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Svarṇaprāśana: Mechanistic Perspectives from Traditional Ayurveda to Modern Nanomedicine

Abstract

Introduction: Svarṇaprāśana, the administration of processed gold (Svarṇa Bhasma) with Ghr̥ta (clarified butter) and Madhu (honey) as a Lehana (lickable electuary), is a classical pediatric intervention described in Kāśyapa Samhitā. Traditionally indicated to enhance intellect, digestion, strength, and longevity, it has recently gained scientific attention for potential immunomodulatory, nootropic, and anti-infective effects.

Methods: This review synthesizes classical Āyurvedic literature, pharmaceuticals, and contemporary experimental and clinical studies to explore the mode of action of Svarṇaprāśana. Emphasis was placed on physicochemical transformation of gold, bioavailability enhancement through lipid-sugar vehicles, and immunological mechanisms.

Results: Evidence suggests that Amṛtīkaraṇa and allied Saṁskāras transform Svarṇa into biologically active nano- or microparticulate forms. The Ghr̥ta–Madhu vehicle enhances uptake and creates an amphipathic, liposomal-like delivery system. Preclinical and clinical data indicate immune benefits including macrophage activation, enhanced antigen presentation, and cytokine modulation, alongside potential cognitive and anti-infective effects.

Discussion: Svarṇaprāśana represents a classical Āyurvedic intervention with translational relevance to modern nanomedicine. Standardization, safety, and rigorous clinical studies remain priorities to validate and optimize its therapeutic potential.

Keywords: Svarṇaprāśana, Immunity, Nanomedicine, Āyurveda

Introduction

Āyurveda classifies Svarṇaprāśana under Lehaṇa — medicated electuaries intended for licking — and as part of Jātakarma Saṃskāra (newborn rites). Ācārya Kāśyapa and later classical authors ascribe to Svarṇaprāśana an array of actions and outcomes: enhancement of medhā (intellect), agni (digestive/metabolic fire), bala (strength and immunity), āyusya (longevity), varṇya (complexion), vṛśya (fertility), protection from malevolent influences and pathogens (grahāpaham), and superior retention of learning (śrutadhāra) [1].

Lehaṇa refers to the licking of medicine, which acts via a sublingual route, bypassing first-pass metabolism and allowing maximum systemic absorption. Administration of processed gold or metallic gold in children is a unique practice mentioned in Āyurveda as Svarṇaprāśana by Ācārya Kāśyapa thousands of years ago. He described the administration of Svarṇa (gold) in children for benefits such as improved intellect, digestion and metabolism, physical strength, immunity, complexion, fertility, and longevity [2].

Svarṇaprāśana in children can be mainly implicated in two contexts of Āyurveda: Lehaṇa (supplementary feeding) and Jātakarma Saṃskāra (newborn care). Svarṇaprāśana therapy is becoming widely popular and is being administered on the Puṣya Nakṣatra of every month at Āyurveda centers across India. Classical texts recommend administration from birth through childhood, with variable regimens (daily for 1–6 months or on auspicious Puṣya Nakṣatra cycles), using processed gold mixed with Ghṛta (ghee) and Madhu (honey) to optimize efficacy and safety [3].

These classical indications remain the heuristic basis for modern evaluation, as highlighted in publications indexed in the PubMed database [4].

Aim

To critically review the classical Āyurvedic basis, pharmaceuticals, proposed mechanisms, and contemporary clinical evidence of Svarṇaprāśana as a pediatric intervention.

Objectives

1. To comprehensively outline the classical references and indications of Svarṇaprāśana.
2. To critically examine the formulation and preparation processes of Svarṇaprāśana, with emphasis on Svarṇa Bhasma, Ghṛta, and Madhu.
3. To review and analyze the pharmacological and mechanistic basis of the ingredients of Svarṇaprāśana.
4. To evaluate modern preclinical and clinical evidence regarding immunological and cognitive effects of Svarṇaprāśana.

