COLLEGIATE BOARD RESOLUTION – RDC No. 843 OF 22 FEBRUARY 2024

Provides for the regularization of food and packaging under the jurisdiction of the Brazilian Health Surveillance System (SNVS) intended for supply within Brazilian territory.

The Collegiate Board of Directors of the Brazilian Health Regulatory Agency, in the use of the attributions vested in it under Article 7, items III and IV, and Article 15, items III and IV of Law no. 9,782 of 26 January 1999, and item VI, paragraph 1 of Article 187 of the Internal Regulation approved by Collegiate Board Resolution – RDC no. 585 of 10 December 2021, adopts the following Resolution, as decided upon in a meeting held on 21 February 2024, and I, Director-President, determine its publication.

CHAPTER I

INITIAL PROVISIONS

Article 1. This Resolution provides for the regularization of food and packaging under the jurisdiction of the Brazilian Health Surveillance System (SNVS, in Portuguese) intended for supply within Brazilian territory.

Sole paragraph. The regularization referred to in the caption of this article includes the procedures for:

I – marketing authorization, post-marketing authorization changes, revalidation of marketing authorization, and cancellation of marketing authorization;

II – notification, change of notification, maintenance of notification, and cancellation of notification; and

III – communication of start of manufacture or import, change of communication, and cancellation of communication.

Article 2. For the purposes of this Resolution, the following definitions apply:

I – food: any substance that is ingested in its natural state, semi-processed or processed, intended for human consumption, including beverages and any other substance used in its production, preparation or treatment, excluding cosmetics, tobacco and substances used solely as medicines;

II – *in natura* food: any food of plant or animal origin, whose immediate consumption requires only the removal of the inedible part and the treatments indicated for its perfect hygiene and conservation;

III – post-marketing authorization changes: changes made to the product's marketing authorization, after it has been granted by Anvisa;

IV – control analysis: analysis performed on the food or packaging after its regularization, when delivered for consumption, with the objective of proving its compliance with the technical regulations that establish its composition, quality, safety, and labeling requirements;

V – notice of start of manufacture or import: communication made by the manufacturer or importer to the competent state or municipal health authority so that it is aware of the products manufactured or imported by them;

VI – packaging: article that is in direct contact with the food and that is intended to protect it from alterations, contamination, and adulteration, from its manufacture until its delivery to the consumer;

VII – food equipment: any article in direct contact with food that is used during the preparation, portioning, storage, commercialization, and consumption of food, such as, for example, containers, machines, conveyor belts, pipes, equipment, accessories, valves and utensils, among others;

VIII – specification: set of documented requirements of a product that describe its characteristics, in a qualitative and quantitative manner, the acceptable variability and the purity requirements, among other parameters, for its standardization and unequivocal identification;

IX – manufacturer: company that has the necessary facilities for the manufacture of food or packaging, authorized by the competent health authority;

X – ingredient: any substance, including food additives, that is used in the manufacture or preparation of food and that is present in the final product in its original or modified form;

XI – health inspection: procedure carried out by the competent health authority to verify compliance with current legislation;

XII – food raw material: any substance in its raw state that, in order to be used as food, needs to undergo treatment and/or transformation of a physical, chemical, or biological nature;

XIII – interdependent requests: two or more simultaneous and directly related post-marketing authorization change requests;

XIV – production: all operations involved in the preparation of a food or packaging, from the receipt of materials, through processing and packaging;

XV – food product: any food derived from food raw material or natural food, with or without the addition of other permitted substances, and which has been obtained through an appropriate technological process;

XVI – manufacturer's representative: a company that is not considered the manufacturer, but has an agreement with the manufacturer for representation before Anvisa; and

XVII – food services: institutional or commercial establishments where the food may or may not be consumed on site, but where the food is handled, prepared, stored, distributed, or displayed for sale, such as restaurants, snack bars, bars, bakeries, food and nutrition units of health services, schools and daycare centers, among others.

