

Control of GPCR Activity, Trafficking, And Localization by Proteins That Interact with GPCRs

Lokesh Kumar^{1*}, Deleshwar Kumar², Jay Kumar Chandra³

¹SSIPSR, Shri Shankaracharya Professional University, Chhattisgarh, India

²KIPS, Shri Shankaracharya Professional University, Bhilai, Chhattisgarh, India

³Raigarh College of Pharmacy, Raigarh, Chhattisgarh, India

*Corresponding Author E-mail: lokesh.pcology@gmail.com

Abstract:

G protein-coupled receptors (GPCRs) are integral in cell-cell communication and physiology, hence represent one of the most promising areas for drug targets. The regulatory proteins involved include GPCR kinases (GRKs), β -arrestins, scaffold proteins, and regulators of G protein signaling (RGS) proteins that modulate desensitization, internalization, recycling, and degradation of the receptors. Disruptions in GPCR signaling lead to numerous diseases in animals such as neurodegenerative diseases, cardiovascular diseases, and cancer. Animal models are very helpful for the study of GPCR regulation, with sophisticated techniques such as fluorescence microscopy, fluorescence resonance energy transfer (FRET), co-immunoprecipitation, and mass spectrometry applied to the study of receptor interaction and trafficking. GEMMs are used to further explain the physiological roles of GPCR-interacting proteins. The study shows the potential importance of targeting these regulatory proteins as a new way forward beyond traditional GPCR agonists and antagonists. This understanding of GPCR signaling in animals provides a new drug discovery path, with precise modulations at the receptor's activity level to improve treatments on veterinary fronts. This means immense implications for the improvement of animal health through next-generation therapeutics that have much greater specificity and fewer off-target effects.

Keywords: G Protein-Coupled Receptors, Trafficking, And Localization, GPCR Regulation, B-Arrestins, GPCR Kinases, RGS Proteins, Animal Models.

1. INTRODUCTION

GPCRs are the largest and most diverse family of membrane proteins in mammals and

other vertebrates, representing a significant fraction of all cellular communication and physiological regulation. They are involved in detecting the most varied extracellular

stimuli, such as neurotransmitters, hormones, peptides, lipids, and sensory signals including light and odors [1]. Activation of GPCRs triggers heterotrimeric G proteins, thereby activating intracellular signaling cascades controlling essential cellular functions, such as metabolism, the immune response,

neurotransmission, and cardiovascular functions. Due to their wide participation in physiological functions, GPCRs are seen as ideal therapeutic targets [2]. Nearly one-third of all FDA-approved drugs are directed against GPCRs, thus underlining the significance of GPCRs in medical science.

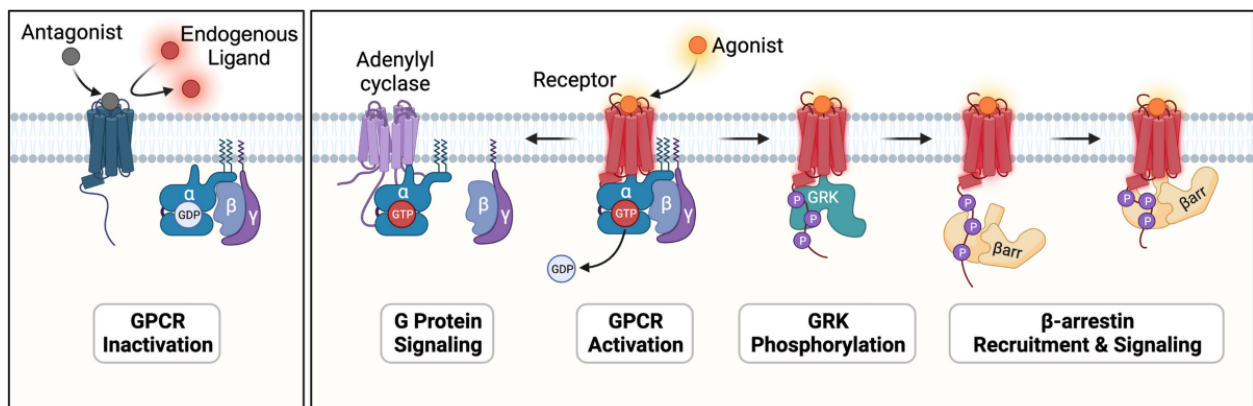


Figure 1: GPCR signaling [3]

GPCR activity does not involve the binding of the ligand only. Several proteins control the receptor's location and trafficking within cells, making them functional in the right environment. Among such proteins include the GPCR kinases (GRKs), β -arrestins, scaffold proteins, and adaptor molecules, which are all pivotal in the desensitization, internalization, recycling, or destruction of receptor activities [4]. They regulate, prolong, or end GPCR signaling to facilitate cellular responses appropriately. These regulatory interactions enable GPCRs to communicate with other pathways within cells for context-dependent and diverse responses [5].

Animals can suffer from neurodegenerative disorders, metabolic syndromes, cardiovascular diseases, and malignancies

because of disruptions in GPCR signaling and trafficking. Altered GPCR trafficking and signaling are associated with Alzheimer's, Parkinson's, and cardiovascular diseases. The knowledge of how GPCR-interacting proteins modulate receptor activity is essential for the development of new therapies. Recent breakthroughs in proteomics, structural biology, and high-resolution imaging have shed light on the molecular mechanics of these interactions, which may open new avenues for drug discovery beyond GPCR agonists and antagonists [6].