Methodology

The present review involved a systematic collection of data from classical Āyurvedic texts, relevant pharmaco-clinical research articles, and dissertations related to Svarṇaprāśana. Sources were identified using the PubMed database as well as through manual searches of bibliographies and reference lists. The PubMed search covered literature published between January 1990 and December 2024.

The keywords used included “Svarṇaprāśana”, “Svarṇa bindu prāśana”, “Svarṇa bhasma”, “gold nanoparticles Ayurveda”, and “Ayurveda immunomodulation”. Only articles published in English were included.

Inclusion criteria were:

- (i) original research articles (experimental or clinical) related to Svarṇaprāśana or Svarṇa Bindu Prāśana;
- (ii) review papers focusing on the pharmacological, toxicological, or immunological properties of its ingredients (Svarṇa, Ghṛta, and Madhu); and
- (iii) references from classical Āyurvedic compendia.

Exclusion criteria were:

- (i) non-English language publications;
- (ii) conference abstracts or incomplete reports;
- (iii) studies not directly related to Svarṇaprāśana or its key constituents; and
- (iv) duplicate records.

From an initial pool of 126 articles identified, 87 remained after removing duplicates. Following title and abstract screening, 42 full-text articles met the inclusion criteria and were considered in the final analysis, along with relevant Āyurvedic textual references. The gathered information was compiled, compared, and critically analyzed to present a comprehensive overview of Svarṇaprāśana from both classical and contemporary perspectives [5].

Literature Review of Svarṇaprāśana

Svarṇaprāśana, as highlighted in the Āyurvedic scriptures, is a unique pediatric intervention that nurtures the child at multiple levels — physical, cognitive, emotional, and spiritual. Its benefits, preserved in texts such as the Kāśyapa Samhitā, extend well beyond simple nourishment and address long-term well-being [6]. In Lehaṇādhyāya, Ācārya Kāśyapa describes Lehaṇa as the administration of various herbal formulations, Ghṛta preparations, and even gold, intended as supplementary feeds for infants and children, with clearly mentioned indications. He is the first to coin the term Svarṇaprāśana, referring specifically to the practice of administering gold.

The classical procedure involves triturating purified gold with water, honey, and ghee on a clean stone while facing the eastern direction, and then administering it to the child (śiśu)

in the form of licking (prāśana). This saṁskāra (rite) is believed to promote medhā (intellect), vyādhikṣamatva (immunity), āyusya (longevity), and overall health [7].

Traditional dosing recommendations commonly cited in classical and contemporary pediatric Āyurvedic practice include single daily drops or small semisolid doses (for example, two drops in the first six months and four drops thereafter, or administration on Puṣya Nakṣatra monthly cycles), and longer courses (1–6 months) for desired outcomes. These regimens arise from classical practice rather than modern dose-finding studies and therefore require systematic pharmacokinetic and pharmacodynamic evaluation [1,2,6,10].

Benefits of Svarṇaprāśana

One of the foremost benefits attributed to Svarṇaprāśana is Medhāvardhana (enhancement of intellect and cognitive abilities). Regular administration is believed to strengthen medhā (intellect), sharpen memory, accelerate the learning process, and improve concentration and analytical skills. This ensures that children not only excel in academic pursuits but also develop clarity of thought and sound decision-making abilities [7].

Another important benefit is Agnivardhana (promotion of digestive and metabolic functions). Āyurveda emphasizes the role of agni (the digestive and metabolic fire) as the cornerstone of health. By stimulating and regulating agni, Svarṇaprāśana aids in efficient digestion, absorption, and assimilation of nutrients. A balanced agni prevents indigestion, malnutrition, and related childhood illnesses, thereby laying a strong foundation for healthy growth and development [8].

