm/dL↓	35.41mg/dL ↓	16.53 mg/dL ↓	1.3 mg/dL ↓	5.47 mg/dL↑
06-06-2025	11.7 gm/dL↑	34.45 mg/dL↓	16.08 mg/dL ↓	1.40 mg/dL↑	5.61 mg/dL↑



II Treatment Plan (Fig 3).

A personalized *Ayurvedic* and Disciplined and Intelligent Person's (DIP) Diet was provided to the patient to

complement the Ayurvedic treatments administered at Jeena Sikho Lifecare Limited Hospital for  $CKD^{[16]}$ :

### • Foods to Avoid:

- · Avoid wheat, refined food, milk and dairy products, coffee, tea, and packed food.
- Do not eat after 8 PM
- Eat solid food in small bites, chewing 32 times for proper digestion.

#### · Hydration:

- Consume 1 litre of alkaline water 3-4 times a day.
- Incorporate herbal tea, living water, and turmeric-infused water into your routine.
- Boil 2 litres of water and reduce it to 1 litre for daily consumption.

#### • Millet Intake:

- Include five types of millet: Foxtail, Barnyard, Little, Kodo, and Browntop.
- Use steel cookware for millet preparation and cook with mustard oil.
- Fasting:

DIET

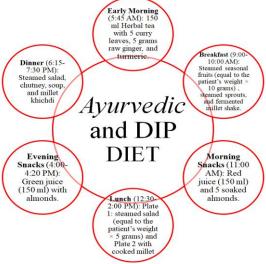
PLA:N

- · One day of fasting is recommended.
- Special Instructions:
- Express gratitude before meals.
- · Practice Vairasana after meals.
- · Take a 10-minute slow walk after meals

Fig 4. Diet Plan.



Fig. 5: Lifestyle Recommendations.



III. *Panchakarma* therapies. Fig 6 Meal Timing and Structure.

# i. Awagah Swedan<sup>[19]</sup>

- A tub was filled with warm water (37–40°C), ensuring the level reached up to the navel.
- The patient was made to sit comfortably in the tub, immersing the body up to the navel region.
- The procedure was continued for 40 minutes.
- After completion, the patient was helped out of the tub and dried with a clean towel.
- The patient was allowed to rest in a warm, closed room.

# ii. Punarnava Gokshur Tail Basti<sup>[20]</sup>

- 80 ml of *Punarnava Gokshur Tail* was warmed by double boiling method to a suitable lukewarm temperature (37-40°C).
- The patient was made to lie down in left lateral position.
- The lubricated nozzle was gently inserted into the rectum.
- The warmed Punarnava Gokshur Tail was administered slowly into the rectum.

- The patient was instructed to retain the oil for as long as comfortably possible.
- The patient was advised to rest in a warm, comfortable room after the procedure.

# iii. Shirodhara with Brahmi oil<sup>[21]</sup>

- The patient was made to lie comfortably in a supine position on the *Droni* (treatment table).
- The forehead and scalp were gently massaged with Brahmi oil.
- Brahmi oil was warmed to a suitable lukewarm temperature (37-40°C).
- The oil was poured in a continuous, steady stream over the center of the forehead (*Ajna Chakra* region) using a *Shirodhara* vessel.
- The procedure was continued for 30 minutes, maintaining a constant oil flow and temperature.
- After completion, excess oil was wiped off, and the patient was allowed to rest for a few minutes.
- The patient was advised to avoid exposure to cold and wind and was given appropriate dietary and lifestyle instructions.

# iv. Guduchi Taila Basti (90 ml)[22]

The patient was kept in left lateral position; *Guduchi Taila* (90 ml) was warmed to body temperature.

- The anal orifice was lubricated, and the warmed oil was administered slowly using a *Basti* syringe with catheter, which was then withdrawn carefully.
- The patient was advised to lie in supine position for 5–10 minutes, and retention of oil was observed under supervision.
- Post-procedure, the patient was instructed to avoid exertion, cold exposure, and suppression of urges, and was advised light, warm, easily digestible food; the procedure was well tolerated.

#### IV. Medicinal Interventions

### a) Avurvedic interventions

The *Ayurvedic* treatment employed in this case CKD Tablet, Yakrit Shoth Har Vati, Chander Vati Tablet, Maha Granthi Har Vati, Dr. BP Tablet, Dr. Kidney Care, Kidney Shuddhi Ark, Kanchnar Guggul, Kidney Care Syrup, Dhatu Poshak Capsule and Divya Shakti Powder (**Table 4**). The description of the medicines is mentioned in **Table 5**.

### a) Allopathic interventions

The patient was diagnosed with hypertension and was taking Amlodipine 5 mg previously which was discontinued after starting *Ayurvedic* treatment.

Table 4: Medications taken during the treatment period.

Date	Medicines	Dosage with Anupana		
	CKD Tablet	1 TAB TDS ( <i>Adhobhakta</i> with <i>koshna jala-</i> After meal with lukewarm water)		
06-01-2025 to	CKD syrup	15 ml BD (Adhobhakta with sama matra koshna jala- After meal with equal amount of lukewarm water)		
15-01-2025	GFR Powder	A teaspoon BD (Adhobhakta with koshna jala)		
(IPD)	Dr. Immune Tablet	1 TAB BD (Adhobhakta with koshna jala)		
	Dr. BP Tablet	1 TAB BD (Adhobhakta with koshna jala)		
	Divya Shakti Powder	Half a teaspoon HS ( <i>Nishikala</i> with <i>koshna jala</i> - Before bed with lukewarm water)		
	Granthi Har Vati	2 TAB BD (Adhobhakta with koshna jala)		
	Divya Shakti Powder	Half a teaspoon HS (Nishikala with koshna jala)		
15-01-2025	CKD Tablet	1 TAB TDS (Adhobhakta with koshna jala)		
	Kidney Shuddhi Ark	15 ml BD (Adhobhakta with sama matra koshna jala)		
(Discharge)	Dr. Immune Tablet	1 TAB BD (Adhobhakta with koshna jala)		
	Dr. BP Tablet	1 TAB BD (Adhobhakta with koshna jala)		
	CKD Tablet	1 TAB TDS (Adhobhakta with koshna jala)		
10-02-2025	Yakrit Shoth Har Vati	2 TAB BD (Adhobhakta with koshna jala)		
	Chandervati	2 TAB BD (Adhobhakta with koshna jala)		
	Maha Granthi Har Vati	2 TAB BD (Adhobhakta with koshna jala)		
	Dr. BP Tablet	2 TAB BD (Adhobhakta with koshna jala)		
18-04-2025	Kidney Care Tablet	1 TAB TDS (Adhobhakta with koshna jala)		
	Kidney Shuddhi Ark	15 ml BD (Adhobhakta with sama matra koshna jala)		
	Chandervati	2 TAB BD (Adhobhakta with koshna jala)		
	Kanchnar Guggulu	2 TAB BD (Adhobhakta with koshna jala)		
	CKD Tablet	1 TAB TDS (Adhobhakta with koshna jala)		
17-05-2025	Kidney Care Syrup	15 ml BD (Adhobhakta with sama matra koshna jala)		
17-05-2025	Chandervati	2 TAB BD (Adhobhakta with koshna jala)		
	Dhatu Poshak Capsule	1 CAP BD (Adhobhakta with koshna jala)		
	Dr. BP Tablet	2 TAB BD (Adhobhakta with koshna jala)		
	Divya Shakti Powder	Half a teaspoon HS (Nishikala with koshna jala)		
	Chandervati	2 TAB BD (Adhobhakta with koshna jala)		
06-06-2025	Dr. Kidney Care Tablet	1 TAB TDS (Adhobhakta with koshna jala)		
00-00-2025	Dr. BP Tablet	2 TAB BD (Adhobhakta with koshna jala)		
	Dhatu Poshak Capsule	2 CAP BD (Adhobhakta with koshna jala)		
	Kidney Shuddhi Ark	15 ml BD (Adhobhakta with sama matra koshna jala)		

Table 5: The Description of Medications taken during the treatment period.

