iJRASET
International Journal For Research in Applied Science and Engineering Technology



INTERNATIONAL JOURNAL FOR RESEARCH

IN APPLIED SCIENCE & ENGINEERING TECHNOLOGY

Volume: 14 **Issue:** IV **Month of publication:** April 2026

DOI: https://doi.org/10.22214/ijraset.2026.80737

www.ijraset.com

Call: 08813907089 | E-mail ID: ijraset@gmail.com

International Journal for Research in Applied Science & Engineering Technology (IJRASET)
ISSN: 2321-9653; IC Value: 45.98; SJ Impact Factor: 7.538
Volume 14 Issue IV Apr 2026- Available at www.ijraset.com

Human-Derived Cellular Components in Cosmetic Applications: Mechanisms, Formulation, and Future Perspectives

Shreyash Chaudhari[1], Sonali K. Shambharkar[2], Dr. Manoj. V. Girase[3]
[1]Student, [2]Assistant Professor, [3]Head of Department, R. C. Patel Institute of Pharmaceutical Education & Research, Shirpur

ABSTRACT: ***In contemporary beauty research, human-derived cellular components have become sophisticated bioactive substances with great potential for anti-aging and skin regeneration [1-3]. These elements, which include fibroblasts and conditioned media, are abundant in extracellular matrix proteins, growth factors, and cytokines that are essential for boosting collagen synthesis, encouraging cellular proliferation, and enhancing skin moisture [4-6]. The molecular mechanisms behind their activity are thoroughly covered in this review, with a focus on pathways related to oxidative stress and tissue repair. reduction, and dermal remodelling[5,7]. In addition, along with important formulation issues for preserving stability and efficacy, the article covers their varied applications in cosmetic formulations, including serums, creams, and skin restoration treatments. The usage of materials produced from humans is also severely assessed in terms of safety, ethical issues, and regulatory considerations. Additionally, new alternatives are investigated as potential avenues for cosmetic innovation in the future, such as biomimetic peptides and plant-derived stem cells [22–24]. All things considered, the development of next-generation cosmeceuticals using human-derived cellular components is a promising strategy, but rigorous assessment of safety and regulatory compliance is still crucial.***

KEYWORDS: ***Human-derived cellular components, Cosmeceuticals, Fibroblasts, Skin regeneration, Anti-aging, Bioactive compounds***

I. INTRODUCTION

With the incorporation of cutting-edge biotechnological advances [1-3] targeted at enhancing skin health and beauty, the global cosmetics industry has experienced a substantial transformation. Biologically active substances that provide improved efficacy are increasingly being added to or used in place of traditional cosmetic formulations, which are mostly made of synthetic and plant-derived chemicals. Human-derived cellular components have emerged [2,4] as a fresh and promising technique in the creation of cosmeceutical products among these developments.

Materials generated from fetal cells, especially fibroblasts and their conditioned media, have garnered a lot of interest because of their great potential for regeneration and capacity to promote skin repair processes. Numerous growth factors, cytokines, and extracellular matrix components are known to be produced by these cells and are essential for preserving the structure and function of the skin. Their anti-aging, skin-rejuvenating, and wound-healing qualities are the main reasons they are employed in cosmetic compositions.

Research into the use of such bioactive compounds, which can increase collagen formation, improve skin suppleness, and lessen wrinkles, has been spurred by the growing need for effective anti-aging remedies. Nevertheless, the use of components produced from fetal cells presents serious ethical, legal, and safety issues notwithstanding their scientific benefits. Their popularity and commercialization are influenced by differences in public perception and regulatory systems around the world.

The purpose of this review is to methodically investigate the usage of chemicals generated from fetal cells in cosmetic goods, including their modes of action, formulation features, uses, safety concerns, and ethical ramifications. The study also emphasizes future prospects and present difficulties in this developing area of cosmetic science.

II. TYPES OF HUMAN-DERIVED CELLULAR COMPONENTS

A. Fetal Fibroblast Cells

Definition:Fetal fibroblasts are mesenchymal cells derived from embryonic or fetal connective tissue.

©IJRASET: All Rights are Reserved

Unlike adult fibroblasts, these cells are in a "privileged" biological state, characterized by a higher rate of proliferation and a unique capability to synthesize Type III collagen (fetal collagen) alongside Type I.

Source: In the cosmetic industry, these are not harvested directly for every product. Instead, they are sourced from immortalized cell lines (e.g., the MRC-5 or WI-38 lines) originally established from legally and ethically consented fetal tissue donations in the 1960s. These cell lines are banked and multiplied in laboratories globally.

Role in Skin Repair: Fetal fibroblasts are the "architects" of scarless healing. They secrete a specific ratio of Type III to Type I collagen. In aging skin, this ratio is skewed; the introduction of fetal fibroblast-derived proteins helps recalibrate the extracellular matrix (ECM), promoting regeneration rather than just "patching" the skin. They also produce high levels of Hyaluronic Acid (HA) and Fibronectin, which enhance the skin's structural integrity.

B. Stem Cell-Derived Extracts

Rather than using live cells (which cannot survive in a cosmetic cream), the industry utilizes the **extracts**—the bioactive molecules produced by stem cells.