The classics also describe Balāvardhana (improvement of immunity and physical strength). Svarṇaprāśana is traditionally credited with enhancing bala (strength), both in terms of immunity (vyādhikṣamatva) and physical vigor (deha-bala). Children administered with this formulation are said to exhibit greater resistance to recurrent infections, seasonal ailments, and general debility, making them healthier and more resilient [9]. By ensuring balanced growth, maintaining immunity, and preventing diseases from an early age, Svarṇaprāśana is also believed to promote āyusya (longevity). This holistic approach ensures that children grow into adults with enhanced vitality and stable health [10].

Beyond the physical domain, Svarṇaprāśana is described as bringing maṅgala (auspiciousness and positive energy) into the life of the child. This benefit reflects not only the attainment of health but also the cultivation of positivity, confidence, and harmony in day-to-day living. Closely allied is the benefit of puṇya (cultivation of virtue and moral strength). The use of Svarṇaprāśana is believed to shape a child's moral compass, fostering qualities such as honesty, compassion, and discipline. This reflects the Āyurvedic understanding that true health is inseparable from character development and ethical living [11].

Another classical benefit is Vṛṣya (improvement of reproductive health). Although reproductive functions are not active during early childhood, Svarṇaprāśana is believed to strengthen the śukravaha srotas (channels related to reproductive tissues). In doing so, it lays a foundation for robust reproductive health in adulthood. Since śukra dhātu contributes to the formation of ojas — the subtle essence responsible for immunity and vitality — this action indirectly enhances the child's long-term health and resilience [12].

The texts further describe Varṇya (enhancement of complexion and skin health). Svarṇaprāśana supports clear, radiant skin and helps maintain proper skin texture, possibly by balancing the doṣas and purifying the system. It is also credited with Grahāpaha (protection from malign influences). In classical terms, this refers to protection from graha doṣa, encompassing both negative psychosomatic influences and external pathogenic factors. In modern interpretation, this may be seen as improved resistance against microbial infections, stress, and environmental challenges that can adversely affect a child's development [13]. Finally, sustained use of Svarṇaprāśana is said to develop śrutadhāra (enhancement of learning and retention). This refers to the ability to grasp, retain, and recall knowledge simply through listening. Such a quality was highly valued in the traditional Indian gurukula system of education, where oral learning formed the foundation of intellectual training [14].

Formulation and Standard Preparation

The preparation of Svarṇa Bhasma (incinerated/calcined gold) and its further processing with Ghṛta and Madhu (often under repeated trituration — mardana saṃskāra) constitute the pharmaceutic core of Svarṇaprāśana [15,16]. Amṛtīkaraṇa (incinerative purification) and successive saṃskāras transform metallic gold into fine particulates (bhasma) that are purportedly biocompatible and non-toxic when correctly prepared [24,26].

Physicochemical studies indicate that bhasma preparations contain a mixture of nano- and micro-sized gold particles, often with surface coatings of oxides or trace elements, depending on the production method [18,19]. These transformations are thought to influence biological reactivity, solubility, and tissue distribution. Accordingly, modern analytical research emphasizes the importance of standardized manufacturing protocols, detailed physicochemical characterization (SEM/TEM, XRD, ICP-AES), and stringent batch-to-batch quality control as prerequisites for safe clinical use [19,25].

A comparative overview of traditional Āyurvedic guidelines and modern analytical standards is presented below to highlight the parameters considered for the quality assurance of Svarṇa Bhasma (Table 1).