Medicine Name	Ingredients	Therapeutic Effects
CKD Syrup	<b>Kasani</b> (Cichorium intybus), <b>Gokshur</b> (Tribulus terrestris), <b>Shatavari</b> (Asparagus racemosus), <b>Giloy</b> (Tinospora cordifolia), <b>Sorbitol</b> , and <b>Shuddh Shilajit</b> (Asphaltum punjabianum)	Raktashodhak (Blood purifier), Virechana (Purgative), Mutral (Diuretic), Agnideepan (Digestive stimulant), Rasayana (Rejuvenator), Shoth har (Antinflammatory), Pitta Shaman (Pitta pacifier), Kaphashodhana (Kapha eliminator), Srotoshodhana (Channel cleanser)
GFR Powder	<b>Punarnava</b> (Boerhavia diffusa), <b>Gokshur</b> (Tribulus terrestris), <b>Kaasni</b> (Cichorium intybus), <b>Bhoomi Amla</b> (Phyllanthus niruri), <b>Badi Hard</b> (Terminalia chebula), <b>Makoy</b> (Solanum nigrum) and <b>Apamarg</b> (Achyranthes aspera)	Mutral (Diuretic), Shoth har (Anti-inflammatory), Virechana (Purgation), Raktaprasadana (Blood purifier), Vatanulomana (Vata regulator), Mutravirechana (Urinary purgation), Rasayana (Rejuvenator), Amapachan (Toxin digestant), Kledahara (Moisture remover), Vrikkadoshahara (Kidney toxin eliminator)
Dr. Immune tablet	Kesar (Crocus sativus), Shudh Kuchla (Strychnos nux-vomica), Ashwagandha Ext. (Withania somnifera), Shatawari Ext. (Asparagus racemosus), Pipali (Piper longum), Tulsi (Ocimum sanctum), Laung (Syzygium aromaticum), Choti Elaichi (Elettaria cardamomum), Sonth (Zingiber officinale), Haldi (Curcuma longa), Loh Bhasma (Ferrum), Swaran Makshik Bhasma (Chalcopyrite), Mukta Shukti Bhasma (Pinctada margaritifera)	Ojas Vardhaka (Vitality enhancer), Rasayana (Rejuvenator), Vyadhi Kshamatva (Immunity booster), Shoth har (Antiinflammatory), Raktashodhak (Blood purifier), Deepan (Appetizer), Balya (Strength promoter)
Granthi Har Vati	Kachnar (Bauhinia variegata), Guggul (Commiphora wightii), Amalki (Phyllanthus emblica), Vibhitik (Terminalia bellirica), Haritiki (Terminalia chebula), Shunti (Zingiber officinale), Marich (Piper nigrum), Pippal (Piper longum), Varuna (Crateva religiosa), Sukshamala, Dalchini (Cinnamomum verum), and Tamal Patar (Cinnamomum tamala)	Lekhana (scraping), Stambhana (astringent), Shoth har (anti-inflammatory), Vedanasthapana (analgesic), Kapha-Vata Shaman (pacifying Kapha and Vata doshas).
Chander Vati Tablet	Pashanbhed (Bergenia ciliata), Varun (Crataeva nurvala), Punarnava (Boerhavia diffusa), Gokhru (Tribulus terrestris), Apamarg (Achyranthes aspera), Haldi (Curcuma longa), Charila (Embelia ribes), Kulthi (Dolichos biflorus), Harad (Terminalia chebula), Bhumiawla (Pyrrosia piloselloides), Giloy (Tinospora cordifolia), Shitalchini (Vernonia cinerea), Anantmool (Hemidesmus indicus), Khas (Vetiveria zizanoides), Yab Kshar (Alkaline substance, botanical origin unclear), Muli Kshar (Raphanus sativus), Kalmi Shora (Sodium bicarbonate), Sajji Kshar (Traditional alkaline substance, botanical origin unclear), Shilajit (Asphaltum), Hajral Yahud (Silicon dioxide), Shwet Parpati (Mercury-based preparation in Ayurvedic medicine).	Vata-Pitta Shaman (Dosha pacifier), Raktashodhan (Blood purifier), Vrikk dhara (Kidney tonic), Shoth har (Anti-inflammatory), Mutral (Diuretic)
Yakrit Shoth Har Vati	Punarnava (Boerhavia diffusa), Kalimirch (Piper nigrum), Pippali (Piper longum), Vayavidanga (Embelia ribes), Devdaru (Cedrus deodara), Kutha Haldi (Picrorhiza kurroa), Chitrak (Plumbago zeylanica), Harad (Terminalia chebula), Bahera (Terminalia chebula, Terminalia bellirica), Amla (Emblica officinalis), Danti (Baliospermum montanum), Chavya (Piper chaba), Indra Jon (Taraxacum officinale), Pippla Mool (Piper longum), Motha Kalajira (Nigella sativa), Kayphal (Myrica esculenta), Kutaki (Picrorhiza kurroa), Nisoth (Operculina turpethum), Saunth (Zingiber officinale), Kakd Singhi (Cucumis sativus), Ajwain (Trachyspermum ammi), Mandur Bhasma (Ferrum).	Raktashodhak ( Blood purifier). Deepan (Appetizer), Pachan (Digestant), Shoth har (Anti- inflammatory), Vata-kapha shamak ( Dosha-balancer), Rasayana (Rejuvenator), Ojovardhak (Immunity enhancer)