Growth Factors (GFs): These are natural proteins capable of stimulating cellular proliferation [6–8] and differentiation. Key GFs include:

- TGF-β (Transforming Growth Factor): Crucial for collagen induction.
- EGF (Epidermal Growth Factor): Accelerates epidermal turnover and barrier repair.
- FGF (Fibroblast Growth Factor): Directly targets the dermis to increase elasticity.

Cytokines:These serve as "messenger molecules" that facilitate cellular communication and inflammation. In order to reduce the micro-inflammation linked to "inflamm-aging," anti-inflammatory cytokines are given priority in cosmetic technologies. This prevents oxidative stress from interfering with the skin's repair processes.

C. Conditioned Media (CM)

The "gold standard" for integrating technology obtained from humans into stable cosmetic compositions is now conditioned media.

What It Includes: CM is the "broth" or supernatant that remains after culturing cells in a lab, such as fetal fibroblasts. It is an intricate concoction that includes:

- Secretome: The full array of GFs, cytokines, and chemokines.
- Exosomes: Micro-vesicles that protect and transport RNA and proteins directly into target skin cells.
- Extracellular Matrix Proteins: Soluble collagen and elastin precursors.

Use in Cosmetics: 1. Formulation Stability: Since CM is cell-free, it can be filtered and stabilized. However, as a Cosmetic Technologist, you must address the thermal stability—these proteins can denature at temperatures above 40°C.

2. Delivery Systems: CM is often encapsulated in liposomes or nanospheres to ensure these large-chain proteins penetrate the stratum corneum to reach the viable epidermis and dermis.

3. Application: Primarily used in "Medical Grade" skincare for post-laser recovery, deep-wrinkle serums, and advanced regenerative night creams.

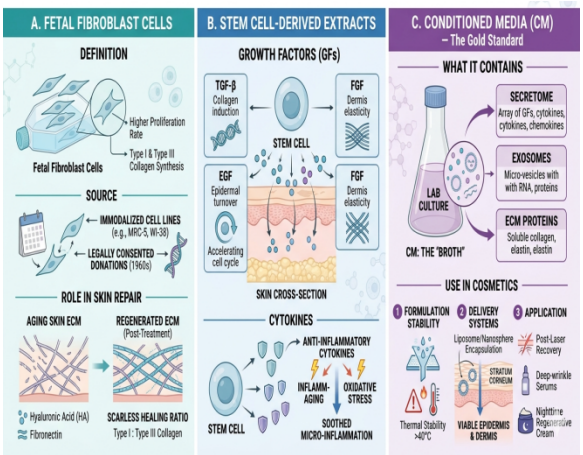


Figure 1 Type of Human-Derived CellularComponent for Skin Regeneration

ISSN: 2321-9653; IC Value: 45.98; SJ Impact Factor: 7.538
Volume 14 Issue IV Apr 2026- Available at www.ijraset.com

III. MECHANISM OF ACTION

Because they are rich in growth factors, cytokines, peptides, and extracellular matrix (ECM) proteins, fetal cell-derived components such as fibroblasts, stem cell extracts, and conditioned media show considerable biological activity. These bioactive compounds have molecular interactions with skin cells and control several signaling pathways related to skin healing, regeneration, and anti-aging.

A. Collagen Synthesis Stimulation

The principal structural protein of the dermis, collagen is in charge of preserving the integrity, strength, and suppleness of the skin. It is known that components produced from fetal fibroblasts secrete a variety of growth factors, including fibroblast growth factor (FGF), transforming growth factor-beta (TGF-β), and epidermal growth factor (EGF), all of which are essential for collagen formation.

These growth factors trigger intracellular signaling cascades, including the TGF-β/Smad pathway, by binding to certain receptors on dermal fibroblasts. Genes that produce collagen, particularly Type I and Type III collagen, are upregulated when this pathway is activated. Furthermore, these factors preserve the extracellular matrix by inhibiting matrix metalloproteinases (MMPs), which are enzymes that break down collagen.

Therefore, increased collagen synthesis [1,3,4] promotes skin firmness, improves dermal density, and lessens the visibility of fine lines and wrinkles.

B. Cell regeneration

A complex blend of bioactive chemicals found in conditioned media produced from fetal cells encourages cellular migration, proliferation, and differentiation [6–8]. These chemicals speed up the natural process of skin renewal by stimulating fibroblasts in the dermis and keratinocytes in the epidermis.

Growth factors like FGF and EGF increase skin cells' mitotic activity, which accelerates cell turnover and replaces aging or damaged cells. Cytokines also control inflammatory reactions, which promotes tissue regeneration and repair.

Improved skin texture, quicker wound healing, and the return of normal skin architecture are all benefits of this increased regenerative activity. Because of their special biological characteristics, fetal cells can facilitate scarless recovery, which increases their importance in cosmetic applications.

C. Anti-aging effects

Several pathways that target both internal and extrinsic aging variables mediate the anti-aging benefits [5,22] of components generated from fetal cells. The promotion of extracellular matrix (ECM) formation, which includes collagen, elastin, and glycosaminoglycans—all crucial for preserving skin elasticity and resilience—is one of the key effects.