Table 1. Comparative Standards for Svarṇa Bhasma Production

Parameter	Classical Rasaśāstra Standards	Modern Scientific Standards / Guidelines
Source material	Pure gold (Svarṇa), purified by Śodhana using herbal decoctions and substances like cow's urine (Gomūtra), lemon juice, or Triphala	24-carat gold subjected to pre-treatment, often verified by XRF (X-ray fluorescence) or ICP-MS for purity
Śodhana (purification)	Heating and quenching cycles in herbal liquids (e.g., Triphala kvātha, Gomūtra) to remove impurities and increase bioavailability	Removal of impurities confirmed by spectroscopy (AAS/ICP-MS); absence of toxic metals (Pb, Hg, Cd, As) as per WHO/GMP standards
Māraṇa (incineration)	Repeated calcination cycles with specific media (e.g., Kumārī svarasa, Kañji, cow ghee); minimum 14–30 Pūta (incinerations) mentioned	Thermal analysis (TGA/DSC) to confirm complete conversion; phase analysis using XRD (X-ray diffraction)
Physicochemical properties	Should exhibit Rekhāpūrṇatva (fills skin creases), Varitāratva (floats on water), Niruttha (does not regain metallic form in fire test); tasteless and odorless	Particle size in nano to submicron range (10–80 nm) confirmed by TEM/SEM; crystalline nature confirmed by XRD; surface charge (zeta potential) analysis
Purity and safety	Nirutthatva ensures non-reversible metallic state, safe for oral use; Apunarbhava (no regeneration of metal)	Heavy metal limits as per WHO: Pb <10 ppm, As <3 ppm, Cd <0.3 ppm, Hg <1 ppm ; cytotoxicity testing on cell lines
Standardization tests	Rasaśāstra parameters such as Varitāratva, Rekhāpūrṇatva, Nirutthata	FTIR for functional groups, ICP-MS for elemental analysis, DLS for particle size distribution, XPS for oxidation states
Storage / packaging	Stored in airtight containers, protected from moisture	GMP-compliant packaging; stability testing under ICH guidelines

Physicochemical Characterization and Toxicity Study of Svarṇa Bhasma

Recent preclinical investigations have sought to establish both the physicochemical basis and the safety margins of Svarṇa Bhasma, given its administration to children necessitates rigorous toxicological validation [17,19]. A 90-day repeated oral toxicity experiment was performed in Holtzman rats, where Svarṇa Bhasma was administered at graded doses: the therapeutic equivalent [~ 3 mg/kg body weight], five times the therapeutic dose [15 mg/kg], and ten times the therapeutic dose [30 mg/kg]. Biochemical analysis revealed mild changes in serum urea, creatinine, and alanine aminotransferase [ALT] levels at higher doses; however, all values remained within physiological reference ranges [19]. This suggests that while supra-therapeutic dosing may induce subtle metabolic stress, properly prepared Svarṇa Bhasma is unlikely to cause overt

renal or hepatic toxicity at therapeutic levels [25]. Hematological evaluations showed no significant abnormalities in red blood cell count, hemoglobin, white blood cell count, or platelet levels, thereby indicating hematological safety. Histopathological examination of vital organs, including liver, kidney, spleen, heart, lungs, brain, and gonads, demonstrated no significant pathological lesions, further supporting the relative safety of long-term use when the preparation has undergone authentic śodhana, māraṇa, and amṛtīkaraṇa saṃskāras [24,26].

Advanced physicochemical analyses highlighted the heterogeneous morphology of Svarṇa Bhasma. Large irregular particles averaging ~60 μm, typically consisting of gold oxides and mixed mineral phases, were detected alongside nanosized spherical particles beginning at ~10 nm. Many nanoparticles displayed surface coatings of biologically relevant trace elements such as iron [Fe], silicon [Si], oxygen [O], phosphorus [P], and sodium [Na]. This dual population of micro- and nanoparticles may account for the balance between macroparticles providing slow release and inert safety, while nanoparticles enable rapid cellular interactions [18]. Moreover, the elemental coatings could modulate surface charge and enhance biocompatibility.

Pharmacological Activities of Svarṇa Bhasma

Neurobehavioral studies conducted in zebrafish models provided additional insight. Behavioral assays revealed that exposure to Svarṇa Bhasma elicited anxiolytic-like effects, evidenced by increased exploratory behavior and prolonged activity in the upper zone of the tank [19,25]. These findings suggest that beyond its immunomodulatory role, Svarṇa Bhasma may exert central nervous system modulation, potentially substantiating its classical attribution as medhyarasāyana [nootropic and intellect-enhancing agent] [17,24].