Chander Vati	Kapoor Kachri (Hedychium spicatum), Vacha (Acorus calamus), Motha (Cyperus rotundus), Kalmegh (Andrographis paniculata), Giloy (Tinospora cordifolia), Devdaru (Cedrus deodara), Desi Haldi (Curcuma longa), Atees (Aconitum heterophyllum), Daru Haldi (Berberis aristata), Pipla Mool (Piper longum root), Chitrak (Plumbago zeylanica), Dhaniya (Coriandrum sativum), Harad (Terminalia chebula), Bahera (Terminalia bellirica), Amla (Phyllanthus emblica), Chavya (Piper chaba), Vayavidang (Embelia ribes), Pippal (Piper longum), Kalimirch (Piper nigrum), Saunth (Zingiber officinale dried ginger), Gaj Pipal (Scindapsus officinalis), Swarn Makshik Bhasm (Gold iron pyrite ash - Ayurvedic preparation), Sajjikshar (Potassium carbonate - traditional alkali preparation), Sendha Namak (Rock salt), Kala Namak (Black salt), Choti Elaichi (Elettaria cardamomum - small cardamom), Dalchini (Cinnamomum verum), Tejpatra (Cinnamomum tamala), Danti (Baliospermum montanum), Nishothra (Operculina turpethum), Vanslochan (Bamboo silica), Loh Bhasm (Iron ash - Ayurvedic preparation), Shilajeet (Asphaltum punjabinum), Guggul (Commiphora wightii).	Raktashodhan (Blood purifier), Pitta Shaman (Pitta pacifier), Deepan (Appetizer), Pachan (Digestant), Vata-Pitta Shaman (Dosha pacifier)
Maha Granthi Har Vati	Parad Bhasm (Mercury), Gandhak (Sulfur), Vang Bhasm (Zinc), Taabr Bhasm (Copper), Kash Bhasm (Potassium), Hartal Bhasm (Realgar), Nilla Thotha (Copper sulfate), Shankh Bhasm (Conch shell powder), Kodi Bhasm (Cuttlefish bone), Loh Bhasm (Iron), Sonth (Zingiber officinale), Kalimirch (Piper nigrum), Pippal (Piper longum), Harad (Terminalia chebula), Bahera (Terminalia bellirica), Amla (Phyllanthus emblica), Chavya (Piper chaba), Kachur (Curcuma zedoaria), Vayavdanga (Tribulus terrestris), Pippla Mool (Piper longum root), Patha (Cyclea peltata), Hau Ber (Ziziphus mauritiana), Vacha (Acorus calamus), Choti Ilaychi (Elettaria cardamomum), Devdaru (Cedrus deodara), Samundar Namak (Rock salt), Senda Namak (Sendha salt), Sambar Namak (Sambhar salt), Vid Namak (Black salt), Kala Namak (Black salt), Vidari (Pueraria tuberosa).	Granthi/Arbud (Cyst/Tumor), Lekhana (Scraping/Reducing excess tissue), Shoth har (Anti- inflammatory), Raktashodhak (Blood purifier), Vedanasthapana (Pain reliever)
Dr. BP Tablet	Shankhpushpi (Convolvulus pluricaulis), Shatavari (Asparagus racemosus), Ashwagandha (Withania somnifera), Brahmi (Bacopa monnieri), Vacha (Acorus calamus), Sarpagandha (Rauvolfia serpentina), Jeera (Cuminum cyminum), Giloy (Tinospora cordifolia), Malabar Nut (Justicia adhatoda), Jatamansi (Nardostachys jatamansi), Mukta Pishti (Purified Pearl Calcium - CaCO <sub>3</sub> ).	Uttara vata Shaman (Pelvic Vata pacifier), Rakta gata pitta Shaman (Bloodborne Pitta pacifier), Raktashodhana (Blood purifier), Vata-pitta Shaman (Vata-Pitta pacifier), Hridaya rog nivaran (Heart disease reliever), Shoth har (Anti-inflammatory),
	<b>Gokshur</b> (Tribulus terrestris), <b>Apamarg</b> (Achyranthes aspera), <b>Mulethi</b> (Glycyrrhiza glabra), <b>Punarnava</b> (Boerhavia diffusa), <b>Varun Chhal</b> (Crataeva nurvala), <b>Sheetal Chini</b> (Piper cubeba)	Mutral (Diuretic), Shoth har (Anti-inflammatory), Ashmarighna (Lithotriptic), Agnivardhak (Carminative), Rasayana (Rejuvenator), Vatanuloman (Carminative)
Kidney Shuddhi Ark	<b>Punarnava</b> (Boerhavia diffusa), <b>Gokshur</b> (Tribulus terrestris), <b>Varuna</b> (Crataeva nurvala), <b>Bhumyamalaki</b> (Phyllanthus niruri), <b>Ashwagandha</b> (Withania somnifera), <b>Amla</b> (Emblica officinalis), <b>Shatavari</b> (Asparagus racemosus), <b>Turmeric</b> (Curcuma longa), <b>Saffron.</b>	Mutral (Diuretic), Shoth har (Anti-inflammatory), Mutravirechana (Urinary purgation), Raktaprasadana (Blood purifier), Kledahara (Moisture remover), Amapachan (Toxin digestant), Vrikk doshahar (Kidney toxin eliminator), Rasayana (Rejuvenator), Vatanuloman (Vata regulator)
Kanchnar Guggulu	Kachnar Chhal (Bauhinia variegata), Haritaki (Terminalia chebula), Bibhitaki (Terminalia bellirica), Amalaki (Emblica officinalis), Shunthi (Zingiber officinale), Marich (Piper nigrum), Pippali (Piper longum), Varun Chhal (Crataeva nurvala), Ela (Elettaria cardamomum), Dalchini (Cinnamomum zeylanicum), Tejpatra (Cinnamomum tamala), Shuddha Guggulu (Commiphora mukul)	Lekhan (scraping/reducing), Shoth har (anti-inflammatory), Deepan-Pachan (digestive and carminative), Medohar (anti- obesity/lipolytic)

Kidney Care Syrup		Vata-Pitta Shaman (Pacifier of Vata and Pitta doshas), Raktashodhak (Blood purifier), Shoth har (Anti-inflammatory), Mutra Vardhak (Promoter of urine flow), Srotoshodhak (Channel purifier)
Dhatu Poshak Capsule	Chuna Shuddh, Shankh Bhasm, Mukta Shukti, Prawal Pishti, Kapardika and Loh	Dhatuposhak (Tissue nourishing), Rasayana (Rejuvenative), Balya (Strengthening), Srotoshodhak (Channel cleansing), Vata-Pitta shaman (Vata and Pitta balancing), shodhak (Detoxifier), Agni Deepan (Digestive fire stimulant), Lekhana (Scraping / Lipolytic)
Divya Shakti Powder	Trikatu, Triphala, Nagarmotha (Cyperus rotundus), Vay Vidang (Embelia ribes), Chhoti Elaichi (Elettaria cardamomum), Tej Patta (Cinnamomum tamala), Laung (Syzygium aromaticum), Nishoth (Operculina turpethum), Sendha Namak, Dhaniya (Coriandrum sativum), Pipla Mool (Piper longum root), Jeera (Cuminum cyminum), Nagkesar (Mesua ferrea), Amarvati (Achyranthes aspera), Anardana (Punica granatum), Badi Elaichi (Amomum subulatum), Hing (Ferula assafoetida), Kachnar (Bauhinia variegata), Ajmod (Trachyspermum ammi), Sazzikhar, Pushkarmool (Inula racemosa), Mishri (Saccharum officinarum)	Ojakshaya (Loss of vitality/immunity), Agnimandya (Low digestive fire), Chakshukshaya (Weak vision), Deepan (Appetizer), Rasayana (Rejuvenator)

### RESULT

Following four months of treatment, the patient showed significant improvement in symptoms, indicating that the interventions applied were effective in managing CKD and hypertension. Additionally, the reduction in

weakness and swelling in B/L legs further supports the efficacy of the *Ayurvedic* approach used in this case. The conditions during the admission and discharge are mentioned in **Table 6**. The DTPA scan reports are mentioned in **Table 7** (**Fig 7**).

Table 6: The conditions during the admission and discharge.

Conditions Dur		ring Admission		During discharg		
Weakness	Severe		Relief			
Swelling in B/L legs	3°			1°		
Weakness Scale						
Pravar Bala / No / Slight Weakness Weakness		Yuktikrit Bala / Mild Weakness	a / Mild M		Akalaja Bala / Severe Weakness	

Table 7: The DTPA scan reports.