1.1. Background Information and Context

GPCRs in animal models exert complicated regulatory processes such as activation, internalization, recycling, or degradation. Important regulatory protein classes include

the G-protein coupled receptor kinases (GRKs), β -arrestins, scaffold proteins and adaptor proteins that control receptor activity through modulation of receptor phosphorylation, desensitization, and intracellular trafficking [7], [8]. These proteins do not only regulate GPCR signaling but also allow the receptor to cross-talk with other pathways thereby creating functional diversity for cellular responses [9]. Animal diseases, neurodegenerative disorders, cardiovascular diseases, and cancer have all been associated with disruptions in GPCR trafficking and signaling. Further progress in the proteomics era of structural biology and imaging has been made by extending our comprehension of GPCR-interacting proteins with a new range of opportunities in targeted therapies applied to animal models for eventual treatment of humans [10].

1.2. Objectives of the Review

This review aims to:

- To look at the important proteins that control GPCR trafficking, activity, and localization in animal models.
- To examine how these proteins, alter cellular distribution and GPCR signaling in animal tissues through molecular mechanisms.
- To talk about how these proteins, regulate GPCR in animal models and the physiological and pathological ramifications.
- highlight new developments and treatment approaches that target

GPCR-interacting proteins, using animal-based disease models.

1.3. Importance of the Topic

Biomedical research, in an animal model, would depend on insight into GPCR activity and regulation of trafficking. GPCRs are the largest class of pharmacological targets in animal studies, and hence it would be possible to advance therapy by further understanding regulation by interacting proteins. For several decades, drug research has focused on agonists and antagonists of GPCR, but regulatory proteins can modulate GPCR action that can be used to develop new therapy [11]. These methods might better animal illnesses by targeting and context-dependently modulating GPCRs. The diversity of animal species and their illness responses opens new opportunities for veterinary therapeutics in research into how regulatory proteins alter GPCR function in animals. Investigating these relationships in animal disease models can be helpful in the development of future human medicines [12]. It is crucial to understand the physiological roles of GPCRs in animal health to improve veterinary disease management and optimize therapies for diverse species.

2. KEY RESEARCH STUDIES ON GPCR

The functions of G protein-coupled receptors (GPCRs) are regulated by several interacting proteins: this is known to control the activity, trafficking, and localization of receptors within cells. Among the crucial regulators of GPCRs, there are arrestins and regulators of G protein signaling (RGS) proteins, which are

to be discussed further according to findings from animal models [13].

🚦 Arrestins: Key Modulators of GPCR Internalization and Signaling

Role in Receptor Internalization and Desensitization Arrestins, especially β -arrestins, play a critical role in regulating the activity of GPCRs after the activation of the receptor. Upon binding of the ligand and activation of the receptor, GPCRs undergo a conformational change that increases their susceptibility to binding by β -arrestin [14]. This interaction is essential for regulating receptor activity and trafficking, particularly in the internalization process.

In animal models, β -arrestins mediate receptor desensitization through the uncoupling of activated GPCRs from G proteins that effectively inhibits further signal transduction. It ensures that the signaling via the receptor remains finite and hence there would be no overstimulation of the cell. Internalization mediated by β -arrestin is clathrin-mediated endocytosis. This involves internalising the receptors that would eventually be recycled to the cell surface or degraded.

- **Desensitization** β -arrestins prevent continued signalling by intracellular pathways. Because they un-Couple GPCRs from their coupled G protein

-

- they contribute to deactivation of signaling cascade [15].

- **Receptor Recycling or Degradation:** The ultimate fate of the internalized GPCRs is decided by β -arrestins, which would either recycle it back to the plasma membrane or send it for degradation. These processes affect the overall receptor availability and, consequently, cellular signaling.

Role in scaffolding Apart from internalizing GPCRs, β -arrestins play the role of a scaffold protein in recruiting other signaling molecules. In the case of animal models, studies have demonstrated β -arrestin to bind Mitogen-Activated Protein Kinases (MAPKs) ERK1/2 and JNK, influencing processes related to cell growth, differentiation, and survival. Such dual functioning by β -arrestins—regulation of GPCR desensitization and recruitment to alternative pathways to modulate cell response—position β -arrestins centrally in control over the cellular response.

Arrestins also perform a scaffolding function, influencing receptor recycling, degradation, and cytoskeletal dynamics [16]. Thus, the significance of these proteins in regulating GPCR trafficking and signaling is further established.

Table 1: Research Studies on GPCR Regulation by Arrestins

References	Title	Topic Covered	Research Study
------------	-------	---------------	----------------

Bahouth, S. W., & Nooh, M. M. (2017) [17]	Barcoding of GPCR trafficking and signaling through the various trafficking roadmaps by compartmentalized signaling networks	GPCR Trafficking and Compartmentalized Signaling	Investigated how arrestins and other proteins modulate GPCR trafficking pathways and their compartmentalized signaling, using various cellular models.
Betke, K. M., Wells, C. A., & Hamm, H. E. (2012) [18]	GPCR mediated regulation of synaptic transmission	GPCR and Synaptic Transmission	Focused on the regulation of GPCRs in synaptic transmission and the role of arrestins in terminating signaling and regulating receptor internalization during neural signaling.
Hanyaloglu, A. C. (2018) [19]	Advances in membrane trafficking and endosomal signaling of G protein-coupled receptors	Membrane Trafficking and Endosomal Signaling	Explored the role of arrestins in endosomal GPCR signaling and the mechanisms behind receptor recycling and degradation in different cell types.
Kunselman, J. M., Lott, J., & Puthenveedu, M. A. (2021) [20]	Mechanisms of selective G protein-coupled receptor localization and trafficking	Selective GPCR Localization and Trafficking	Discussed how arrestins and other regulatory proteins mediate selective localization and trafficking of GPCRs, contributing to proper receptor function in cellular contexts.
Liccardo, F., Luini, A., & Di Martino, R. (2022) [21]	Endomembrane-based signaling by GPCRs and G-proteins	GPCR Endomembrane Signaling	Focused on how arrestins play a role in GPCR signaling at endosomal membranes, further enhancing the