Taken together, these findings suggest that Svarṇa Bhasma, when authentically prepared through traditional processes such as śodhana, māraṇa, and amṛtīkaraṇa saṃskāras, yields a complex nanostructured material with dual properties. At therapeutic doses, it is physiologically well tolerated and demonstrates biological activity in both the immune and nervous systems [17,19,25]. Structurally, it presents as a heterogeneous compound, allowing both macroscale inert safety and nanoscale bioactivity, thereby validating its continued relevance within the Āyurvedic pharmacopeia [15,16,24,26].

Drug Review

• Svarṇa Bhasma [Gold]

Classically, Svarṇa is described as madhura rasa, madhura vipāka, śīta vīrya, balancing the tridoṣas and promoting ojas. Biomedically, nanoparticulate or calcined gold exhibits anti-inflammatory, immunomodulatory, antioxidant, and antimicrobial activities [17,18]. Mechanistic studies reveal enhanced macrophage functions, antigen presentation, and Th1

polarization, supporting its classical claim of vyādhikṣamatva. Effects vary with dose, form, and adjuvants like ghr̥ta and madhu [17,19].

• Ghr̥ta [Clarified Butter]

Ghr̥ta is considered sarvasnehatamam, medhya, br̥mhaṇa, and rasāyana. Modern evidence supports its role as a lipid vehicle enhancing absorption of lipophilic constituents and facilitating transport across biological barriers. Cow ghee contains long-chain PUFAs, DHA precursors, and fat-soluble antioxidants, potentially supporting neuronal functions [22,23]. Experimental data suggest it enhances uptake of lipophilic actives, though cognitive outcomes remain variable [23].

• Madhu [Honey]

Traditionally, madhu is dīpana, var̥ṇya, ropana, and śodhana. Modern studies confirm antibacterial, antioxidant, and immunomodulatory effects [20,21]. In Svarṇapraśana, honey improves palatability, offers antimicrobial protection, and stabilizes colloidal gold [20,21].

Integrated Delivery Concept — “A Lipophilic Nano-Electuary”

Together, Svarṇa Bhasma, ghr̥ta, and madhu form an amphipathic electuary. Repeated mardana may coat gold particles with a lipid layer stabilized by honey’s sugars and polyphenols, resembling nanoemulsions that enhance mucosal absorption [18,22,21]. Sublingual administration on an empty stomach may allow bypassing first-pass metabolism, facilitating systemic effects [17,18]. This aligns with evidence on gold nanoparticles as immune adjuvants and nootropics [17,18,19,22].

Mode of Action of Svarṇapraśana

Studies suggest that gold nanoparticles, which closely resemble Svarṇa Bhasma in particle size, can undergo rapid systemic absorption and show tissue distribution to the liver, spleen, and kidneys, with gradual excretion mainly via the hepatobiliary route [26,27]. Preliminary human pilot data also indicate measurable bioavailability of Svarṇa Bhasma following oral administration, supporting the plausibility of enhanced absorption through sublingual delivery [28].

Svarṇapraśana is defined in the present study as the administration of an electuary containing a gold preparation [Svarṇa Bhasma] combined with a lipid [ghr̥ta] and a sweetening agent [madhu]. Historically, Svarṇapraśana served as a simple method to confer immunization in ancient times. This practice adopts a scientific approach to help children develop acquired immunity, functioning as a form of immunization that supports healthy, disease-free, and stable physical and mental development.