Date	13-01	-2025	07-06-2025		
Kidney	Right Left		Right	Left	
GFR	31.0 ml/min	34.8 ml/min	33.3 ml/min	34.8 ml/min	
Split function	47% 53%		49%	51%	
Global GFR	65.8 n	nl/min	68.1 ml/min		

# **DISCUSSION**

This case report details the integrative *Ayurvedic* management of a 36-year-old male patient diagnosed with CKD and associated hypertension. Upon presentation, the patient exhibited symptoms including generalized weakness and bilateral oedema, which are commonly observed in progressive renal dysfunction. The therapeutic approach involved a combination of

Ayurvedic internal medications and dietary modifications, customized to address the patient's Prakriti (constitution), Doshic imbalance, and disease stage. The underlying Ayurvedic pathogenesis (Samprapti) of the condition, which integrates the concepts of Dosha Dushti, Dhatu Kshaya, and Srotorodh, has been illustrated in Fig. 8, referencing classical Ayurvedic frameworks. [23,24]

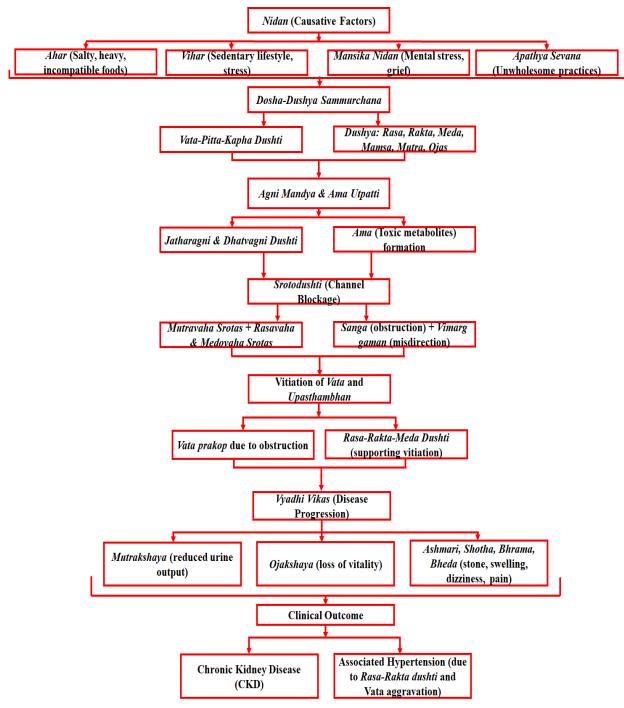


Fig. 8: The samprapti for this study.

### 1. The Samprapti

CKD with Hypertension is a result of *Nidan Sevana* (consumption of causative factors) such as *Ati-Lavana*, *Guru*, *Abhishyandi Ahara*, and *Avyaayama*, *Ratrijagarana*, leading to *Tridosha Prakopa*, predominantly *Vata-Kapha*. [25] This leads to *Agnimandya* (suppression of *Jatharagni* and *Dhatvagni*), causing *Ama Utpatti*. [26] The accumulated *Ama* obstructs the *Mutravaha Srotas*, along with *Rasavaha* and *Medovaha Srotas*, resulting in *Srotorodha*. [27] This *Margavarana* (obstruction) provokes *Vata*, particularly *Apana Vayu*, disturbing its normal *Gati* and causing *Mutrakshaya*, *Mutraghata*, and *Mutradourbalya*. [28]

The continuous obstruction and dosha-dushya sammurchana vitiate Rasa, Rakta, Meda, Mamsa, Mutra, and Ojas, leading to Ojakshaya and Dhatukshaya. [29] The Rakta vaha srotodushti contributes to Uchcha Raktachap (Hypertension) due to increased Rasa-Rakta abaddhata. Clinically, this manifests as Shoth, Shrama, Phenil mutrata, Mutra alpata, Bhrama, Raktapradurbhava. [30] Over time, the disease progresses into a Krchchhra-sadhya vyadhi requiring a Shaman and Rasayana based approach.[31]

### 2. The Nidan

In Ayurveda, Nidan refers to the causative factors responsible for the initiation and progression of disease. [32] In the context of CKD with Hypertension, common Nidanas include Ati-Lavana Sevan (excessive salt intake), guru and Abhojya Ahara (heavy and incompatible food), Ajirna Bhojan (eating with poor digestion), Mandagni (weak digestive fire), Avyaayaam (lack of exercise), Ratrijagarana (night waking), Divaswapa (day sleep), and Manasik Nidan such as Chinta (worry) and Shoka (grief). [33] These factors aggravate Tridoshas, especially Vata and Kapha, impair Agni, and contribute to Ama Utpatti, leading to Srotodushti and eventually resulting in CKD and Hypertension. [34]

Nidan Parivarjan is the foremost principle of treatment in Ayurveda, which emphasizes the elimination or avoidance of these causative factors to break the disease cycle. In this case, it involves adopting a Pathya Ahar-Vihar (wholesome diet and lifestyle), reducing salt and protein intake, avoiding sedentary habits, maintaining regular sleep, and managing stress through Manasik Shuddhi (mental clarity). Nidan Parivarjan not only helps to prevent disease progression but also enhances the effectiveness of therapeutic interventions and supports Dosha Shaman and Dhatu Poshana (balance and nourishment of body tissues).

### 3. The effects of *Panchakarma* therapies

In CKD, Panchakarma therapies such as Awagah Swedana, Punarnava Gokshur Taila Basti, Shirodhara with Brahmi oil, and Guduchi Taila Basti provide multidimensional benefits. Awagah Swedana promotes gentle sweating, enhances circulation, aids detoxification, reduces fluid retention, and alleviates stiffness while providing relaxation. [19] Punarnava Gokshur Taila Basti delivers local anti-inflammatory and diuretic effects, pacifies Vata and Kapha, facilitates elimination of toxins, and supports renal tissue accumulated regeneration.<sup>[20]</sup> Shirodhara with Brahmi oil induces deep relaxation, balances Vata, exerts neuroprotective and antioxidant effects, and helps regulate blood pressure and sympathetic activity, indirectly benefiting renal function. [21] Guduchi Taila Basti acts as a Rasayana and immunomodulator, promotes detoxification, enhances metabolic efficiency, and supports kidney tissue repair. [22] Collectively, these therapies improve fluid balance, reduce inflammation, rejuvenate renal tissues, and enhance overall systemic and mental well-being in CKD patients.

