

## Dravyaguna Principles As Predictive Tools In Ayurvedic Pharmacology: A Review

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### ABSTRACT

Ayurveda, the ancient Indian system of medicine, offers a comprehensive pharmacological framework grounded in natural principles that emphasize the interplay between the properties of substances and their effects on the human body. The branch of Dravyaguna Vijnana (Ayurvedic pharmacology) encompasses the detailed study of Rasa (taste), Guna (qualities), Veerya (potency), Vipaka (post-digestive transformation), and Prabhava (specific action). These principles not only classify substances but also serve as predictive tools for assessing their therapeutic potential, even in the absence of experimental validation. Unlike modern pharmacology, which often focuses on isolated chemical constituents and receptor-specific actions, the Ayurvedic model is holistic, integrative, and tailored to individual constitution (Prakriti). This review critically explores each of the Panchavidha Gunas—highlighting their theoretical foundations, clinical significance, and potential correlations with modern pharmacodynamic and pharmacokinetic concepts. It further discusses how these principles can guide empirical drug formulation and contribute to the growing field of personalized and integrative medicine. Contemporary research supports the potential of Dravyaguna parameters in identifying bioactivity, forecasting therapeutic outcomes, and understanding herb–drug interactions. Moreover, advancements in pharmacogenomics and systems biology provide a promising platform to scientifically validate these ancient concepts. By integrating traditional Ayurvedic wisdom with modern scientific methodologies, Dravyaguna principles can play a pivotal role in bridging ethnopharmacology and modern drug discovery. The review underscores the relevance of these timeless concepts in developing safer, more effective, and personalized therapeutic interventions in today's evolving medical landscape.

**Keywords:** Dravyaguna, Rasa, Guna, Veerya, Vipaka, Prabhava, Ayurvedic pharmacology.

### 1. INTRODUCTION

Ayurveda, the ancient Indian system of medicine, offers a unique and comprehensive approach to health and healing that emphasizes the interrelationship between the body, mind, and environment. One of its most significant contributions lies in its individualized and holistic approach to pharmacology, encapsulated within the discipline of *Dravyaguna Vijnana*—the science of the properties and actions of medicinal substances. Unlike modern pharmacology, which often isolates active chemical constituents and studies their effects through quantitative and receptor-specific models, *Dravyaguna* focuses on understanding the entirety of a substance, including its sensory attributes, energetic properties, and post-digestive effects on the body. *Dravyaguna Vijnana* encompasses the detailed study of medicinal substances derived from plant, animal, and mineral origins. These substances are evaluated based on five principal attributes: *Rasa* (taste), *Guna* (quality), *Veerya* (potency), *Vipaka* (post-digestive transformation), and *Prabhava* (specific or unique effect). These parameters collectively

guide the Ayurvedic physician in predicting the drug's pharmacological activity, physiological compatibility, therapeutic utility, and potential interactions with the individual's constitution (*Prakriti*). This framework allows Ayurveda to offer customized treatment plans, which align with the current global trend toward personalized medicine [1].

Each principle in *Dravyaguna* plays a predictive role. *Rasa*, or taste, provides immediate sensory feedback and is considered a primary indicator of the drug's physiological action. For instance, *Tikta Rasa* (bitter taste) is associated with detoxifying and antimicrobial properties, while *Madhura Rasa* (sweet taste) is nourishing and anabolic in nature. *Guna*, the qualitative attribute, includes twenty physical properties (e.g., heavy, light, dry, unctuous), which help predict the substance's physiological effects, much like how molecular weight and solubility influence drug behavior in modern pharmacokinetics. *Veerya* refers to the drug's potency or bio-energy, typically classified as *Ushna* (hot) or *Sheeta* (cold), determining whether a substance is stimulating or sedative. This concept parallels the thermogenic or thermolytic properties observed in modern drugs. *Vipaka* describes the long-term effect a substance exerts after undergoing digestion and metabolism, impacting the tissue level and influencing dosha balance and waste elimination. Lastly, *Prabhava* is the specific or extraordinary action of a substance that cannot be explained by other parameters, akin to receptor-specific actions or idiosyncratic drug responses in modern pharmacology [2].