Additionally, by lowering reactive oxygen species (ROS) produced by environmental stressors including pollution and UV radiation, these components demonstrate antioxidant qualities. They stop cellular damage, protein breakdown, and early aging by reducing oxidative stress.

Bioactives produced from fetal cells also alter cellular signaling pathways that enhance the firmness and suppleness of the skin. Wrinkles, fine lines, and sagging are visibly reduced as a result, giving the skin a more youthful appearance.

D. Skin hydration

Maintaining healthy and effective skin depends heavily on skin moisture. By strengthening the skin barrier and encouraging the production of natural moisturizing factors (NMFs), components obtained from fetal cells help to increase moisture.

Keratinocytes are stimulated by cytokines and growth factors to create vital proteins and lipids like filaggrin, which are crucial for preserving the stratum corneum's integrity. This lowers transepidermal water loss (TEWL) and fortifies the skin barrier.

Furthermore, conditioned media's water-binding ability is improved by the presence of glycosaminoglycans and compounds that resemble hyaluronic acid, which increases moisture retention. As a result, the skin becomes smoother, softer, and more hydrated overall.

The mechanism of action is illustrated in Fig. 2.

©IJRASET: All Rights are Reserved | SJ Impact Factor 7.538 | ISRA Journal Impact Factor 7.894 |

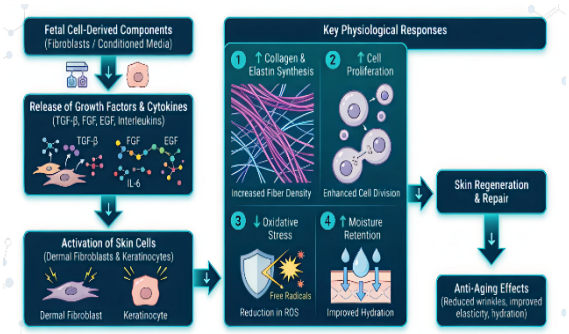


Figure 2Mechanism of action of fetal cell-derived components

IV. APPLICATIONS IN COSMETIC PRODUCTS

Because of their regenerative, anti-aging, and reparative qualities, fetal cell-derived components—in particular, fibroblast cultures, conditioned media, and bioactive extracts—have become more important in contemporary cosmetic formulations. When added to topical solutions, these ingredients' abundance of growth factors, cytokines, peptides, and extracellular matrix (ECM) proteins provide a variety of functional advantages. As will be covered below, they are used in a variety of cosmetic and cosmeceutical product categories.

Table 1: Components and Their Functions in Cosmetic Applications

Component	Function	Cosmetic Use
Fibroblasts	Stimulates synthesis of Collagen Type I & III and Elastin fibers.	Anti-aging creams, firming lotions.
Growth Factors *(EGF, FGF, TGF-β)*	Promotes rapid cell proliferation, differentiation, and tissue regeneration.	High-potency serums, recovery essences.
Conditioned Media	Provides a complex secretome for deep skin repair and rejuvenation.	Post-procedure repair products, night creams.
Cytokines	Regulates inflammatory pathways and mediates cellular communication.	Sensitive skin formulations, soothing balms.
ECM Proteins	Provides immediate structural support and scaffolds the skin matrix.	Instant-lift and firming formulations.

Table I summarizes the key components and their roles in cosmetic applications.

A. Anti-Aging Creams

One of the most common uses of chemicals obtained from fetal cells is in anti-aging lotions. These formulas are especially made to lessen wrinkles, fine lines, and skin elasticity loss—all of which are obvious indicators of aging.

Through the action of growth factors such as fibroblast growth factor (FGF) and transforming growth factor-beta (TGF-β), components produced from fetal fibroblasts improve the production of collagen and elastin in the dermal layer. The skin becomes firmer and more structurally sound as a result. These bioactive compounds also improve the overall texture of the skin by promoting cellular turnover and healing damaged skin tissues.

Fetal cell-derived extracts' antioxidant qualities are also essential for counteracting reactive oxygen species (ROS), which cause premature aging. Regular use of these creams might therefore result in skin that is firmer, smoother, and looks younger.

B. Serums and Essences

High concentrations of active ingredients are delivered straight into the skin's deeper layers via serums and essences, which are lightweight, quickly absorbing compositions. Because of their low molecular weight and strong bioactivity, components produced from fetal cells are especially well suited for these formulations.

These products frequently include conditioned media enhanced with growth factors, cytokines, and peptides that operate on target cells like fibroblasts and keratinocytes by penetrating the epidermis. As a result, extracellular matrix formation is stimulated, cell proliferation is increased, and hydration is improved.

Additionally, serums and essences offer focused treatment for particular skin issues like dullness, uneven skin tone, and early indications of aging. They are the perfect delivery strategy for fetal cell-derived bioactives in sophisticated skincare regimens because of their quick absorption and great efficacy.

C. Skin Repair Products

In order to restore injured or stressed skin brought on by environmental causes including UV radiation, pollution, and chemical exposure, skin restoration treatments containing components generated from fetal cells are frequently employed. These products are made to encourage skin barrier renewal and healing.

By promoting cell division and boosting the production of structural proteins, the bioactive compounds found in fetal cell extracts quicken the healing process. While cytokines control inflammation and aid in tissue repair, growth factors like epidermal growth factor (EGF) are crucial in encouraging epidermal regeneration.