# 4. The effects of Ayurvedic medications

The Ayurvedic management of CKD with hypertension targets multiple levels of the disease Samprapti, aiming to break the pathological chain. Chander Vati Tablet acts as an Agnivardhak and Mutral, helping to correct Agnimandya, digest Ama, and restore the flow through Mutravaha srotas. Yakrit Shoth Har Vati works as a

Shothahara and Raktashodhak, supporting liver function (Yakrituttejaka) and aiding in the removal of metabolic toxins that contribute to Rakta Dushti and hypertension. Maha Granthi Har Vati acts through Lekhan and Kapha-Shaman, reducing Granthi (fibrotic/cystic formations) and *Meda Dushti*, thereby supporting kidney structure. Dr. BP Tablet helps to control hypertension through Vatanuloman, Raktashodhan, and Hridaya Balya properties, pacifying Vyana Vayu and stabilizing Rakta Dhatu. Dr. Kidney Care offers a broad-spectrum effect with Mutral, Shoth-har, Ashmarighna, and Rasayana actions, targeting Srotorodh, oedema, urinary flow, and *Dhatu* rejuvenation. Kidney Shuddhi Ark supports detoxification and corrects Mutravaha Srotodushti, while Kanchnar Guggul aids in reducing glandular swelling and Meda Granthi, acting as a Lekhan and Shoth-har. Kidney Care Syrup supports urine flow, soothes Pitta, and balances Vata-Kapha. Dhatu Poshak Capsule acts as a Rasayana and Balya, promoting Dhatu Pushti and addressing Ojakshaya, while Divya Shakti Powder supports overall systemic balance, Agni Deepana, and Dhatu nourishment. CKD Syrup provides Raktashodhak, Mutral, Virechana, Shoth-har, Pitta pacifying, Kapha eliminating, and Srotoshodhana actions. GFR Powder aids diuresis, anti-inflammation, urinary purgation, Vata regulation, detoxification, and renal rejuvenation. Dr. Immune Tablet enhances Ojas, immunity, strength, digestion, blood purification, and Rasayana effects, while Granthi Har Vati contributes Lekhana, Stambhana, Shoth-har, analgesic, and Kapha-Vata pacifying actions. Together, these formulations act synergistically to restore Agni, eliminate Ama, correct Srotodushti, pacify Vata-Pitta, rejuvenate affected Dhatus, improve renal structure and function, and effectively break the Samprapti of CKD with hypertension.

The Rasa Panchak of key Ayurvedic herbs used in CKD with hypertension reflects their potent therapeutic action. Most, like Punarnava<sup>[36]</sup>, Gokshur<sup>[37]</sup>, and Guduchi<sup>[38]</sup>, are Tikta (bitter) and Kashaya (astringent) in Rasa, with Laghu (light) and Ruksha (dry) Guna, Sheet (cool) or Ushna (hot) Veerya, and Katu Vipak. These properties help reduce Kapha and Pitta, promote diuresis, detoxification, and support renal tissue repair. Their Prabhava (specific action) includes Mutral (diuretic) and Shoth-har (anti-inflammatory) effects, aligning with the Ayurvedic approach to managing kidney dysfunction and associated hypertension, which is mentioned in **Table 8**.

Table 8: The Ras Panchaka of the common key ingredients.

anchaka or	anchaka of the common key ingredients.						
Ingredient	Rasa (Taste)	Guna (Qualities)	Veerya (Potency)	Vipaka (Post- digestive Effect)	Prabhava (Specific Action)	Present In	
<b>Punarnava</b> (Boerhavia diffusa)	Tikta, Kashaya (bitter, astringent)	Laghu , Ruksha (light, dry)	Ushna (hot)	Katu (pungent)	Mutral , Shoth har	Chander Vati, Yakrit Shoth Har Vati	
<b>Palash</b> (Butea monosperma)	Tikta , Kashaya (bitter, astringent)	Laghu , Ruksha (light, dry)	Ushna (hot)	Katu (pungent)	Shoth har , Mutral	Dr. Kidney Care, Divya Shakti Powder	
<b>Daruharidra</b> (Berberis aristata)	Tikta, Kashaya (bitter, astringent)	Laghu , Ruksha (light, dry)	Ushna (hot)	Katu (pungent)	Krimighna , Raktashodhak	Chander Vati, Yakrit Shoth Har Vati	
<b>Guduchi</b> (Tinospora cordifolia)	Tikta (bitter)	Laghu, Snigdha (light, unctuous)	Ushna (hot)	Madhur (sweet)	Rasayana , Tridoshaghna	Chander Vati , Dr. Kidney Care	
<b>Haridra</b> (Curcuma longa)	Tikta , Katu (bitter, pungent)	Laghu , Ruksha (light, dry)	Ushna (hot)	Katu (pungent)	Shoth har , Raktashodhak	Chander Vati , Yakrit Shoth Har Vati	
<b>Shilajit</b> (Asphaltum)	Katu, Tikta (pungent, bitter)	Laghu , Ruksha (light, dry)	Ushna (hot)	Katu (pungent)	Yogavahi , Rasayana , Vrishya	Chander Vati, Dr. Kidney Care	



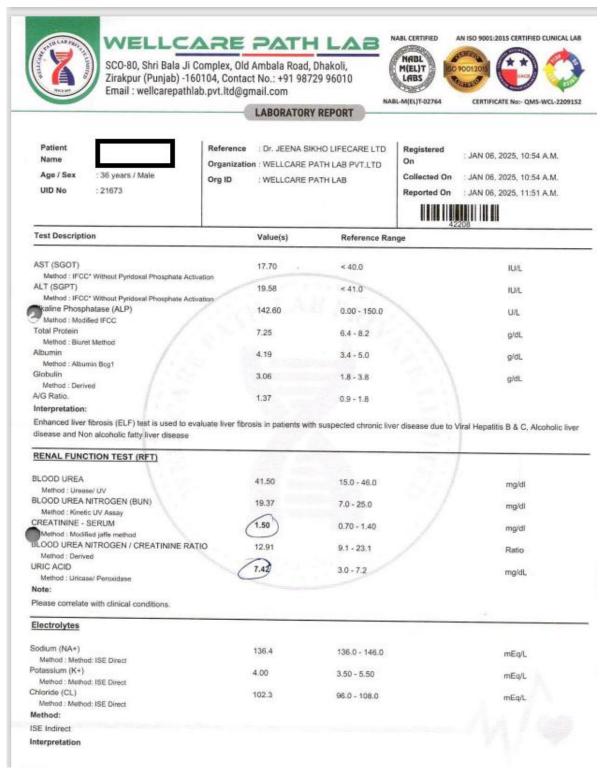


Fig 2: The laboratory investigation reports before and after treatment.



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CERTIFICATE No:- OMS-WCL-2209152



# LABORATORY REPORT

Patient Name Age / Sex 36 years / Mai UID No 21673

: Dr. JEENA SIKHO LIFECARE LTD Organization: WELLCARE PATH LAB PVT.LTD Org ID

: WELLCARE PATH LAB

Registered

JAN 13, 2025, 09:21 A.M.

: JAN 13, 2025, 09:21 A.M. Collected On Reported On : JAN 13, 2025, 10:08 A.M.