The integration of these five principles allows Ayurvedic pharmacology to go beyond symptom management, offering a predictive and preventive model of therapy. This approach provides a deeper understanding of drug-body interaction that modern pharmacology is only recently beginning to explore through the lens of pharmacogenomics, metabolomics, and systems biology. In recent years, scientific interest in traditional medicine has led to efforts to validate these Ayurvedic principles using contemporary tools. The alignment of *Dravyaguna* with modern pharmacological models—such as taste-receptor studies, molecular docking simulations, and genomic correlations—demonstrates its potential as a predictive model for both drug discovery and therapeutic customization. Therefore, a critical review of *Dravyaguna* principles not only enhances our understanding of Ayurvedic pharmacology but also contributes to the evolution of an integrative and personalized approach to global healthcare.

## 2. METHODOLOGY

This review article is based on classical Ayurvedic texts, peer-reviewed journals, modern pharmacological literature, and current research on Ayurvedic drug mechanisms. Databases such as PubMed, AYUSH Research Portal, and Scopus were searched using keywords like “*Dravyaguna*”, “*Ayurveda pharmacology*”, and “*predictive tools in Ayurveda*”. Textual references were drawn from Charaka Samhita, Sushruta Samhita, and Bhavaprakasha Nighantu, among others.

## 3. CONCEPTUAL FRAMEWORK OF DRAVYAGUNA

### Definition and Scope

*Dravyaguna*, as defined in Ayurvedic classics, refers to the study of the pharmacodynamic and pharmacokinetic aspects of a substance. “*Dravyam guna karmatmakam*” indicates that the action (*karma*) is directly determined by the inherent qualities (*guna*) present in the substance (*dravya*) [3].

### Predictive Nature

The principles of *Dravyaguna* allow Ayurvedic physicians to predict the actions of a new or lesser-known drug based on its sensory and energetic attributes. This anticipatory approach is especially useful in empirical drug formulation and personalized therapeutics [4].

## 4. THE PANCHAVIDHA DRAVYAGUNA PRINCIPLES AS PREDICTIVE TOOLS [TABLE 1]

### Rasa (Taste)

*Rasa* is the initial sensory perception experienced upon ingestion. There are six *Rasas*—*Madhura* (sweet), *Amla* (sour), *Lavana* (salty), *Katu* (pungent), *Tikta* (bitter), and *Kashaya* (astringent). Each *Rasa* has distinct actions on doshas and tissues. For example, *Tikta Rasa* is known for its antimicrobial and detoxifying effects [5]. *Rasa* is correlated with chemical composition; for instance, the bitterness of alkaloids correlates with *Tikta Rasa* [6].

### Guna (Qualities)

There are twenty *Gunas* in Ayurveda, which are considered responsible for physiological and pathological modifications. *Gunas* like *Laghu* (light), *Snigdha* (unctuous), and *Guru* (heavy) determine the bioavailability and metabolism of a drug. For instance, *Laghu Guna* promotes quick absorption and action, much like hydrophilic molecules in modern pharmacokinetics [7].

### Veerya (Potency)

*Veerya* represents the active principle or energy of the drug, generally classified as *Ushna* (hot) and *Sheeta* (cold). *Ushna Veerya* drugs like *Pippali* (*Piper longum*) stimulate digestion and metabolism, mimicking thermogenic and stimulant drugs

in modern pharmacology [8].

### Vipaka (Post-digestive Effect)

Vipaka is the effect of a drug after digestion and metabolism. It influences tissue-level activity and long-term physiological outcomes. For example, drugs with Madhura Vipaka are generally anabolic and nourishing, whereas Katu Vipaka drugs are catabolic [9].