			versatility of GPCR signaling in cellular compartments.
Paek, J., Kalocsay, M., Staus, D. P., Wingler, L., Pascolutti, R., Paulo, J. A., ... & Kruse, A. C. (2017) [22]	Multidimensional tracking of GPCR signaling via peroxidase-catalyzed proximity labeling	GPCR Signaling Tracking	Provided insights into tracking arrestin-GPCR interactions and receptor signaling pathways using proximity labeling techniques to study receptor endocytosis and signaling dynamics.
Wright, S. C., Lukasheva, V., Le Gouill, C., Kobayashi, H., Breton, B., Mailhot-Larouche, S., ... & Bouvier, M. (2021) [23]	BRET-based effector membrane translocation assay monitors GPCR-promoted and endocytosis-mediated Gq activation at early endosomes	GPCR Activation and Endocytosis	This study used BRET technology to investigate how arrestins regulate GPCR-promoted Gq activation during receptor internalization and subsequent signaling events.

Regulators of G Protein Signaling (RGS): Modulating GPCR Signaling and Trafficking

RGS Proteins in G Protein Deactivation RGS proteins represent another important category of proteins involved in the regulation of GPCR activity. These proteins are crucial in controlling the strength and duration of GPCR-mediated signaling by promoting the hydrolysis of GTP bound to G protein α subunits. This accelerates the deactivation of G proteins, thereby turning off the signaling cascade initiated by GPCR activation [24].

Studies with animal models have shown that the function of RGS proteins regulates cell response to activation by GPCRs tightly. For example, their function is important in the attenuation of overactivated signals through converting GTP back into GDP so as to stop potential pathological events, for example cardiac failure or disordered cell division. RGS Proteins and GPCR Trafficking More recent studies with animal models have also indicated that RGS proteins modulate GPCR trafficking [25]. For instance, the RGS protein RGS2 has been implicated in the trafficking of GPCRs between the plasma membrane and intracellular compartments. Such regulation

is important for receptor internalization and recycling.

RGS proteins exert their function at the modulation site of GPCR-G-protein interactions that ultimately affect signaling spatiality, that is the cellular location as well as the magnitude of signal events. Determining the cell-type-specific cellular responses based on localized receptor expression requires significant regulation of spatial location of signaling via GPCR's. Pathophysiological Implications of Dysregulated RGS Proteins Overexpression or mutation of RGS proteins has been linked to a number of pathophysiological conditions, ranging from heart failure and cancer to neurological disorders in the models using animals. RGS proteins are integral for balancing GPCR signaling and, therefore dysregulation of the proteins leads to abnormal activation of receptors that contribute to progressing to disease.

3. METHODOLOGIES AND FINDINGS IN GPCR REGULATION BY INTERACTING PROTEINS

Much of the research into G-protein coupled receptors (GPCRs) and the regulation of these receptors by interacting proteins has gone forward with the use of animal models. These models provide a physiological context essential for understanding the complex roles of proteins involved in GPCR trafficking, localization, and signaling [26]. The next section is devoted to discussing methodologies and some key findings, mainly focusing on strategies for animal-based approaches to studying GPCR regulation.

A. Fluorescence Microscopy and FRET in Animal Models

One of the techniques for investigating the GPCR regulatory processes that have recently been quite common is fluorescence microscopy, which makes special mention of the fluorescence resonance energy transfer or FRET method.

Animal-based Application

- G-protein-coupled receptors can be labeled with fluorescent markers in genetically modified mice or rats to observe their behavior in living animals.
- Using FRET, scientists can observe the interactions of GPCRs with their interacting proteins in a specific tissue (such as brain, liver, or heart). This will allow for the monitoring of receptor internalization and desensitization processes in real time under various physiological states [27].

Such as, FRET studies have in fact illuminated how β -arrestins associate with the internalizing GPCRs and further altered the recycling of the receptor. These discoveries give this insight into the regulation of GPCRs in living systems in terms of its temporal and spatial principles.

B. Co-immunoprecipitation (Co-IP) in Animal Models

Co-immunoprecipitation is a highly sensitive method that allows for the identification of physical interactions between GPCRs and their interacting proteins [28]. It can be used to examine protein complexes in the study of

receptor trafficking and signaling in animal-based studies.

Animal-based Application:

- It is also possible to get animal tissues such as brain slices or heart muscle, in genetically engineered mice or rats that express tagged GPCRs.
- Specific antibodies for pulling down GPCRs or their interactors are employed, and with the help of this, investigators can identify *in vivo* the whole range of proteins associated with GPCRs.

Recent advances in research by using Co-IP in animal models have highlighted several critical proteins involved in the process of GPCR internalization, recycling, and signaling: β -arrestins, G proteins, and AP-2 adaptor proteins. This will be significant in understanding the regulation of GPCR function in the whole-organism context.

C. Mass Spectrometry in Animal Models

MS is a highly sensitive technique used to identify proteins and their interacting molecules [29]. If coupled with immunoprecipitation, the technique can be applied to identify novel interacting proteins in the protein interactome of GPCRs in animal models, thus revealing some proteins that control the function of GPCRs.

Animal-based Application:

- MS can be utilized for the examination of animal tissues like knockout or transgenic mouse models in order to

find proteins engaged in GPCR trafficking and signaling.

- The method has also been applied in the identification of regulatory proteins, including kinases, phosphatases, and ubiquitination enzymes that regulate GPCR stability and activity *in vivo*.