Additionally, by improving lipid production and lowering transepidermal water loss (TEWL), these products aid in the restoration of the integrity of the skin barrier. As a result, the skin is more resilient, less sensitive, and heals from injuries more quickly.

D. Scar and Wound Healing

Scar reduction and wound healing products are among the most important uses of components obtained from fetal cells. Fetal cells have special regenerative qualities, such as the capacity to accelerate recovery with little scarring—a process known as "scarless healing."

Growth factors and cytokines found in fetal cell-conditioned medium promote cellular migration, angiogenesis (the creation of new blood vessels), and collagen remodeling—all processes necessary for efficient wound healing. Additionally, these elements control the inflammatory response, avoiding the development of excessive fibrosis and scar tissue.

These products are used in cosmetic applications to lessen the visibility of surgical scars, acne scars, and other skin flaws. They are quite useful in both therapeutic and cosmetic dermatology because of their capacity to enhance skin texture and encourage consistent healing.

V. FORMULATION ASPECTS

The biological nature and sensitivity of components obtained from fetal cells necessitate careful attention while formulating cosmetic products. Peptides, growth factors, cytokines, and fibroblast-conditioned medium are among the components that are most vulnerable to chemical and environmental deterioration. In order to guarantee product efficacy, it is crucial to preserve their stability, compatibility, and safety during formulation.

©IJRASET: All Rights are Reserved | SJ Impact Factor 7.538 | ISRA Journal Impact Factor 7.894 |

A. Stability issues

Fetal cells produce bioactive compounds that are inherently unstable and sensitive to environmental factors such as temperature, light, oxygen, and pH. These elements, particularly proteins and peptides, are susceptible to denaturation and degradation under unfavorable conditions, which leads to the loss of biological function.

High temperatures during manufacture or storage can damage the structural integrity of proteins, and exposure to UV light can cause photodegradation. Reactive oxygen species-induced oxidative stress may further alter the molecular structure of these bioactives. Additionally, the durability of such components depends on maintaining an optimal pH range, which is often close to physiological pH.

To get around these problems, sophisticated formulation strategies like liposomal delivery systems, microencapsulation, and antioxidant incorporation are employed. By stopping degradation, these techniques increase the shelf life of sensitive bioactive compounds.

B. Incorporation in creams/serums

When adding components made from fetal cells to cosmetic products, creams and serums are frequently utilized as effective delivery systems.

Creams provide a suitable medium for the uniform dispersion of bioactive components and are often prepared as water-in-oil (W/O) or oil-in-water (O/W) emulsions. However, because of their heat sensitivity, these components are frequently added during the cooling phase of formulation, typically below 40°C, to prevent denaturation. Careful selection of stabilizers and emulsifiers is necessary to maintain formulation stability.

Serums are lightweight, mostly aqueous or gel-based, and enable higher concentrations of active substances and improved skin penetration. These formulations facilitate the rapid absorption of growth factors and peptides. However, due to their limited protective matrix, additional stabilizing techniques such encapsulation may be required to prevent deterioration of the active components.

C. Compatibility with other ingredients

The compatibility of components derived from fetal cells with other formulation ingredients is a significant factor influencing the stability and efficacy of a product. These bioactives must retain their chemical and physical stability while excipients like emulsifiers, preservatives, fragrances, and other active ingredients are present.

Among the chemicals that may destabilize proteins and reduce their effectiveness are strong acids, alcohols, and reactive compounds. Interactions between incompatible chemicals can cause bioactive molecules to precipitate, agglomerate, or degrade.

To ensure compatibility, formulators must conduct thorough stability and compatibility studies. The functional integrity of components made from fetal cells depends on the maintenance of an optimal pH environment and the use of mild, non-reactive excipients.

D. Preservation challenges

Preservation is a crucial part of formulations using fetal cell-derived products since they are vulnerable to microbial contamination. The proteins and other nutrients in these components may promote microbial growth if they are not well managed.

Standard preservatives may pose problems since some compounds can denature proteins or interfere with the activity of bioactive molecules. Therefore, the selection of preservatives must provide both safety and compatibility while offering effective antibacterial protection.

Alternative preservation methods include airless containers, natural antibacterial agents, and mild preservatives. These techniques extend product shelf life and lessen contamination. To ensure the product's quality and safety, strict adherence to hygienic manufacturing practices is also required.

VI. SAFETY AND TOXICOLOGICAL ASPECTS

Because of their biological origin, the use of components produced from fetal cells in cosmetic compositions requires a careful assessment of safety and toxicological characteristics. These elements, which include growth factors, cytokines, and fibroblast-conditioned media, need to be evaluated for any negative impacts on human skin and general health. A thorough safety evaluation includes risk analysis, clinical evaluations, in vitro and in vivo research, and compliance with regulations.

©IJRASET: All Rights are Reserved | SJ Impact Factor 7.538 | ISRA Journal Impact Factor 7.894 |

A. Skin irritation studies

Skin irritation tests are essential to determine whether components made from fetal cells have the potential to cause irritation or sensitization when applied topically. These studies frequently employ both in vitro models, such as reconstructed human epidermis, and in vivo patch testing on human participants.