### Prabhava (Specific Action)

Prabhava represents an inexplicable or unique effect that cannot be predicted by other parameters. It is analogous to the receptor-specific actions or idiosyncratic reactions observed in allopathic medicine. For instance, the anti-diabetic effect of Gudmar (*Gymnema sylvestre*) is classified under Prabhava [10].

**Table 1: Panchavidha Dravyaguna Principles and Their Predictive Significance**

Principle	Definition	Examples	Modern Correlation	Predictive Significance
<b>Rasa</b> (Taste)	Initial sensory perception experienced during ingestion. There are six types: Madhura, Amla, Lavana, Katu, Tikta, and Kashaya.	Tikta Rasa – bitter herbs like Neem ( <i>Azadirachta indica</i> )	Correlates with chemical constituents (e.g., bitterness of alkaloids)	Indicates primary action on doshas and tissues; predicts antimicrobial, detoxifying, or nourishing actions
<b>Guna</b> (Qualities)	Twenty inherent physical properties of a substance such as Laghu (light), Snigdha (unctuous), Guru (heavy), etc.	Laghu Guna – herbs like Nagarmotha ( <i>Cyperus rotundus</i> )	Similar to molecular size, polarity, solubility	Predicts absorption, metabolism, and systemic distribution
<b>Veerya</b> (Potency)	The active principle or energetic effect of a substance, generally Ushna (hot) or Sheeta (cold).	Ushna Veerya – Pippali ( <i>Piper longum</i> )	Comparable to thermogenic or stimulant effects	Determines drug's impact on metabolic rate and physiological stimulation or sedation
<b>Vipaka</b> (Post-digestive Effect)	Transformation and effect of the substance after digestion, influencing tissues and long-term effects.	Madhura Vipaka – Haritaki ( <i>Terminalia chebula</i> )	Comparable to metabolic end-products and anabolic/catabolic effects	Predicts tissue nutrition, waste elimination, and hormonal-like long-term impact
<b>Prabhava</b> (Specific Action)	Inexplicable or unique action not accountable by other principles.	Anti-diabetic action of Gudmar ( <i>Gymnema sylvestre</i> )	Analogous to receptor-specific or idiosyncratic actions	Predicts special therapeutic actions beyond observable properties

## 5. MODERN CORRELATIONS AND PREDICTIVE APPLICATIONS

The classical framework of *Dravyaguna* has long guided Ayurvedic pharmacological practice. However, with the advent of modern biomedical sciences, efforts are increasingly being made to contextualize and validate these principles through contemporary scientific methodologies. This section explores the correlations of *Dravyaguna* principles with modern pharmacodynamics, personalized medicine, and pharmacogenomics, thereby highlighting its predictive potential in present-day drug development and therapeutics.

### Correlation with Pharmacodynamics

The understanding of drug action in modern pharmacology often revolves around receptor-ligand interactions, bioavailability, enzyme modulation, and signal transduction pathways. These pharmacodynamic principles can be meaningfully correlated with Ayurvedic parameters. For instance, *Rasa* (taste) may be viewed through the lens of molecular docking and receptor binding affinities, as certain phytochemicals with specific tastes are known to activate or inhibit biological receptors. For example, the bitter taste associated with *Tikta Rasa* corresponds to the presence of alkaloids, which are often antimicrobial and anti-inflammatory [11].

Similarly, *Guna* (qualities) can be correlated with molecular size, solubility, and lipophilicity—factors that influence a

substance's absorption, distribution, and interaction with cellular membranes. For instance, substances with *Laghu Guna* (lightness) may mimic hydrophilic molecules that are readily absorbed and exhibit faster onset of action. *Veerya* (potency), reflecting the energetic effect of a substance, may be understood in terms of a drug's thermogenic or vasodilatory properties, similar to stimulant or antipyretic medications in modern pharmacology.

Thus, *Dravyaguna* principles can serve as qualitative predictors of pharmacodynamic behavior, and their integration with systems biology and cheminformatics can offer a holistic understanding of drug actions.