For example, MS-based analysis in animal models has helped to reveal post-translational modifications, including those by phosphorylation and ubiquitination, of GPCRs that play crucial roles in their regulation during endocytosis, signaling, and recycling processes.

D. Genetically Engineered Mouse Models and RNA Interference (RNAi)

Genetically engineered mouse models are of utmost value for studying the functional roles of interacting proteins in the context of the whole organism. Such models allow the disruption of specific genes involved in GPCR regulation and make it possible to study their physiological roles *in vivo* [30].

Animal-based Application:

- Knock-out or knock-in mouse models were used to dissect the consequences of ablation or alterations in genes encoding inter-acting proteins, such as β -arrestins, G proteins, or adaptor proteins such as AP-2.
- Use RNA interference (RNAi) in animal models to knockdown genes responsible for the regulation of GPCRs to understand the functional outcome of a

disrupted specific protein-protein interaction.

For example, in animal models, RNAi-based studies have revealed the crucial role of β -arrestins in regulating receptor desensitization and internalization. It was shown that the functions of β -arrestins extend not only to the regulation of receptors but also to activating alternative pathways for signaling that have a positive impact on cell survival and gene expression.

Key Findings in Animal-Based GPCR Regulation Research:

- β -arrestins as Scaffolding Proteins:** Studies in animal models have further identified that β -arrestins, along with participating in receptor desensitization and internalization, also work as scaffolding proteins that can support alternative signaling pathways, influence survival of cells, gene expression, and resensitization of receptors.

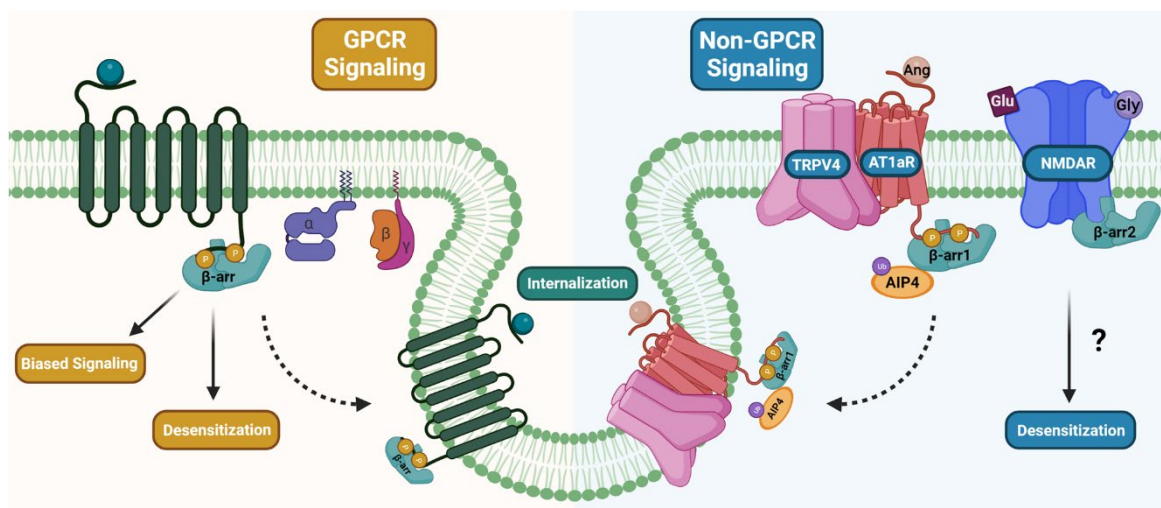


Figure 2: β -arrestins [31]

- Role of G Proteins and RGS Proteins:** Research with animals has defined the role of G proteins and their regulators, like RGS proteins, in the modulation of GPCR signaling in terms of both duration and intensity. The RGS proteins control the deactivation of G proteins, thereby ensuring proper outcomes in different tissues.
- GPCR proteins use adaptor proteins like AP-2** in regulating their trafficking for internalization and recycling. This, therefore, influences the availability of receptors at the cell surface, and as a result, it determines the responsiveness of the cells to such external signals.
- Physiological roles of in-vivo interacting proteins:** The ability to

genetically modify mouse models through RNAi methodologies has allowed animals to provide definitive in-vivo evidence for many of the pivotal roles of protein-protein interacting proteins, β -arrestin, G protein, and their adaptor proteins during GPCR biology. Such data have been confirmed to indicate modifications through these interactions toward receptor localization dynamics of overall cellular responses as well.

4. THEMATIC ANALYSIS OF GPCR REGULATION: TRAFFICKING, RECYCLING, DEGRADATION, AND LOCALIZATION

The importance of animal-based models lies in studying the regulation of GPCRs, such as understanding the trafficking, recycling, degradation, and localization processes involved in these molecules [32]. Animal models offer the facility for controlled experiments to explore intricate molecular mechanisms controlling GPCR behavior in living organisms. Here's a summary of how animal-based experiments can be utilized to understand GPCR regulation:

1. Trafficking and Internalization in Animal Models

Animal models, including genetically engineered mice, have become instrumental in studies concerning GPCR trafficking and internalization. In particular, the knockout mice with the deficiency of the key protein like β -arrestins or AP-2 will help in testing the role played by these proteins in receptor

internalization. For such studies, the movement of tagged GPCRs can be monitored within tissues or cells isolated from animals through live-cell imaging and fluorescence microscopy techniques. Such models permit researchers to observe the GPCR endocytosis dynamics under physiological conditions.

A typical experiment would be to activate a specific GPCR in an animal model, and then observe the internalization of the receptor using fluorescence resonance energy transfer or other imaging methods. The contribution of specific proteins, such as β -arrestins or dynamin, in the process could be assessed by comparison of wild-type animals with knockouts lacking such proteins.