The evaluation's primary focus is on parameters including erythema, edema, and any inflammatory response following administration. Bioactive proteins and cytokines derived from fetal cells may elicit immune-mediated responses in sensitive individuals. Nonetheless, most formulations are designed to be biocompatible and non-irritating when taken in appropriate amounts.

Additionally, repeated application studies are conducted to assess cumulative irritation and long-term skin compatibility. When these ingredients are correctly manufactured and purified, the risk of irritation is significantly reduced.

B. Clinical safety

Clinical safety assessment uses controlled human research to evaluate the overall safety, tolerability, and efficacy of cosmetic products containing components derived from fetal cells. These studies are conducted under dermatological supervision and follow ethical guidelines.

Among the factors assessed are skin compatibility, the incidence of adverse responses, and overall user acceptance. Clinical trials may also evaluate other benefits such enhanced skin hydration, elasticity, and texture.

The safety profile of these components is greatly influenced by their source, extraction method, and degree of purification. High-quality processing and controlled manufacturing processes are essential to ensuring that the final product is safe for human use.

C. Risk assessment

Risk assessment is a systematic process that looks at potential hazards associated with using materials made from fetal cells in cosmetic products. It includes hazard identification, exposure assessment, dose-response analysis, and risk characterization.

Potential risks include immunogenic reactions, pollution, or unanticipated biological effects caused by the presence of active biomolecules. The frequency of application and the concentration of active ingredients have a significant impact on overall risk.

Toxicological data, including studies on acute and chronic exposure, is analyzed to establish safe consumption limits. A margin of safety (MoS) is calculated to ensure that the product can be used under normal conditions without causing harm.

D. Regulatory safety requirements

The use of human-derived materials in cosmetics is tightly restricted in many places. Regulatory agencies have established guidelines to ensure the products' quality, safety, and ethical acceptability.

In general, safety rules concerning ingredient safety, labeling, and production processes must be followed by cosmetics. India's cosmetic laws are governed by the Drugs and Cosmetics Act and the Bureau of Indian Standards (BIS). International regulatory systems, including those in the US and the EU, impose further limitations on the use of chemicals derived from humans.

Manufacturers must submit safety data, toxicological profiles, and evidence of non-toxicity before a product is approved. Ethical considerations like sourcing and permission are also taken into consideration in some domains.

VII. ETHICAL AND REGULATORY CONCERNS

A. Ethical issues of fetal cell use

The primary ethical concern is the source of components made from fetal cells. These cells are typically produced from fetal tissue, which raises complex moral and ethical questions about consent, the source of the tissue, and the circumstances surrounding its acquisition.

There are questions about whether donors have given their informed consent and whether the use of such biological resources complies with ethical standards. Additionally, the commercialization of products derived from human fetal tissue is sometimes viewed as problematic, particularly in countries where the sanctity of human life is profoundly rooted in religion or culture.

Another ethical concern is labeling and marketing methods' transparency. The prospect that customers may not fully understand the origins of some substances raises questions about informed choice. Transparent communication, precise documentation, and ethical sources are essential to solving these issues.

©IJRASET: All Rights are Reserved | SJ Impact Factor 7.538 | ISRA Journal Impact Factor 7.894 |

B. Public perception

The popularity and success of cosmetic products containing components obtained from fetal cells are significantly influenced by public perception. Because of ethical, cultural, and religious sensitivities, the usage of human-derived materials, particularly those of fetal origin, may result in unfavorable consumer views.

Customers' desire to utilize these items may be impacted by their association of such components with contentious behaviors. However, because these compounds have been shown to have anti-aging and skin-regenerating properties, some groups of people may accept them.

Public trust can be increased through ethical sourcing, transparent ingredient disclosure, and appropriate education regarding the nature and safety of these ingredients. Marketing plans need to be carefully crafted to respect customer sensitivities and steer clear of deceptive claims.

C. Regulations in:

1) India

Under the Drugs and Cosmetics Act, 1940, and Rules, 1945, the Central Drugs Standard Control Organization oversees the regulation of cosmetic items in India. Additionally, the Bureau of Indian Standards offers guidelines for the quality and safety of cosmetics.

Although fetal cell-derived compounds in cosmetics are not specifically covered by any rules, manufacturers must guarantee product safety, appropriate labeling, and adherence to general cosmetic regulations. Additional attention may be applied to the usage of materials originated from humans, especially in relation to safety documentation and ethical sourcing.

2) USA

Under the Federal Food, Drug, and Cosmetic Act, the Food and Drug Administration oversees cosmetic items in the United States. Manufacturers are in charge of guaranteeing the safety of their products and materials, however the FDA does not need pre-market approval for cosmetics.

Although it is not expressly forbidden, the use of human-derived materials—including components produced from fetal cells—must adhere to safety and labeling regulations. Although they are not explicitly regulated, ethical factors may have an impact on industry practices and market acceptance.

3) Europe

Under EU Cosmetics Regulation (EC) No. 1223/2009, the European Commission regulates cosmetic products in Europe. Strict safety, quality, and labeling regulations are enforced by this system.

Consumer safety and ethical issues are highly valued in the European Union. Cosmetics makers are required to give thorough safety evaluations and paperwork, and the usage of materials derived from humans is closely monitored. Furthermore, the approval of substances obtained from fetal cells is frequently restricted in the European market due to ethical issues and public sensitivities.