### Personalized Medicine

Ayurveda's emphasis on *Prakriti*—the unique psychosomatic constitution of an individual—provides a foundational model for personalized medicine. The interplay between *Dravyaguna* and *Prakriti* is central to determining therapeutic suitability, dosage, and duration of treatment for a given patient. For example, an individual with a *Pitta Prakriti* (predominantly fire element) may respond adversely to *Ushna Veerya* (hot potency) drugs and benefit more from *Sheeta Veerya* (cool potency) substances. This personalized approach anticipates the current shift in modern medicine toward tailoring treatments based on an individual's genetic and metabolic profile [12].

Moreover, Ayurveda's focus on long-term balance, rather than short-term symptomatic relief, aligns with contemporary interests in preventative care and holistic wellness. The predictive application of *Dravyaguna* based on individual constitution makes it a valuable complement to genomic and phenotypic profiling in modern healthcare.

### Pharmacogenomics

Emerging research has shown that certain Ayurvedic herbs influence gene expression, protein translation, and epigenetic modulation. For instance, compounds in *Ashwagandha* (*Withania somnifera*) and *Turmeric* (*Curcuma longa*) have been found to interact with transcription factors like NF-κB and modulate inflammatory gene networks.

The Ayurvedic concept of *Prabhava*, which explains specific drug actions unaccounted for by other principles, could be interpreted as reflecting pharmacogenomic variability—where the same drug exerts different effects based on individual genetic makeup [13]. This suggests that *Dravyaguna* incorporates, albeit implicitly, elements of personalized molecular medicine.

The convergence of *Dravyaguna* with pharmacogenomic science offers a novel paradigm wherein classical Ayurvedic concepts can be used to guide biomarker-based drug selection, enhance safety profiles, and reduce adverse drug reactions through individualized therapeutic strategies [Table 2].

**Table 2: Modern Correlations and Predictive Applications of Dravyaguna Principles**

Ayurvedic Concept	Modern Correlation	Example	Predictive Application
<b>Rasa</b> (Taste)	Molecular docking, receptor-ligand affinity	Bitter Rasa (Tikta) linked with alkaloids and antimicrobial activity	Predicts primary action and receptor interaction of compounds
<b>Guna</b> (Qualities)	Molecular weight, solubility, lipophilicity	<i>Laghu Guna</i> correlates with hydrophilic, fast-absorbing drugs	Predicts absorption, metabolism, and systemic distribution
<b>Veerya</b> (Potency)	Thermogenic effects, metabolic stimulation	<i>Ushna Veerya</i> of Pippali mimics stimulant or thermogenic drugs	Predicts metabolic stimulation or suppression
<b>Vipaka</b> (post-digestive effect)	Metabolic end-products, long-term pharmacodynamics	<i>Madhura Vipaka</i> drugs show anabolic, nourishing effects	Predicts long-term impact on tissues, hormones, and metabolism
<b>Prabhava</b> (Specific action)	Pharmacogenomics, receptor specificity, epigenetics	Anti-diabetic action of Gudmar ( <i>Gymnema sylvestre</i> )	Predicts unique drug actions, possibly genotype-specific
<b>Prakriti</b> (Body constitution)	Personalized medicine, genotype-phenotype profiling	<i>Pitta Prakriti</i> individuals respond better to <i>Sheeta Veerya</i> drugs	Aids in personalized drug selection and dosage
<b>Ayurvedic formulation logic</b>	Systems biology, polyherbal synergy	Multi-herb combinations as seen in <i>Triphala</i> , <i>Chyawanprash</i>	Supports multi-targeted therapy for chronic and lifestyle

Ayurvedic Concept	Modern Correlation	Example	Predictive Application
			disorders

## 6. DISCUSSION

The predictive potential of *Dravyaguna* lies in its holistic and integrative framework, which assesses the pharmacological properties of substances through a composite understanding of organoleptic, energetic, and metabolic dimensions. Unlike modern pharmacology, which largely concentrates on identifying and isolating single active constituents and targeting specific molecular receptors, *Dravyaguna* emphasizes the synergistic interaction of multiple attributes—*Rasa* (taste), *Guna* (qualities), *Veerya* (potency), *Vipaka* (post-digestive transformation), and *Prabhava* (specific action). This multi-dimensional approach is particularly significant in the management of complex, chronic, and lifestyle disorders such as diabetes, arthritis, and metabolic syndrome, where mono-targeted interventions often yield limited results.