Test Description	Value(s)	Reference Range					
RENAL FUNCTION TEST (RFT)							
BLOOD UREA Method : Urease/ UV	30.40	15.0 - 46.0	mg/dl				
DOD UREA NITROGEN (BUN)	14.19	7.0 - 25.0	mg/dl				
Method : Kinetic UV Assay			50.000				
CREATININE - SERUM  Method : Modified jaffe method	1.48	0.70 - 1.40	mg/dl				
BLOOD UREA NITROGEN / CREATININE RATIO Method : Derived	9.59	9.1 - 23.1	Ratio				
URIC ACID	6.79	3.0 - 7.2	mg/dL				
Method : Uricase/ Peroxidase		100	rigide				
Note:							
Please correlate with clinical conditions.							
Electrolytes	ALF V						
Sodium (NA+) Method : Method: ISE Direct	139.3	136.0 - 146.0	mEq/L				
Potassium (K+) Method: ISE Direct	3.74	3,50 - 5,50	mEq/L				
Chloride (CL) Method: Method: ISE Direct	103.2	96.0 - 108.0	mEq/L				
Method:							
ISE Indirect							
erpretation							

Sodium measurements are used in the diagnosis and treatment of aldosteronism (excessive secretion of the hormone aldosterone), diabetes insipidus (chronic excretion of large amounts of dilute urine, accompanied by extreme thirst), adrenal hypertension. Addison's disease (caused by destruction of the adrenal glands), dehydration, inappropriate antidiuretic hormone secretion, or other diseases involving electrolyte imbalance. Potassium measurements are used to monitor electrolyte balance in the diagnosis and treatment of disease conditions characterized by low or high blood potassium levels. Chloride measurements are used in the diagnosis and treatment of electrolyte and metabolic disorders such as cystic fibrosis and diabetic acidosis

"END OF REPORT"

Dr. Ankit Aggarwal (Consultant Pathologist)

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neptring result are for the information and for interpretation of the referring doctor only. • If the result of the test (n) are alarming or unexpected, the patient is advised to contact the laboratory immediately for soldie remedial advice. • This reports is not valid for medico-legal purposes. • Wellcare Clinical Lab out its employees assume any liability to for any loss or damage that may be incurred by any person as a result of presuming meaning or contents of the report. • It is Presumed that the tests performed on the specimen belong to the patient; names or identified. • Results of tests may vary from laboratory and also in some parameter from et to firm for the same patient. Only such medical professional who enderstand reporting units, reference assigned and initiations or technologies should interpret result. • Reports valid unit stamped by labs authorized signatory.

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NABL-M(EL)T-02764

QMS-WCL-2209152

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### LABORATORY REPORT

Patient Name

Age / Sex

UID No : 23539

: 36 years / Male

Reference : Dr. JEENA SIKHO LIFECARE LTD

Organization: WELLCARE PATH LAB PVT.LTD

: WELLCARE PATH LAB

Registered

On

: FEB 10, 2025, 01:21 P.M.

Collected On : FEB 10, 2025, 01:21 P.M. Reported On : FEB 10, 2025, 02:12 P.M.

Test Description	Value(s)	Reference Range	
Complete Blood Count(CBC)			
Hemoglobin (HB)  Method : Cynmeth Photometric Measurement	11.6	13.0 - 17.0	g/dL
Total Leucocytes Count (TLC)	10800	4000 - 11000	/cmm
Method : Electrical Impedance	10000	4000 - 11000	/cmm
Metriod : Electrical Impedance			
DIFFERENTIAL COUNT			
Neutrophils	59	40 - 75	%
Method : VCSn Technology			
Lymphocytes	32	20 - 45	%
Method : VCSn Technology		The state of the s	
Monocytes	05	2 - 10	%
Method : VCSn Technology			
Eosinophils	04	1 - 6	%
Method : VCSn Technology			00 00 00 00 00
Basophils	00	0 - 1	%
Total RBC Count	4.36	3.50 - 6.50	Mill/Cumm
Method : Electrical Impedance			
Platelet Count	3.02	1.50 - 4.50	Lacs/Cumm
Method : VCSn Technology			
PCV/HCT	37.9	35.0 - 47.0	%
Method : Calculated			
Red cell distribution width (RDW)	14.5	13.0 - 18.0	%
Method : Electrical Impedance			
ean corpuscular volume (MCV)	87.0	76.0 - 96.0	fl
Method : Electrical Impedance		1045	
Mean Corpuscular Hemoglobin (MCH)  Method : Calculated	26.6	27.0 - 32.0	pg
Mean Corpuscular Hemoglobin Concentration(MCHC)	30.5	30.0 - 35.0	%
Method : Calculated	7		

Microscopy, Fully Automated Hematology Analyser alfa swelab double chamber 3 Part

RENAL FUNCTION	N TEST (RFT)
----------------	--------------

BLOOD UREA	50.36	15.0 - 46.0	mg/dl
Method : Urease/ UV			
BLOOD UREA NITROGEN (BUN)	23.50	7.0 - 25.0	mg/dl
Method : Kinetic UV Assay			
CREATININE - SERUM	2.11	0.70 - 1.40	mg/dl
Method: Modified jaffe method			

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## LABORATORY REPORT

Patient Name

UID No

Age / Sex : 36 years / Male

: 23539

: Dr. JEENA SIKHO LIFECARE I TD Reference Organization: WELLCARE PATH LAB PVT.LTD

: WELLCARE PATH LAB

Registered : FEB 10, 2025, 01:21 P.M. On

Collected On : FEB 10, 2025, 01:21 P.M. Reported On : FEB 10, 2025, 02:12 P.M.

			45553
Test Description	Value(s)	Reference Range	
BLOOD UREA NITROGEN / CREATININE RATIO Method : Derived	11.14	9.1 - 23.1	Ratio
URIC ACID	8.46	3.0 - 7.2	mg/dL
Method : Uricase/ Peroxidase			3
Note:	CARL FOR MARKET		
Please correlate with clinical conditions.			
. icase correlate with clinical cortainors.			
Electrolytes	A 1		
Sodium (NA+)	136.2	136.0 - 146.0	mEg/L
Method : Method: ISE Direct		100.0	meq/E
Potassium (K+)	4.47	3.50 - 5.50	mEg/L
Method : Method: ISE Direct			meqre
Chloride (CL)	105.2	96.0 - 108.0	mEq/L
Method : Method: ISE Direct			meq/E
Method:			9
ISE Indirect			
			1

# Interpretation

Sodium measurements are used in the diagnosis and treatment of aldosteronism (excessive secretion of the hormone aldosterone), diabetes insipidus (chronic excretion of large amounts of dilute urine, accompanied by extreme thirst), adrenal hypertension, Addison's disease (caused by destruction of the adrenal glands), dehydration, inappropriate antidiuretic hormone secretion, or other diseases involving electrolyte imbalance. Potassium measurements are used to monitor electrolyte balance in the diagnosis and treatment of disease conditions characterized by low or high blood potassium levels. Chloride measurements are used in the diagnosis and treatment of electrolyte and metabolic disorders such as cystic fibrosis and diabetic acidosis

# C/E Complete Urine Examination

# URINE ROUTINE AND MICROSCOPIC EXAMINATION

# PHYSICAL EXAMINATION:

Colour of Urine	Pale Yellow	Straw to Yellow	/HPF
Visually Appearance	Clear	Expected Clear	/HPF
Reaction (pH)	Acidic 5.5	5.0 - 8.0	/HPF
Specific Gravity	1.010	1.000 - 1.030	/HPF
Protein	Absent	Expected Absent	/HPF
Glucose	Absent	Expected Absent	/HPF
MICROSCOPIC EXAMINATION			
Pus Cells	1 - 2	0 - 2	/HPF
Epithelial Cells	2 - 3	Expected Absent	/HPF
Red Blood Cells (RBC).	Absent	Expected Absent	/HPF
Casts	Absent	Expected Absent	/HPF

### **CONDITIONS OF LABORATORY TESTING & REPORTING**

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# LABORATORY REPORT

Patient Name

Age / Sex 36 years / Male

**UHID No** : 1052025 Reference : Dr. JEENA SIKHO LIFECARE LTD Organization: WELLCARE PATH LAB PVT.LTD

: WELLCARE PATH LAB

Registered

On

: JUN 06, 2025, 11:34 A.M.