VIII. ADVANTAGES AND LIMITATIONS

A. Advantages:

1) High efficacy

Growth factors, cytokines, peptides, and extracellular matrix proteins all contribute to the high biological activity of components produced from fetal cells. These bioactive substances have chemical interactions with skin cells that promote tissue healing, increase cellular proliferation [6–8], and stimulate collagen formation [1,3,4].

Their capacity to target several routes at once leads to better skin hydration, elasticity, and texture. These compounds have more noticeable and quick benefits than traditional cosmetic ingredients, especially in anti-aging and skin-rejuvenation applications.

2) Natural origin

The growing consumer demand for natural and bio-based beauty products is in line with the classification of fetal cell-derived compounds as biologically derived materials. These components are derived from biological sources and replicate natural processes that take place within the human body, in contrast to synthetic compounds.

When processed and manufactured correctly, this natural origin helps to improve biocompatibility and lower the likelihood of negative reactions. Furthermore, the move toward sophisticated biotechnology-based cosmetics is supported by the use of such chemicals.

©IJRASET: All Rights are Reserved | SJ Impact Factor 7.538 | ISRA Journal Impact Factor 7.894 |

3) Regenerative potential

The remarkable regeneration potential of components produced from fetal cells is one of their biggest benefits. Growth factors and signaling chemicals that support cell division, proliferation, and tissue remodeling are abundant in these materials.

Fetal cells are excellent in healing damaged skin and restoring its structural integrity because of their reputation for promoting quick, scarless healing. This regenerative capacity is very useful in formulations for skin regeneration, wound healing, and anti-aging treatments.

B. Limitations:

1) Ethical concerns

The origin and procurement of materials obtained from fetal cells provide significant ethical concerns. Public acceptance and regulatory decisions may be influenced by worries about informed consent, ethical issues, and the commercialization of biological resources originating from humans.

Such items may not be widely accepted in some areas due to cultural and religious beliefs. To solve these issues, ethical compliance and source transparency are crucial.

2) High cost

Complex and costly biotechnological processes are involved in the extraction, processing, purification, and stabilization of components obtained from fetal cells. Production expenses are further increased by maintaining sterile conditions, guaranteeing quality control, and putting advanced delivery systems in place.

Because of this, cosmetics made with these substances are frequently more expensive than traditional formulations, which may prevent a wider range of consumers from using them.

3) Regulatory restrictions

varied countries have varied regulations regarding the use of materials obtained from humans in cosmetics. The usage or commercialization of components produced from fetal cells may be limited by stringent safety evaluations, paperwork, and ethical considerations.

Manufacturers may find it difficult to get clearances in some areas due to regulatory ambiguity or a lack of clear guidelines. Product development and market expansion may be hampered by these limitations.

IX. FUTURE PERSPECTIVES

1) Alternative technologies (plant stem cells)

A viable and morally acceptable substitute for biological components derived from humans is plant stem cell technology. These cells, which are abundant in antioxidants, polyphenols, and other bioactive substances, are derived from plant sources such fruits, roots, and leaves.

By reducing free radical damage, extending cell life, and boosting skin vitality, plant stem cells provide protective and regenerative effects on the skin. They are frequently used to anti-aging formulas to increase the firmness, elasticity, and moisture of the skin. Additionally, their non-human origin avoids ethical issues and lowers the possibility of immunogenic reactions, increasing their acceptability to regulators and customers.

2) Synthetic biotechnology

Because synthetic biotechnology makes it possible to produce bioactive compounds using laboratory-based methods, it is a major achievement in the cosmetics sector. Peptides, growth factors, and other functional molecules that replicate the actions of naturally occurring biological substances are designed and synthesized using this method.

These synthetic bioactives provide a number of benefits, such as constant performance, regulated composition, and high purity. Additionally, they solve the ethical and safety issues related to materials obtained from fetal cells by doing away with the requirement for biological sourcing. The potential for creating specific and useful cosmetic compounds is further increased by the application of peptide engineering and recombinant DNA technology.

3) Market trends

Growing consumer knowledge and desire for safe, natural, and ethically derived goods are driving a major transition in the worldwide cosmetics sector. Clean-label cosmetics, which prioritize sustainability, transparency, and the lack of contentious ingredients, are becoming more and more popular.

Manufacturers are adopting alternative technologies like plant stem cells and synthetic bioactives as a result of consumers' growing preference for cruelty-free, vegan, and ecologically friendly products. Furthermore, the creation of novel goods that satisfy ethical and performance standards is being aided by developments in formulation science and marketing tactics.

©IJRASET: All Rights are Reserved | SJ Impact Factor 7.538 | ISRA Journal Impact Factor 7.894 |

Therefore, in order to ensure both product efficacy and consumer trust, the future of cosmetic formulations is probably going to concentrate on combining cutting-edge biotechnology with sustainable methods.

X. CONCLUSION

Human-derived cellular components offer novel approaches to skin regeneration and anti-aging applications, marking a major breakthrough in cosmetic research. Their abundance of growth factors, cytokines, and extracellular matrix proteins promotes better cellular proliferation, increased collagen synthesis, and efficient dermal remodeling, which results in noticeable changes in the texture, elasticity, and moisture of the skin. These bioactive ingredients' increasing significance in contemporary cosmeceuticals is demonstrated by their incorporation into cosmetic formulations such serums, creams, and repair treatments.