By enabling predictions of therapeutic action through sensory perception and energetic principles, *Dravyaguna* provides an anticipatory model of drug behavior. For instance, the use of *Tikta Rasa* and *Laghu Guna* drugs in skin disorders or the application of *Sheeta Veerya* in inflammatory conditions reflects a pattern-based logic rather than a reductionist model. Such pattern recognition is aligned with modern systems biology, where the interaction of multiple pathways and molecular targets is increasingly being recognized in pharmacological research.

Modern tools and methodologies offer promising avenues to scientifically evaluate and quantify *Dravyaguna* principles. Quantitative Structure Activity Relationship (QSAR) models can be employed to correlate molecular structures of phytoconstituents with traditional attributes like *Rasa* or *Veerya*. For example, molecular docking studies have begun to link bitter-tasting compounds with anti-inflammatory receptor binding, supporting the predictive role of *Tikta Rasa*. Metabolomics allows profiling of drug metabolites in various physiological environments, potentially validating the Ayurvedic concept of *Vipaka*. Artificial intelligence (AI) and machine learning algorithms can further aid in mapping complex herb–herb and herb–host interactions, contributing to an evidence-based interpretation of classical Ayurvedic pharmacology.

Despite this potential, several limitations hinder the full-scale integration of *Dravyaguna* into modern pharmacological discourse. One of the primary challenges is the lack of standardization in Ayurvedic raw materials and formulations. Variability in drug sourcing, seasonal influence, and processing techniques can significantly alter the pharmacological profile of a substance. Additionally, translating qualitative descriptors like *Snigdha* (unctuous) or *Guru* (heavy) into measurable physicochemical parameters requires sophisticated and context-sensitive analytical methods.

Another challenge is the limited documentation of controlled clinical trials based on *Dravyaguna* principles. While classical Ayurvedic texts provide descriptive accounts of drug action, the absence of statistically significant data in standardized populations limits acceptance in scientific forums. However, recent collaborative efforts between Ayurvedic institutions and biomedical research centers are paving the way for cross-disciplinary research that may help overcome these barriers.

## 7. CONCLUSION

The principles of *Dravyaguna* offer a time-tested and holistic framework for predicting the pharmacological behavior of Ayurvedic drugs. Through the systematic assessment of *Rasa* (taste), *Guna* (qualities), *Veerya* (potency), *Vipaka* (post-digestive effect), and *Prabhava* (specific action), Ayurvedic pharmacology provides an intuitive yet structured approach to forecast therapeutic outcomes. These parameters not only guide drug selection but also offer predictive insights into their efficacy and safety, even in the absence of large-scale clinical trials. In the current era of integrative medicine, the convergence of *Dravyaguna* with modern pharmacological sciences—such as receptor biology, metabolomics, pharmacogenomics, and systems biology—presents a promising pathway for developing more comprehensive, personalized, and effective treatment strategies. The Prakriti-based personalization aligns seamlessly with modern trends in individualized medicine, making *Dravyaguna* highly relevant in contemporary clinical practice. However, the predictive power of *Dravyaguna* will be truly realized when its qualitative principles are standardized, validated, and correlated with quantitative scientific parameters. This requires interdisciplinary research, modern analytical tools, and robust clinical studies. When integrated responsibly, *Dravyaguna* can significantly contribute to evidence-based Ayurveda and the future of global pharmacology.

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## 9. CONFLICT OF INTEREST

None declared.

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