CHAPTER II

GENERAL REQUIREMENTS FOR THE REGULARIZATION OF FOOD AND PACKAGING

Article 3. The regularization of food and packaging must be carried out prior to its supply through the following procedures:

I – marketing authorization by Anvisa, in the case of the categories listed in Annex I of Normative Instruction – IN No. 281 of 22 February 2024;

II – notification to Anvisa, in the case of the categories listed in Annex II of Normative Instruction – IN No. 281 of 22 February 2024; or

III – communication of the start of manufacture or import to the health authority of the State, Federal District, or Municipality, in the case of the categories listed in Annex III of Normative Instruction – IN No. 281 of 22 February 2024.

Sole paragraph. The categories described in Annex IV of Normative Instruction – IN No. 281 of 22 February 2024, which are exempt from regularization with the SNVS, are exempt from the provisions in the caption of this article.

Article 4. The regularization referred to in Article 3 of this Resolution must be carried out:

I – by the manufacturer's head office, the manufacturer's representative or the importer, in the case of the categories referred to in items I and II of Article 3 of this Resolution; or

II – by the manufacturer or importer, in the case of the categories referred to in item III of Article 3 of this Resolution.

Paragraph 1. In cases of imports on behalf of a third party or by order, the importer referred to in items I and II must be the purchaser or the orderer, respectively.

Paragraph 2. The company that requests regularization shall be its holder.

Article 5. The regularization holder is responsible for:

I – ensuring the veracity, accuracy, and updating of the information presented in the regularization procedures;

II – being responsible for ensuring the health requirements of composition, quality, safety, and labeling established for the regularized product;

III – requesting changes to the information provided in the regularization process whenever the product undergoes modification on its own initiative, on the manufacturer's initiative, due to an update of the legislation, or by determination of the health authority;

IV – requesting revalidation of marketing authorization or maintenance of the notification, whenever there is an interest in continuing to make the product available on the market; and

V – requesting cancellation of the regularization, when there is no longer any interest in offering the product.

Article 6. In the case of the categories referred to in items I and II of Article 3 of this Resolution, the regularization holder must:

I – register in Anvisa’s system and keep the related information updated; and

II – inform, at the time of the regularization request, all manufacturers of the product that perform production, quality control, and storage activities, indicating their respective responsibilities.

Paragraph 1. All manufacturers that perform production, quality control, and storage activities of the products referred to in the caption of this article must be registered in Anvisa's system and are responsible for keeping the declared information up to date.

Paragraph 2. The registration of Brazilian companies must be carried out by the company itself, in accordance with the procedure manual available on Anvisa's website.

Paragraph 3. The registration of international companies must be carried out by Anvisa based on the information presented at the time of the regularization request.

Paragraph 4. Updates to the registrations of Brazilian and international companies must be requested through a specific petition protocol.

Article 7. In the case of the categories referred to in items I and II of Article 3 of this Resolution, different presentations of the same product may be requested in a single regularization request when they differ by at least one of the following variations:

I – types or concentration of food additives used in the technological functions of coloring or flavoring;

II – types or concentration of other ingredients used to provide color, flavor, or aroma to the product; or

III – types of or packaging materials.

Article 8. In case the categories referred to in items I and II of Article 3 of this Resolution have different manufacturers or different brands, such information must be presented in a single regularization request, not configuring, in these cases, different presentations.

Article 9. The documents proving that the product meets the requirements of the specific applicable health standards must be made available to the health authority, when requested.

Article 10. After regularization of the products of the categories referred to in items II and III of Article 3 of this Resolution, the competent health authority may indicate the need to perform control analysis.

CHAPTER III

SPECIFIC REQUIREMENTS FOR MARKETING AUTHORIZATION AND RENEWAL, CANCELLATION AND POST-MARKETING AUTHORIZATION CHANGES

Section I

Marketing authorization

Article 11. The marketing authorization request must be filed with Anvisa by means of a petition with a specific subject code for the category of the product in question.

Paragraph 1. The petition referred to in the caption of this article must be accompanied by the documents described in Annexes V and VI of Normative Instruction – IN No. 281 of 22 February 2024.

Paragraph 2. If the product object of marketing authorization has more than one presentation, the petition referred to in the caption of this article must be accompanied by information on all presentations.

Article 12. The marketing authorization of the product shall be granted by means of publication in the Federal Official Gazette (DOU, in Portuguese).