The use of human-derived materials presents significant safety, ethical, and regulatory issues that must be carefully addressed to assure consumer protection and compliance with international standards, notwithstanding their apparent usefulness. These delicate biomolecules are now more stable and delivered thanks to formulation technology advancements, but more study is needed to maximize their long-term safety and effectiveness.

Future prospects in this field are focused on the creation of safer and more sustainable substitutes, such as biomimetic substances and stem cells generated from plants, which might provide similar advantages with less moral dilemmas. All things considered, human-derived cellular components have a great deal of potential to influence next-generation cosmetics—as long as scientific advancement is matched with moral obligation and legal compliance.

XI. ACKNOWLEDGEMENT

The authors express their sincere gratitude to R. C. Patel Institute of Pharmaceutical Education & Research, Shirpur (Autonomous Institute) for providing the necessary facilities, guidance, and academic support to carry out this review work successfully.

The authors are highly thankful to Ms. Sonali K. Shambharkar (Guide) for her valuable guidance, continuous support, and encouragement throughout the preparation of this manuscript. Her insightful suggestions greatly contributed to the quality of this work.

The authors also extend their heartfelt thanks to Dr. M. V. Girase (Co-guide) for his constant motivation, expert advice, and technical assistance during the completion of this review article.

Finally, the authors acknowledge all faculty members and colleagues who directly or indirectly contributed to the successful completion of this work.