2. Recycling and Degradation in Animal Models

After internalization, GPCRs are either recycled to the cell surface or targeted for degradation. These two processes may be balanced using animal models. Mouse models carrying mutations in recycling proteins, such as Rab GTPases, or in degradation proteins, such as E3 ubiquitin ligases, can be used to determine how these proteins regulate receptor fate [33].

For instance, the use of tracking of internalized receptors in living animals post activation for subsequent assessment whether these receptors were recycled back to the membrane or degraded in lysosomes has proven to be widely applied. GPCRs can be fluorescently tagged, and real-time measurement of receptor recycling or degradation efficiency can be ascertained

using tissues from the animals. Besides that, specific pathways can be disrupted in animal models using pharmacological inhibitors or RNA interference to study the downstream effects on GPCR trafficking and degradation.

3. Localization and Signaling in Animal Models

The localization of GPCRs in lipid rafts within specific cellular compartments is

essential for their signaling outcome. This study can be done in an animal model by introducing GPCRs into various tissues using biochemical techniques for the isolation of lipid raft fractions. The interaction of GPCRs with components of rafts, such as caveolins and flotillins, during co-localization studies employing living animals and fluorescence microscopy.

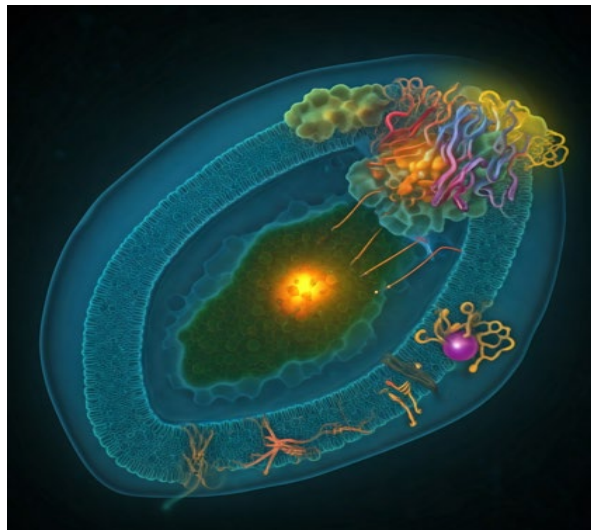


Figure 3: GPCR Localization: An Animal Model Perspective [34]

For example, through the use of transgenic animals that express tagged GPCRs in particular cell types, *in vivo* visualization of receptor localization can be possible. This could especially be valuable for tissues like the brain where GPCR signaling plays a crucial role in neuronal communication. Animal model studies can clarify the mechanism through which the lipid rafts' location of GPCRs can impact signaling pathways such as specific G protein recruitment and other downstream molecules.

5. DISCUSSION

5.1. Interpretation and Analysis of Findings

The review further states that the regulation of GPCR (G-Protein Coupled Receptor) in animals is complex, since interacting proteins are necessary for receptor activity control, trafficking, and cellular responses. GPCRs are involved in various physiological processes, and precisely, the network of proteins in their operation determines how these receptors work in animal cells. β -Arrestins, GPCR kinases, scaffold proteins, and adaptor molecules regulate the GPCRs most

dominantly [35]. The mechanisms ensure proper GPCRs are activated precisely in the right instances as some of them like desensitization, internalization, recycling, and even degradation occur appropriately.

β -Arrestins play a pivotal role in terminating G protein-mediated signaling in animal cells. Bound to phosphorylated receptors, they block the further activation of G proteins, effectively switching off the pathway. Additionally, β -arrestins bring other signaling molecules to the membrane, activating alternative pathways like MAPKs, which influence various important cellular functions, including growth, differentiation, and survival, in diverse tissues throughout animals. Regulators of G Protein Signaling (RGS) proteins accelerate the deactivation of G proteins and ensure that the signaling is terminated immediately to avoid overstimulation of the cell while maintaining homeostasis.

In particular, the review emphasizes the complicated regulation of GPCR signaling in animals. Such proteins are involved in not only regulating the timing of the function of a

receptor but also modulating spatial dynamics of trafficking receptors within a cell [36]. Thus, they help to manage where exactly a receptor locates in a variety of cellular compartments such that the relevant signals occur there. Such spatial regulation is important for the context-dependent nature of GPCR signaling in different tissues and in various physiological environments, which have to be fine-tuned to meet the particular needs of the animal.

The work, also, points out the therapeutic value in targeting the proteins as potential modulators for veterinary drugs and animal health. While traditional drug development targets the GPCRs directly through agonists and antagonists, the modulation of the interacting proteins for GPCR regulation is a more alternative and refined approach. Targeting the proteins involved in GPCR signaling and trafficking could lead to novel therapeutic strategies for diseases in animals caused by dysregulated GPCR activity, from neurological disorders to cardiovascular and immune disorders.

Table 2: Key Regulatory Proteins Involved in GPCR Signaling in Animals [37]

Regulatory Protein	Function	Effect on GPCR Signaling	Potential Therapeutic Target
β-Arrestins	Bind to phosphorylated receptors, terminating G protein signaling	Prevent further activation of G proteins; initiate alternative signaling pathways like MAPKs.	Targeting β -arrestin interactions for alternative signaling regulation.