Collected On : JUN 06, 2025, 11:34 A.M. Reported On : JUN 06, 2025, 01:19 P.M.

est Description	Value(s)	Reference Range	
Complete Blood Count(CBC)			
Hemoglobin (HB)	11.7	13.0 - 17.0	g/dL
Method : Cynmeth Photometric Measurement			4577507
Total Leucocytes Count (TLC)	8500	4000 - 11000	/cmm
lethod : Electrical Impedance			
DIFFERENTIAL COUNT			
Veutrophils	68	40 - 75	%
Method : VCSn Technology			
ymphocytes	26	20 - 45	%
Method : VCSn Technology			1
Monocytes	04	2 - 10	%
Method : VCSn Technology			
Eosinophils	02	1 - 6	%
Method : VCSn Technology			
Basophils	00	0 - 1	%
otal RBC Count	3.97	3.50 - 6.50	Mill/Cumm
Method : Electrical Impedance			
Platelet Count	2.02	1.50 - 4.50	Lacs/Cumm
Method : VCSn Technology			/
PCV/HCT	37.4	35.0 - 47.0	%
Method : Calculated			
Red cell distribution width (RDW)	12.9	13.0 - 18.0	%
Method : Electrical Impedance			
lean corpuscular volume (MCV)	94.1	76.0 - 96.0	fl
lethod : Electrical Impedance		and the second	
Mean Corpuscular Hemoglobin (MCH)	29.5	27.0 - 32.0	pg
Method : Calculated		9 7	
Mean Corpuscular Hemoglobin Concentration(MCHC)	31.4	30.0 - 35.0	%

Microscopy, Fully Automated Hematology Analyser alfa swelab double chamber 3 Part

# RENAL FUNCTION TEST (RFT)

BLOOD UREA	34.45	15.0 - 46.0	mg/dl
Method : Urease/ UV			
BLOOD UREA NITROGEN (BUN)	16.08	7.0 - 25.0	mg/dl
Method : Kinetic UV Assay			
CREATININE - SERUM	1.40	0.70 - 1.40	mg/dl
Method : Modified jaffe method			

### **CONDITIONS OF LABORATORY TESTING & REPORTING**

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The reporting result are for the information and for interpretation of the referring doctor only. • If the result of the test (s) are alarming or unexpected, the patient is advised to contact the laboratory immediately for possible remedial advice. • This reports is not valid for medico-legal purposes. • Wellcare Path Lab not its employees assume any liability to for any loss or damage that may be incurred by any person as a result of presuming the meaning or contents of the report. • It is Presumed that the tests performed on the specimen belong to the patient: names or identified. • Results of tests may vary from laboratory to laboratory and also in some parameter from time to time for the same patient. Only such medical professional who understand reporting units, reference ranges and limitations or technologies should interpret result. • Reports valid until stamped by labs authorized signatory.

NOT VALID FOR MEDICO LEGAL PURPOSE | EMERGENCY 24 HOURS | TIMINGS : 8.00 AM TO 8.00 PM

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SCO-80, Shri Bala Ji Complex, Old Ambala Road, Dhakoli, Zirakpur (Pb) -160104, Contact No.: +91 98729 96010 Email : wellcarepathlab.pvt.ltd@gmail.com





CERTIFICATE No.: QMS-WCL-2209152

g/dL

We are enrolled with CMC EQAS & AIIMS EQAS External Quality Assurance, We are running CMC EQAS & AIIMS Quality Controls daily a day

# LABORATORY REPORT

Patient
Name
Age / Sex : 36 years / Male
UHID No : 1052025

Reference : Dr. JEENA SIKHO LIFECARE LTD

Organization : WELLCARE PATH LAB PVT.LTD

Org ID : WELLCARE PATH LAB

Registered : JUN 06, 2025, 11:34 A.M.

Collected On : JUN 06, 2025, 11:34 A.M.

Reported On : JUN 06, 2025, 01:19 P.M.

Test Description	Value(s)	Reference Range	
BLOOD UREA NITROGEN / CREATININE RATIO Method : Derived	11.49	9.1 - 23.1	Ratio
URIC ACID	5.61	3.0 - 7.2	mg/dL
Method : Uricase/ Peroxidase			mgrac
Note:		AR BEING STORY	
se correlate with clinical conditions.			
Liver Function Test (LFT)			
Total Bilirubin	0.73	0.20 - 1.00	mg/dL
Method : Vanadate : oxidation			3
Direct Bilirubin	0.35	0.00 - 0.60	mg/dL
Method : Vanadate : oxidation			
Indirect Bilirubin	0.38	0.00 - 0.80	mg/dL
Method : Derived			
AST (SGOT)	24.26	< 40.0	IU/L
Method : IFCC* Without Pyridoxal Phosphate Activation		10.0	
ALT (SGPT)	17.56	< 41.0	IU/L
Method : IFCC* Without Pyridoxal Phosphate Activation			
Alkaline Phosphatase (ALP)	109.81	0.00 - 150.0	U/L
Method : Modified IFCC			
Total Protein	6.86	6.4 - 8.2	g/dL
Method : Biuret Method			3
Albumin	3.95	3.4 - 5.0	g/dL
Method : Albumin Bcg1		. /	<b>3</b>

A/G Ratio.
Interpretation:

ISE Indirect

ulin

ethod : Derived

Enhanced liver fibrosis (ELF) test is used to evaluate liver fibrosis in patients with suspected chronic liver disease due to Viral Hepatitis B & C, Alcoholic liver disease and Non alcoholic fatty liver disease

1.8 - 3.8

0.9 - 1.8

Electrolytes			
Sodium (NA+)	140.0	136.0 - 146.0	mEq/L
Method : Method: ISE Direct			(1000 mm)
Potassium (K+)	4.42	3.50 - 5.50	mEq/L
Method : Method: ISE Direct			1
Chloride (CL)	104.0	96.0 - 108.0	mEa/L
Method : Method: ISE Direct			4
Method:			

2.91

1 36

CONDITIONS OF LABORATORY TESTING & REPORTING

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NOT VALID FOR MEDICO LEGAL PURPOSE | EMERGENCY 24 HOURS | TIMINGS : 8.00 AM TO 8.00 PM

Fig 7 The DTPA scan

AERB Reg. No. 21-NMLICENSE-639460



NAME

REG. N

# INDIAN INSTITUTE OF NUCLEAR MEDICINE & SCANNING

(A Unit of Pan Rock Management India Pvt. Ltd.)