Paragraph 1. The product may only be offered on the market after publication of the marketing authorization in the DOU.

Paragraph 2. The labeling of the food granted marketing authorization must contain the information "Food granted marketing authorization by Anvisa:", followed by the full registration number published in the DOU, observing the requirements of Article 8 of Collegiate Board Resolution – RDC no. 727 of 1 July 2022.

Paragraph 3. The marketing authorization shall be valid for 5 (five) years from its date of publication in the DOU.

Section II

Marketing authorization revalidation

Article 13. The request for marketing authorization revalidation must be filed:

I – by means of a petition with a specific subject code for the category of the product in question; and

II – at most twelve months and at least three months before the expiration of the marketing authorization.

Paragraph 1. The petition referred to in item I of this article must be accompanied by the documents described in Annex VII of Normative Instruction – IN No. 281 of 22 February 2024.

Paragraph 2. Filing the petition after the deadline provided for in item II of this article implies its rejection.

Article 14. Anvisa shall grant automatic revalidation when the marketing authorization revalidation petition:

I – is filed in a timely manner, in accordance with item II of Article 13 of this Resolution; and

II – is pending a decision on the marketing authorization expiration date.

Sole paragraph. Automatic revalidation does not prevent Anvisa from continuing to analyze the petition.

Article 15. Marketing authorization revalidation and automatic revalidation of product marketing authorization shall be granted by means of publication in the DOU.

Sole paragraph. The publication of the revalidations referred to in the caption of this article implies maintaining the initial marketing authorization number of the product.

Section III

Cancellation of marketing authorization

Article 16. The cancellation of the marketing authorization of a product shall be carried out when:

I – the holder requests its cancellation, in accordance with item V of Article 5 of this Resolution;

II – the petition for revalidation of marketing authorization is rejected by Anvisa; or

III – the holder does not request or requests revalidation of marketing authorization late, in accordance with Article 13 of this Resolution.

Paragraph 1. The cancellation of the marketing authorization shall be carried out by means of its publication in the DOU.

Paragraph 2. In the case provided for in item II of this article, the marketing authorization cancellation shall be published only after its expiration and the maintenance of rejection decision by the administrative appeals instances.

Paragraph 3. In the cases provided for in item III of this article, the marketing authorization cancellation shall be published only after its expiration due to expiry.

Paragraph 4. The cancellation referred to in the caption of this article shall be carried out without prejudice to other actions or measures provided for in health legislation.

Section IV

Post-marketing authorization changes

Article 17. The request for a post-marketing authorization change must be filed by means of a petition with a specific subject code for each type of change.

Sole paragraph. The types of post-marketing authorization change, their purpose and the documents that must accompany the petitions referred to in the caption of this article are described in Annexes VIII and IX of Normative Instruction – IN No. 281, of 22 February 2024.

Article 18. Interdependent post-marketing authorization change petitions must be filed and analyzed jointly.

Sole paragraph. The rejection of one of the petitions referred to in the caption of this article implies the rejection of the others.

Article 19. The conditions for implementing each type of post-marketing authorization change are defined in Annex VIII of Normative Instruction – IN No. 281 of 22 February 2024, and may be carried out:

I – without prior authorization from Anvisa;

II – after the established analysis period; or

III – after analysis and favorable decision by Anvisa.

Paragraph 1. The changes referred to in item I of this article must be implemented immediately after the petition is filed with Anvisa.

Paragraph 2. The changes referred to in item II of this article may be implemented after the established period has elapsed, provided that there is no manifestation from Anvisa.

Paragraph 3. In the case of post-marketing authorization changes implemented based on item II of this article whose petitions are subsequently analyzed and rejected by Anvisa, the conditions prior to the filing of the petition must be reestablished.

Paragraph 4. In the case of post-marketing authorization changes analyzed and approved by Anvisa:

I – the deadline for implementation is up to 180 (one hundred and eighty) days, after the publication of the decision in the DOU; and

II – the production of batches under different conditions is not permitted, after the production of the first batch of the product with the post-marketing authorization changes.

Paragraph 5. Interdependent post-marketing authorization changes must be implemented in full and in a single act.