GPCR Kinases (GRKs)	Phosphorylate activated GPCRs, marking them for desensitization	Control the activation and desensitization of GPCRs, ensuring timely termination of signaling.	Inhibition of GRK activity to prolong GPCR signaling in specific animal diseases.
Regulators of G Protein Signaling (RGS)	Accelerate the deactivation of G proteins	Ensure rapid termination of signaling, preventing overstimulation and maintaining cellular balance.	Modulation of RGS proteins to regulate G protein signaling pathways.
Scaffold Proteins	Provide structural support for signaling complexes	Coordinate the assembly of protein complexes, ensuring that signaling occurs at the correct cellular location.	Targeting scaffold protein interactions to modify GPCR trafficking.
Adaptor Molecules	Facilitate the assembly of protein complexes and receptor trafficking	Ensure proper receptor localization and internalization, affecting the spatial dynamics of signaling.	Modulating adaptor protein function to influence receptor localization and internalization.

Furthermore, the integral understanding of the functions of these regulatory proteins in the future would result in the focus of veterinary therapeutic practices on selectively targeting particular proteins or interactions to modulate GPCR signaling in animals. This could provide a more contextual and precise strategy in improving their health to better manage diseases associated with GPCR dysfunction in various animal species.

5.2. Implications and Significance

Such implications have very strong importance in the field of veterinary medicine and animal health [38]. Here, GPCR (G-Protein Coupled Receptor) signaling plays a

key role in controlling different physiological functions. In animals, GPCRs play important roles in processes including neurotransmission, immune response, and cardiovascular function. Given the central position of GPCRs in preserving normal animal physiology, dysregulation of GPCR signaling has been implicated in a wide array of diseases affecting animals, from neurological disorders to cardiovascular diseases and even cancer.

In animals, GPCR signaling misregulation has been implicated in several diseases. For example, in neurodegenerative diseases in dogs, such as canine cognitive dysfunction or

certain models of Parkinson's disease, aberrant GPCR activity is thought to play a role in the progression of these diseases. GPCR signaling has been demonstrated to modulate heart function and responsiveness to therapy in cardiovascular diseases in pets, including heart failure or arrhythmias. In veterinary oncology, GPCR dysfunction has been implicated in the process of tumor metastasis in animals with cancer, where inappropriate signaling promotes the growth of the tumor and makes it resistant to chemotherapy [39]. There is a possibility that targeting the regulatory proteins involved in GPCR signaling may correct the dysregulated pathways and provide new therapeutic interventions for these animal diseases.

The findings open the door for developing therapies not only directly on the GPCRs but also on the regulatory proteins that control their activity. This may provide more targeted and precise modulation of GPCR signaling in animals. In contrast to traditional treatments, which are often non-specific to modulate GPCR activity, this approach would enable more refined control over the regulatory proteins involved in receptor regulation. This could lead to fewer off-target effects in veterinary drugs and better therapeutic outcomes, especially for complex diseases in animals where there is a requirement for precise modulation of GPCR signaling.

Moreover, the capacity to regulate GPCR signaling through regulatory proteins might have very broad applications in the discovery of animal drugs. Among the most significant and diverse families of drug targets in veterinary medicine are GPCRs. A deeper

understanding on how to modulate these pathways could lead to the development of the next generation of veterinary drugs for a wide variety of diseases in animals [40]. This drug development approach would be particularly useful in the treatment of relatively hard-to-manage conditions with traditional GPCR-targeting drugs, some of which include certain forms of cancer, autoimmune diseases in animals, and neurological disorders. By targeting the regulatory proteins controlling GPCR activity, we can design more effective and targeted therapy for many diseases affecting animals, making them healthier and better off overall.

5.3. Gaps and Suggestions for Future Research Directions

Gaps in Current Research

1. **Unexplored Molecular Mechanisms:** Significant advancements have been made, but there is still an incomplete appreciation of the molecular mechanisms guiding animal GPCRs and their regulatory proteins. Although some of these regulatory proteins have been identified in different animal species, the spectrum of interacting proteins, and their functions, are incompletely delineated. This gap in knowledge confines our capacity to define the way these proteins regulate GPCR activity, localization, and trafficking in animal cells.
2. **Role of Post-Translational Modifications:** In animals, the extent

of post-translational modifications such as phosphorylation, ubiquitination, and SUMOylation of GPCRs and its interaction with other regulatory proteins remains to be fully understood. It could be crucial in modulating the receptor function and dynamics of interactions, and thus a closer understanding might open novel avenues for drug development in veterinary medicine against diseases in animals.

- 3. Variation in Animals-Specific Tissues:** Signaling through GPCRs varies from one tissue to another in animals. The tissue-specific functions of regulatory proteins such as β -arrestins may have an influence on receptor signaling, which could differ in animals depending on the tissue involved, including the brain, heart, or liver. These tissue-specific roles, especially in animal models, could provide insight into more customized veterinary therapies. However, the current research does not sufficiently evaluate the variation in these roles across different animal tissues and physiological conditions.
- 4. Sufficient In Vivo Animal Model Validation:** Though the animal models have provided valuable insights, there is a need for further in vivo validation of GPCR-interacting proteins and their effects on disease progression in animal species. The models in use may not exactly represent the pathophysiological

conditions found in animals with natural diseases. Thus, the need to develop models that more closely resemble the diseases found in animals is important in understanding the precise mechanisms at play.

- 5. Technological Limitations in Animal Research:** The animal-based research on GPCR-regulatory protein interactions can be advanced by more sophisticated technologies. Techniques like proteomics, structural biology, and live-cell imaging have demonstrated potential but remain largely unexploited in the context of animal GPCR signaling. Enhanced application of these technologies may further reveal new protein partners and more clearly elucidate the molecular mechanisms underlying GPCR regulation in animals.