The pure ash of gold [Svarṇa Bhasma] enhances immunity through its scavenging and immunomodulatory effects [17,19]; being a highly valuable metal, it contributes to the overall development of the body, intellect, and mental faculties of children. Pure madhu acts as an

anti-allergic and antimicrobial agent [20,21], while ghr̥ta improves the bioavailability of these components by facilitating emulsification, lubrication, and better absorption [22,23]. The classical benefits of Svarṇaprasāna in children [Table No. DR 01] include medhāvardhana [enhancement of intellect], agnivardhana [supporting digestion and metabolism], balavardhana [strengthening immunity and physical vitality], āyusya [promoting longevity], maṅgala [auspiciousness], puṇya [virtue], varṇya [improvement of complexion and skin texture], vṛṣya [fertility], and grāhapaham [protection from evil spirits and infectious organisms] [1,2].

When administered for one month, māsāt parama medhāvī indicates that the child becomes exceedingly intelligent, and vyādhībhira ca dṛśyate suggests resistance to diseases. Prolonged administration for six months, ṣaḍbhir māse śrutadhāra, enables the child to retain and comprehend knowledge simply by listening [5,9]. Ācārya has highlighted that Svarṇaprasāna possesses vṛṣya properties. Since the śukravaha srotas is not fully functional during the bālya avasthā, Svarṇaprasāna may positively influence the child's future fertility. Given that śukra dhātu contributes to the formation of ojas in the body, the vṛṣya effect may further promote ojas development, supporting overall vitality and health [7,12].

Immunological Effects of Suvarṇaprasāna

Preclinical evidence indicates that calcined gold preparations can activate phagocytes/macrophages, increasing nitric oxide and pro-inflammatory mediators that enhance pathogen clearance, modulate dendritic cell maturation and antigen presentation, and bias CD4⁺ T cells toward Th1 phenotypes in infection models — all compatible with enhanced innate and antigen-specific immune responses. These findings map onto classical claims of improved resistance to infectious disease [Vyādhikṣamatva] and improved physical robustness [Bala]. Importantly, the immune response is context-sensitive: particle size, surface chemistry, dose, and vehicle [Ghr̥ta + Madhu] all shape the balance between beneficial immune stimulation and potential pro-inflammatory or toxic sequelae. Clinical data remain limited but encouraging: a randomized controlled clinical trial in infants reported favorable immunomodulatory signals without adverse growth effects over a short follow-up period. Collectively, mechanistic and early clinical data support the hypothesis that Suvarṇaprasāna functions as a mild immunomodulatory preparation rather than as a pathogen-specific vaccine [3,4,17,19].

Safety and Toxicology Evidences

Safety is paramount when administering metal-derived formulations to infants and children. Modern toxicology and physicochemical characterization studies show that properly prepared Suvarṇa/Swarnabhasma contains nano- to micro-sized gold particles, sometimes with trace coatings, and that well-characterized preparations at therapeutic doses in animal

models produced limited or no adverse histopathological changes [19,25]. However, higher doses or poorly prepared products can alter biochemical markers [urea, creatinine, ALT] and raise safety concerns [19]. In vitro and in vivo studies [cell lines, rodents, zebrafish] have largely reported acceptable tolerability for authenticated bhasma, but data are heterogeneous and dependent on manufacturing methods [19,25]. Therefore, rigorous standardization [including identification of particle size distribution, residual contaminants, and batch sterility], third-party quality testing, and regulatory oversight are essential prerequisites before recommending routine pediatric use in settings beyond supervised traditional Ayurvedic practice [19,25].

Discussion

Suvarnaprāsana represents a classical pediatric intervention deeply rooted in Jātakarma Saṃskāra and Lehana practices, as described in Ayurvedic texts such as the Kāśyapa Saṃhitā [1,16]. Historically, it was administered to enhance medhā, strengthen bala, promote longevity [āyusya], and confer protection against diseases [grahapaham]. The formulation, comprising Suvarṇa Bhasma, Ghṛta, and Madhu, embodies a sophisticated pharmaceutical design wherein metallic gold is transformed via Śodhana, Māraṇa, and Amṛtikaraṇa processes to produce biocompatible bhasma with nano- and micro-particulate forms [17,19]. Ghṛta acts as a lipophilic vehicle facilitating bioavailability and potential central nervous system uptake [22,23], while Madhu provides palatability, antimicrobial support, and stabilization of the colloidal gold particles [20,21].