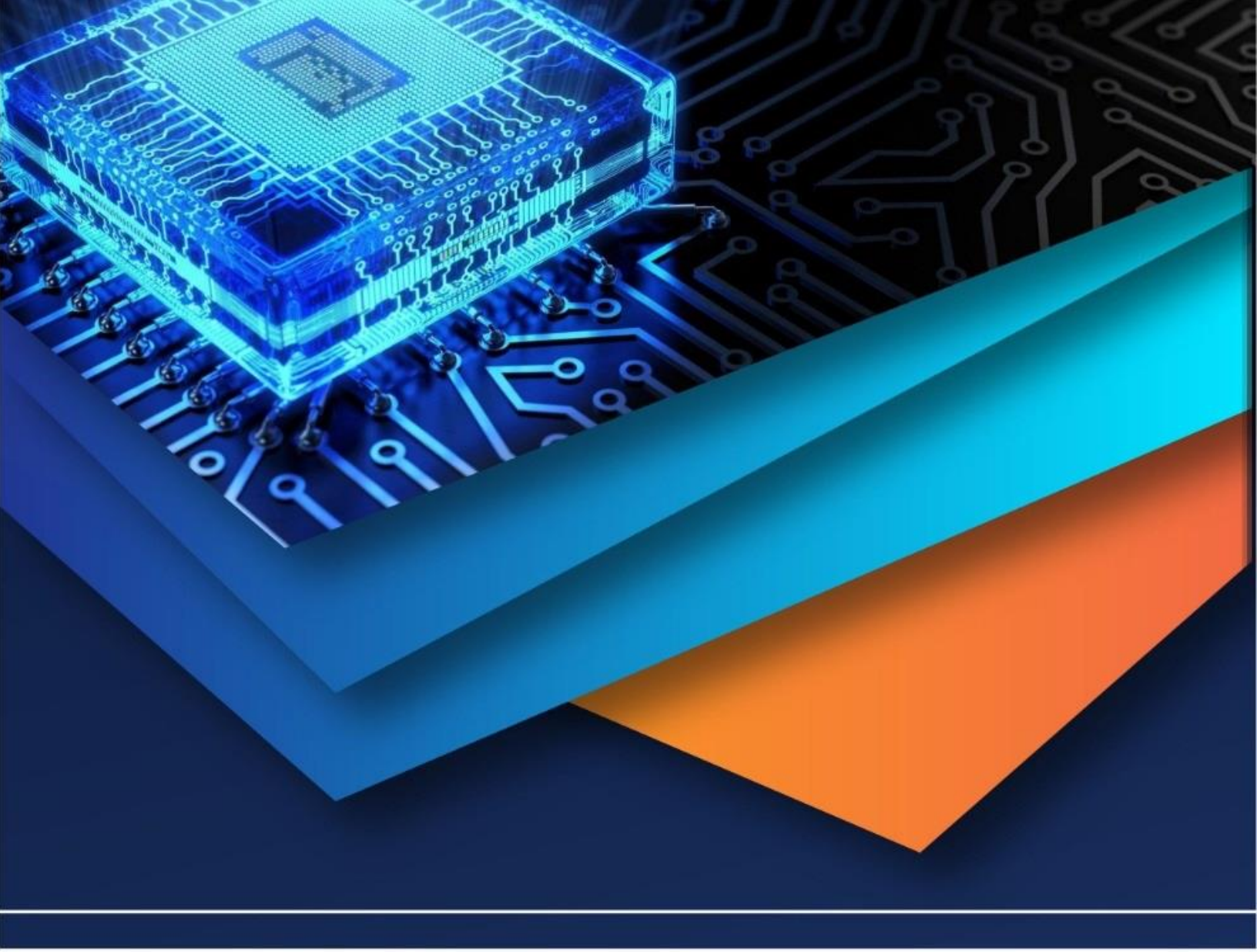
REFERENCES

[1] Kim DH, Lee SY, Kim SM. Effects of growth factors on skin regeneration. J Dermatol Sci. 2019;95(2):85–92.
[2] Lee HJ, Lim IJ, Lee MC. The role of fibroblasts in skin aging and repair. Int J Mol Sci. 2016;17(3):456–462.
[3] Uitto J. The role of collagen in skin aging. J Invest Dermatol. 2008;128(6):1415–1417.
[4] Quan T, Fisher GJ. Role of age-associated alterations in dermal fibroblasts. J Invest Dermatol. 2015;135(2):464–471.
[5] Rittié L, Fisher GJ. Natural and sun-induced aging of human skin. Cold Spring Harb Perspect Med. 2015;5(1):a015370.
[6] Park BS, Jang KA, Sung JH. Adipose-derived stem cells and their secretory factors in skin rejuvenation. Dermatol Surg. 2008;34(10):1323–1326.
[7] Kim WS, Park BS, Sung JH. Wound healing effect of adipose-derived stem cells. J Dermatol Sci. 2007;48(1):15–24.
[8] Chen L, Tredget EE, Wu PY. Paracrine factors of stem cells in wound healing. J Cell Physiol. 2008;217(3):749–758.
[9] Werner S, Grose R. Regulation of wound healing by growth factors and cytokines. Physiol Rev. 2003;83(3):835–870.
[10] Mast BA, Schultz GS. Interactions of cytokines, growth factors, and proteases in wound healing. Wound Repair Regen. 1996;4(4):411–420.
[11] Eming SA, Krieg T, Davidson JM. Inflammation in wound repair: molecular and cellular mechanisms. J Invest Dermatol. 2007;127(3):514–525.
[12] Gurtner GC, Werner S, Barrandon Y. Wound repair and regeneration. Nature. 2008;453(7193):314–321.
[13] Shukla A, Rasik AM, Dhawan BN. Growth factors in wound healing. Indian J Exp Biol. 1999;37(3):247–254.
[14] Phan TT, Lim IJ, Aalami O. Fetal fibroblasts exhibit enhanced healing properties. J Cell Biol. 2001;152(5):1119–1128.
[15] Lorenz HP, Longaker MT. Scarless skin wound repair in the fetus. N Engl J Med. 2008;339(12):813–817.
[16] Pipino C, Pandolfi A. Role of human fibroblasts in tissue regeneration and repair. Int J Mol Sci. 2015;16(5):11607–11623.
[17] Fuchs E. Skin stem cells and their role in regeneration. Nature. 2007;445(7130):834–842.
[18] Blanpain C, Fuchs E. Epidermal homeostasis and regeneration. Annu Rev Cell Dev Biol. 2009;25:163–192.
[19] Ghazizadeh S, Taichman LB. Organization of stem cells in human epidermis. J Invest Dermatol. 2005;124(2):367–372.
[20] Proksch E, Brandner JM, Jensen JM. Skin barrier function and repair. Exp Dermatol. 2008;17(12):1063–1072.
[21] Lintner K, Peschard O. Biologically active peptides in cosmetic formulations. Int J Cosmet Sci. 2000;22(3):207–218.
[22] Farris PK. Cosmeceutical growth factors: scientific evidence and clinical applications. Clin Dermatol. 2009;27(5):502–07.
[23] Gold MH. Use of growth factors in cosmetic dermatology. J Clin Aesthet Dermatol. 2011;4(5):30–34.
[24] Kim J, Kim WS, Park SH. Conditioned media from stem cells in cosmetic applications. J Cosmet Dermatol. 2013;12(2):96–102.
[25] Elsner P, Maibach HI. Cosmeceuticals and active cosmetics. Dermatol Ther. 2005;18(5):402–407.

©IJRASET: All Rights are Reserved | SJ Impact Factor 7.538 | ISRA Journal Impact Factor 7.894 |

[26] Parolini O, Soncini M. Human placenta-derived stem cells and their potential in regenerative medicine. Stem Cells Dev. 2006;15(5):785–795.
[27] Badiavas EV, Falanga V. Treatment of chronic wounds with stem cells. Expert OpinBiol Ther. 2003;3(2):175–182.
[28] European Commission. Regulation (EC) No 1223/2009 on cosmetic products. Off J Eur Union. 2009;342:59–209.
[29] US Food and Drug Administration. Guidance for industry: cosmetic safety. FDA. 2020.
[30] Bureau of Indian Standards. Indian standards for cosmetics. BIS. 2016.

©IJRASET: All Rights are Reserved | SJ Impact Factor 7.538 | ISRA Journal Impact Factor 7.894 |



10.22214/IJRASET

45.98

IMPACT FACTOR: 7.129

IMPACT FACTOR: 7.429

INTERNATIONAL JOURNAL FOR RESEARCH

IN APPLIED SCIENCE & ENGINEERING TECHNOLOGY

Call : 08813907089 (24*7 Support on Whatsapp)