Suggestions for Future Research Directions

- 1. Comprehensive Analysis of the Post-Translational Modification in Animals:** Future studies into animal models will be based on post-translational modifications. In-depth analysis about how phosphorylation, ubiquitination, and SUMOylation influence animal GPCR function may come up with newer therapeutic approaches against veterinary diseases of GPCR origins. This could develop better treatments against neurological diseases in dogs and feline heart failure.

2. **Tissue-Specific Animal Model Studies:** This is one of the significant areas of research: how regulatory proteins influence GPCR signaling in animal tissues. Determination of whether β -arrestins are involved in heart function in dogs or brain transmission in rats may reveal tissue-specific functions of the regulatory proteins. Such studies will be applied to design more targeted and effective medicines for animals, thus reducing off-target effects and increasing precision.
3. **Advanced Animal Models:** Develop more robust animal models that better approximate human situations. Improvement of canine cancer models, horse neurological illnesses, and feline heart diseases may better approximate human situations for researchers to understand how GPCR-interacting proteins affect the progression of animal illness and improve therapeutic methods.
4. **Enhanced techniques in proteomics, structural biology, and live-cell imaging:** Improved proteomics, structural biology, and live-cell imaging approaches will help in future research with animal-based GPCR studies. Such approaches could be adopted for studying GPCRs interactions with their regulatory proteins within various tissues of animals that reveal molecular mechanisms for health or sickness in animals. In the study with live-cell

imaging of animals, some unsuspected receptor trafficking and interaction might emerge.

5. **Identifying Targets for Veterinary Drug Development:** Armed with a deeper understanding of how regulatory networks modulate animal GPCR activity, future research should identify druggable targets. This may ultimately result in novel veterinary drugs that modulate the activity of regulatory proteins, offering new therapeutic avenues for GPCR-related disorders. It could revolutionize the treatment of pet cancer, autoimmune, and neurological diseases by developing small molecules or biologics that regulate regulatory proteins.

6. CONCLUSION

The significance of GPCRs and their regulatory proteins in maintaining cellular signaling homeostasis is underscored. Advanced animal models have extensively explored the complex mechanisms of GPCR signaling, trafficking, and regulation, providing crucial insights into the molecular interactions governing these processes. The presence of β -arrestins, GRKs, and RGS proteins in regulating receptor activity, desensitization, and recycling unfolds a potential as pharmacological targets for illnesses associated with inappropriately regulated GPCR signaling. Such findings go beyond a deeper understanding of GPCR biology and open up new possibilities in targeted interventions in veterinary medicine, offering more accurate and effective

treatments in the veterinary field of animals' treatment. Such attention to regulatory proteins provides new prospects for modulation of GPCR signaling in the way towards a new generation of drugs, side effects will be fewer and therapeutically effectiveness higher.

REFERENCES

1. Crilly, S. E., & Puthenveedu, M. A. (2021). Compartmentalized GPCR signaling from intracellular membranes. *The Journal of Membrane Biology*, 254, 259-271.
2. Irannejad, R., Pessino, V., Mika, D., Huang, B., Wedegaertner, P. B., Conti, M., & Von Zastrow, M. (2017). Functional selectivity of GPCR-directed drug action through location bias. *Nature chemical biology*, 13(7), 799-806.
3. Maurice, P., Guillaume, J. L., Benleulmi-Chaachoua, A., Daulat, A. M., Kamal, M., & Jockers, R. (2011). GPCR-interacting proteins, major players of GPCR function. *Advances in pharmacology*, 62, 349-380.
4. Milligan, G. (2010). The role of dimerisation in the cellular trafficking of G-protein-coupled receptors. *Current opinion in pharmacology*, 10(1), 23-29.
5. Lobingier, B. T., & von Zastrow, M. (2019). When trafficking and signaling mix: How subcellular location shapes G protein-coupled receptor activation of heterotrimeric G proteins. *Traffic*, 20(2), 130-136.
6. Jean-Alphonse, F., & Hanyaloglu, A. C. (2011). Regulation of GPCR signal networks via membrane trafficking. *Molecular and cellular endocrinology*, 331(2), 205-214.
7. Fu, Q., & Xiang, Y. K. (2015). Trafficking of β -adrenergic receptors: Implications in intracellular receptor signaling. *Progress in molecular biology and translational science*, 132, 151-188.
8. Van Anthony, M. V., Cuevas, S., Zheng, X., & Jose, P. A. (2016). Localization and signaling of GPCRs in lipid rafts. In *Methods in cell biology* (Vol. 132, pp. 3-23). Academic Press.
9. Toneatti, R., Shin, J. M., Shah, U. H., Mayer, C. R., Saunders, J. M., Fribourg, M., ... & González-Maeso, J. (2020). Interclass GPCR heteromerization affects localization and trafficking. *Science signaling*, 13(654), eaaw3122.
10. Romero, G., von Zastrow, M., & Friedman, P. A. (2011). Role of PDZ proteins in regulating trafficking, signaling, and function of GPCRs: means, motif, and opportunity. *Advances in pharmacology*, 62, 279-314.
11. Ray, K. (2015). Calcium-sensing receptor: trafficking, endocytosis, recycling, and importance of