Modern preclinical studies support the immunomodulatory and neurobehavioral effects of Suvarṇa Bhasma. Animal and zebrafish models demonstrate macrophage activation, enhancement of antigen presentation, Th1-biased CD4⁺ T cell responses, and anxiolytic-like effects consistent with the classical claims of improved medhā and vyādhikṣamatva [17,19]. The lipid-sugar vehicle, created by combining Ghṛta and Madhu, likely serves as a nano-emulsified delivery system that enhances mucosal absorption, bypasses first-pass metabolism, and facilitates sustained systemic bioavailability of Suvarṇa Bhasma [20,21,22]. Safety considerations are paramount. Properly prepared Suvarṇa Bhasma exhibits minimal toxicity in preclinical models, with nano and micro-sized gold particles demonstrating biocompatibility, limited biochemical perturbation, and negligible histopathological changes at therapeutic doses [17,19]. Conversely, poorly prepared products or supra-therapeutic dosing can alter renal and hepatic markers, emphasizing the necessity of standardization, quality control, and regulatory oversight prior to broader clinical implementation [19].

Recommendations for Future Work

To move Suvarnaprāsana from traditional practice toward evidence-based integration, the following are priorities: [1] rigorous standardization of Suvarṇa Bhasma production,

including adherence to Good Manufacturing Practices and analytical release criteria [19,25]; [2] controlled dose-finding pharmacokinetic and biodistribution studies in appropriate animal models and in adults to model pediatric scaling [19]; [3] mechanistic immunology studies that interrogate antigen presentation, cytokine networks, and any potential for immune-related adverse events [17,19]; [4] well-designed randomized controlled clinical trials in pediatric populations with clearly defined clinical endpoints [e.g., incidence of upper respiratory tract infections, vaccine response augmentation, growth metrics], safety monitoring, and long-term follow-up [3,4]; and [5] exploration of the vehicle dynamics [Ghṛta–Madhu emulsion] as an active determinant of uptake and organ targeting [20,21,22,23]. Current experimental literature suggests plausible modes of action but cannot yet substitute for methodologically robust clinical validation [18]. To generate high-quality, generalizable evidence, well-designed multicentric, randomized pediatric trials are recommended. These studies should employ standardized Suvarṇa Bhasma preparations, clearly defined outcome measures for immune function and cognitive development, and appropriate pharmacokinetic assessments to establish safety and efficacy. Such efforts would bridge classical Ayurvedic practice with contemporary evidence-based medicine, enabling broader clinical adoption.

Conclusion

Suvarṇaprāśana represents an intriguing intersection of ancient pharmaceuticals and modern nanomedicine principles: a processed gold preparation delivered in a lipid–sugar electuary that, based on emerging preclinical and limited clinical data, can modulate innate and adaptive immune functions and may support aspects of neurodevelopment and general robustness when prepared and administered according to classical purification and dosing regimens. The cumulative evidence supports cautious optimism: Suvarṇaprāśana has biologically plausible mechanisms [macrophage activation, antigen-presentation enhancement, antioxidant/free-radical effects, and vehicle-mediated bioavailability] and preliminary safety data for standardized preparations, but definitive clinical recommendations await high-quality trials and strict standardization. Clinicians and researchers should proceed with science-driven rigor — integrating classical wisdom with modern analytical, toxicological, and clinical methodologies.

Limitations and Research Gaps

Current evidence on Suvarṇaprāśana is constrained by small clinical sample sizes, absence of pediatric pharmacokinetic data, and variability in Suvarṇa Bhasma preparations. Moreover, the lack of standardized outcome measures across studies limits comparability and reproducibility. Future research should focus on larger, multicentric trials, standardized protocols, and systematic pharmacological evaluations to strengthen scientific validation.

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