SEX: M DATE: 13/01/2025

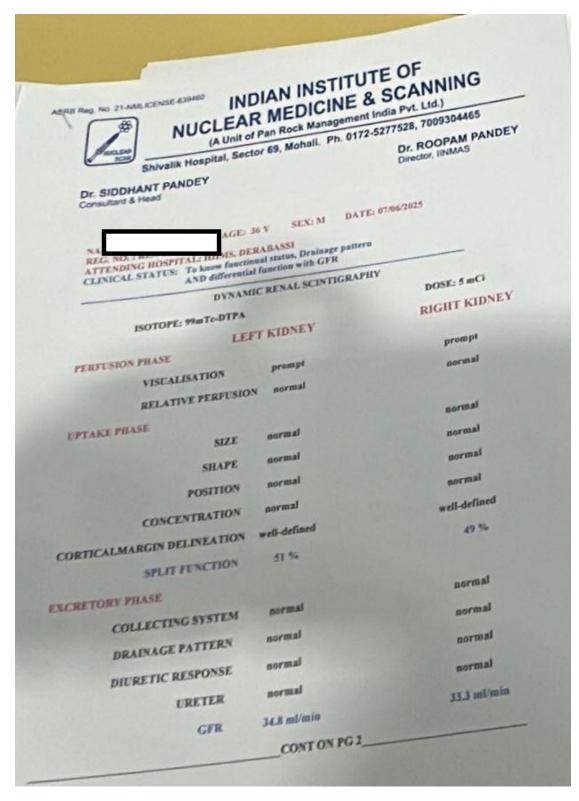
Shivalik Hospital, Sector 69, Mohali. Ph. 0172-5277528, 7009304465

AGE: 36 Y

Dr. SIDDHANT PANDEY Consultant & Head Dr. ROOPAM PANDEY, MD Director, IINMAS

ATTENDING HOSPITAL: HIIMS, DERABASSI CLINICAL STATUS: To know functional status, Drainage pattern AND differential function with GFR DYNAMIC RENAL SCINTIGRAPHY ISOTOPE: 99mTc-DTPA DOSE: 5 mCi LEFT KIDNEY RIGHT KIDNEY PERFUSION PHASE VISUALISATION prompt prompt RELATIVE PERFUSION normal normal UPTAKE PHASE SIZE normal normal. SHAPE normal normal POSITION normal CONCENTRATION normal normal well-defined CORTICALMARGIN DELINEATION well-defined SPLIT FUNCTION 53 % 47 % EXCRETORY PHASE normal COLLECTING SYSTEM normal DRAINAGE PATTERN normal normal DIURETIC RESPONSE normal normal URETER normal normal 34.8 ml/min 31.0 ml/min CONT ON PG 2

NOT VALID FOR MEDICO-LEGAL PURPOSE



### 5. The effects of Ahar-Vihar

The *Ayurvedic* and DIP dietary guidelines offers a practical framework for interrupting this chain of events. A millet-based diet comprising five types, Foxtail, Kodo, Little, Browntop, and Barnyard millets, serves as a key dietary intervention. <sup>[39]</sup> These millets are *Laghu* (light to digest), *Tridoshahar* (balancing to all three *Doshas*), and naturally low in sodium and phosphorus, thus reducing the metabolic load on the kidneys. The regular

consumption of steamed salads, seasonal fruits, fermented millet shakes, and sprouts promotes *Deepan* and *Pachan* (enhancement of digestion and metabolism), thereby strengthening the digestive fire (*Agni*) and reducing the formation of *Ama*. [40] Excluding aggravating foods such as wheat, refined food, milk and dairy products, coffee, tea, and processed or packed items prevents further *Kapha* and *Meda Dushti*, which are known to obstruct *Srotas* and increase toxicity. [41]

Including herbal teas made from curry leaves, raw ginger, and turmeric offers *Shoth har* (anti-inflammatory), *Mutral* (diuretic), and *Rasayana* (rejuvenative) benefits, which are critical for vascular and renal health. [42] Hydration through 1 liter of alkaline water consumed 3–4 times a day, as well as turmeric-infused water, boiled water reduced to half its volume, and living water, helps balance the body's pH, reduce systemic inflammation, and flush out toxins, thereby supporting kidney function. [43]

The lifestyle component (Vihar) of this regimen emphasizes the importance of daily yoga, particularly Sukhasan and Sukshm Pranayaam, which pacify Vata, especially Vyana Vayu, and enhance the parasympathetic response, contributing to better blood pressure control. [44] Mindful practices such as expressing gratitude before meals improve Manasik Bhava (mental state), while chewing food 32 times ensures proper digestion and assimilation. [45] The recommendation to eat before 8 PM aligns with the body's circadian rhythm and promotes optimal digestion. [46] Performing Vajrasan after meals and taking a 10-minute slow walk further supports digestion and prevents postprandial blood sugar spikes. [47] One day of fasting per week serves as Langhan therapy, allowing the digestive system to rest, enhancing Agni, reducing Ama, and clearing Srotorodh through internal detoxification. [48]

From a physicochemical standpoint, this approach also addresses key factors associated with CKD and hypertension. It helps lower oxidative stress, corrects chronic low-grade metabolic acidosis, improves endothelial function, and downregulates proinflammatory cytokines. [49] Millets provide a low-glycemic, high-fiber profile that regulates insulin levels and supports cardiovascular health. [50] Alkaline hydration and herbal infusions support electrolyte balance and kidney filtration. [51] Cooking millets in steel cookware with mustard oil adds further anti-inflammatory and circulatory benefits.

# FUTURE RESEARCH PERSPECTIVES

This study was conducted on a 36-year-old male patient with CKD and hypertension. While the findings were encouraging, the conclusions are limited due to the single-patient design. To strengthen the evidence base, future research must include larger sample sizes through well-structured, randomized controlled trials. These studies should aim to evaluate the long-term efficacy, safety, and reproducibility of the *Ayurvedic* therapies and lifestyle modifications used. Establishing standardized clinical protocols through such research will be essential for incorporating these interventions into broader clinical practice and enhancing patient outcomes in CKD with hypertension.

### CONCLUSION

The following conclusions can be drawn from this case study on treating CKD with hypertension using Ayurvedic treatments with previously prescribed allopathic medications:

**Symptoms:** At the first visit, the patient presented with symptoms such as weakness and swelling in B/L legs. After the treatments followed by *Ayurvedic* care, the patient showed significant improvement. The conditions and symptoms were reduced, and no new symptoms were reported, indicating notable improvement in kidney function and overall well-being.

**Vitals:** The patient's vital parameters revealed a marked transition from hypertensive to more stable readings over time. On 18 April, 2025, the highest blood pressure recorded was 180/110 mmHg, indicating severe hypertension. By 06 June, 2025, the blood pressure had normalized to 120/90 mmHg, suggesting improved cardiovascular control. These findings reflect positive outcomes in managing hypertension and autonomic balance during the course of treatment.

Investigations: The case study demonstrated notable improvements in renal function and key biochemical parameters in a 36-year-old male patient with chronic kidney disease (CKD) and hypertension following Ayurvedic interventions. The GFR of the right kidney improved from 31.0 ml/min on 13-01-2025 to 33.3 ml/min on 07-06-2025, while the left kidney GFR remained stable at 34.8 ml/min. Consequently, the global GFR increased from 65.8 ml/min to 68.1 ml/min, indicating enhanced overall kidney function. Biochemical parameters also reflected clinical improvement. Serum creatinine reduced significantly from 2.11 mg/dL to 1.40 mg/dL. Urea levels peaked at 59.93 mg/dL but declined steadily to 34.45 mg/dL, and blood urea nitrogen (BUN) dropped from 27.97 mg/dL to 16.08 mg/dL. Additionally, hemoglobin improved from 10.7 gm/dL to 11.7 gm/dL, and uric acid stabilized within manageable limits. These findings collectively suggest a positive trend in renal function and systemic biochemical balance with Ayurvedic management.

The study concludes that combining *Ayurvedic* therapies for CKD led to beneficial outcomes, such as reduced symptoms, improved vital parameters, and enhanced lab results.

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