- interacting proteins. *Progress in molecular biology and translational science*, 132, 127-150.
12. Bhosle, V. K., Rivera, J. C., & Chemtob, S. (2019). New insights into mechanisms of nuclear translocation of G-protein coupled receptors. *Small GTPases*, 10(4), 254-263.
 13. Bockaert, J., Perroy, J., Bécamel, C., Marin, P., & Fagni, L. (2010). GPCR interacting proteins (GIPs) in the nervous system: Roles in physiology and pathologies. *Annual review of pharmacology and toxicology*, 50(1), 89-109.
 14. Eichel, K., & von Zastrow, M. (2018). Subcellular organization of GPCR signaling. *Trends in Pharmacological Sciences*, 39(2), 200-208.
 15. Weinberg, Z. Y., & Puthenveedu, M. A. (2019). Regulation of G protein-coupled receptor signaling by plasma membrane organization and endocytosis. *Traffic*, 20(2), 121-129.
 16. Pavlos, N. J., & Friedman, P. A. (2017). GPCR signaling and trafficking: the long and short of it. *Trends in Endocrinology & Metabolism*, 28(3), 213-226.
 17. Bahouth, S. W., & Nooh, M. M. (2017). Barcoding of GPCR trafficking and signaling through the various trafficking roadmaps by compartmentalized signaling networks. *Cellular signalling*, 36, 42-55.
 18. Betke, K. M., Wells, C. A., & Hamm, H. E. (2012). GPCR mediated regulation of synaptic transmission. *Progress in neurobiology*, 96(3), 304-321.
 19. Hanyaloglu, A. C. (2018). Advances in membrane trafficking and endosomal signaling of G protein-coupled receptors. *International review of cell and molecular biology*, 339, 93-131.
 20. Kunselman, J. M., Lott, J., & Puthenveedu, M. A. (2021). Mechanisms of selective G protein-coupled receptor localization and trafficking. *Current opinion in cell biology*, 71, 158-165.
 21. Liccardo, F., Luini, A., & Di Martino, R. (2022). Endomembrane-based signaling by GPCRs and G-proteins. *Cells*, 11(3), 528.
 22. Paek, J., Kalocsay, M., Staus, D. P., Wingler, L., Pascolutti, R., Paulo, J. A., ... & Kruse, A. C. (2017). Multidimensional tracking of GPCR signaling via peroxidase-catalyzed proximity labeling. *Cell*, 169(2), 338-349.
 23. Wright, S. C., Lukasheva, V., Le Gouill, C., Kobayashi, H., Breton, B., Mailhot-Larouche, S., ... & Bouvier, M. (2021). BRET-based effector membrane translocation assay monitors GPCR-promoted and endocytosis-mediated Gq activation at early endosomes. *Proceedings of the*

- National Academy of Sciences, 118(20), e2025846118.
24. Rozenfeld, R., & Devi, L. A. (2011). Exploring a role for heteromerization in GPCR signalling specificity. *Biochemical Journal*, 433(1), 11-18.
25. Tian, X., Kang, D. S., & Benovic, J. L. (2014). β -arrestins and G protein-coupled receptor trafficking. *Arrestins-Pharmacology and Therapeutic Potential*, 173-186.
26. Tsvetanova, N. G., Irannejad, R., & von Zastrow, M. (2015). G protein-coupled receptor (GPCR) signaling via heterotrimeric G proteins from endosomes. *Journal of Biological Chemistry*, 290(11), 6689-6696.
27. Kang, D. S., Tian, X., & Benovic, J. L. (2014). Role of β -arrestins and arrestin domain-containing proteins in G protein-coupled receptor trafficking. *Current opinion in cell biology*, 27, 63-71.
28. Kennedy, J. E., & Marchese, A. (2015). Regulation of GPCR trafficking by ubiquitin. *Progress in molecular biology and translational science*, 132, 15-38.
29. Calebiro, D., & Koszegi, Z. (2019). The subcellular dynamics of GPCR signaling. *Molecular and cellular endocrinology*, 483, 24-30.
30. Dunn, H. A., & Ferguson, S. S. (2015). PDZ protein regulation of G protein-coupled receptor trafficking and signaling pathways. *Molecular pharmacology*, 88(4), 624-639.
31. Gurevich, V. V., & Gurevich, E. V. (2019). GPCR signaling regulation: the role of GRKs and arrestins. *Frontiers in pharmacology*, 10, 125.
32. Calebiro, D., & Godbole, A. (2018). Internalization of G-protein-coupled receptors: Implication in receptor function, physiology and diseases. *Best practice & research Clinical endocrinology & metabolism*, 32(2), 83-91.
33. Calebiro, D., Nikolaev, V. O., Persani, L., & Lohse, M. J. (2010). Signaling by internalized G-protein-coupled receptors. *Trends in pharmacological sciences*, 31(5), 221-228.
34. Varandas, K. C., Irannejad, R., & von Zastrow, M. (2016). Retromer endosome exit domains serve multiple trafficking destinations and regulate local G protein activation by GPCRs. *Current Biology*, 26(23), 3129-3142.
35. Hislop, J. N., & von Zastrow, M. (2011). Role of ubiquitination in endocytic trafficking of G-protein-coupled receptors. *Traffic*, 12(2), 137-148.
36. Wang, G., & Wu, G. (2012). Small GTPase regulation of GPCR anterograde trafficking. *Trends in pharmacological sciences*, 33(1), 28-34.

37. Xu, X., Meckel, T., Brzostowski, J. A., Yan, J., Meier-Schellersheim, M., & Jin, T. (2010). Coupling mechanism of a GPCR and a heterotrimeric G protein during chemoattractant gradient sensing in Dictyostelium. *Science signaling*, 3(141), ra71-ra71.
38. Irannejad, R., & Von Zastrow, M. (2014). GPCR signaling along the endocytic pathway. *Current opinion in cell biology*, 27, 109-116.
39. Lämmermann, T., & Kastenmüller, W. (2019). Concepts of GPCR-controlled navigation in the immune system. *Immunological reviews*, 289(1), 205-231.
40. Magalhaes, A. C., Dunn, H., & Ferguson, S. S. (2012). Regulation of GPCR activity, trafficking and localization by GPCR-interacting proteins. *British journal of pharmacology*, 165(6), 1717-1736.