Article 20. Unless otherwise provided, in the case of updating the legislation that implies post-marketing authorization changes, the holder must file the respective petitions in a timely manner, in order to ensure that all necessary adjustments to the product are implemented within the adjustment period provided, in full and in a single act, considering the provisions of articles 17 to 19 of this Resolution.

CHAPTER IV

SPECIFIC REQUIREMENTS FOR NOTIFICATION AND EVALUATION, MAINTENANCE, CANCELLATION, AND CHANGES TO NOTIFICATION

Section I

Notification

Article 21. The notification must be filed with Anvisa by means of a petition with a specific subject code for the category of the product in question.

Paragraph 1. The petition referred to in the caption of this article must be accompanied by the documents described in Annex X of Normative Instruction – IN No. 281 of 22 February 2024.

Paragraph 2. If the notified product has more than one presentation, the petition must be accompanied by information about all presentations.

Paragraph 3. The notification that meets the requirements of this Resolution shall be made automatically, not preceded by analysis by Anvisa.

Paragraph 4. The product may only be offered on the market after the notification has been filed.

Paragraph 5. The labeling of the notified food must contain the information "Food notified with Anvisa:", followed by the full number of the notification process.

Section II

Notification analysis

Article 22. The notification of the product may be subject to analysis by Anvisa at any time.

Paragraph 1. Anvisa may request additional information to that recorded in the notification, in order to support the evaluation of the adequacy of the notified product to the legislation.

Paragraph 2. When inaccuracies are found in the notified information, Anvisa may:

I – inform the holder on the corrections necessary in the notification of the product and the respective deadline for adaptation;

II – proceed with the corrections and inform the holder on the adaptations to be implemented in the product and the respective deadline for adaptation; or

III – cancel the notification of the product and inform the holder on the measures to be adopted, in the case of inaccuracies that represent a risk to consumer's health.

Paragraph 3. The corrective measures referred to in Paragraph 2 of this article shall be carried out without prejudice to other actions or measures provided for in health legislation.

Section III

Maintenance of notification

Article 23. The request for maintenance of notification must be filed:

I – by means of a single petition declaring interest in the continued marketing of all products notified by the holder; and

II – between the first day of October and the last day of December of each year defined for the expression of interest.

Paragraph 1. The years for the expression of interest are defined by successively adding 5 (five) years to the year this Resolution enters into force.

Paragraph 2. If the request for maintenance of notification is not filed in accordance with the provisions in the caption of this article, the notification shall be inactivated.

Paragraph 3. After the inactivation of the notification, the holder may file, at any time, the request for reactivation by means of an individual petition for each product.

Section IV

Cancellation of notification

Article 24. The cancellation of a product's notification will be carried out when:

I – the holder requests its cancellation, in accordance with item V of Article 5 of this Resolution; or

II – inaccuracies that pose a risk to the population's health are identified, in accordance with item III of Paragraph 2 of Article 22 of this Resolution.

Sole paragraph. The cancellation referred to in the caption of this article shall be carried out without prejudice to other actions or measures provided for in health legislation.

Section V

Amendment to notification

Article 25. The request for amendment to notification must be filed by means of a petition for amendment of notification.

Paragraph 1. The petition referred to in the caption of this article must be accompanied by the documents described in Annex X of Normative Instruction – IN No. 281 of 22 February 2024, which were impacted by the proposed amendment.

Paragraph 2. The offer of the changed product may only be initiated after the notification amendment petition has been filed.

Paragraph 3. The request referred to in the caption of this article may not change the product category.

CHAPTER V

COMMUNICATIONS OF START OF MANUFACTURE OR IMPORT AND CHANGE AND CANCELLATION OF THE COMMUNICATIONS

Section I

Communications of start of manufacture or import

Article 26. The notification of start of manufacture or import must be made to the competent Health Surveillance body, by means of the form contained in Annex XI of Normative Instruction – IN No. 281 of 22 February 2024, duly completed.

Paragraph 1. When the same product has different manufacturers or importers, each manufacturer or importer must make the protocol referred to in the caption of this article with the respective competent health authority.

Paragraph 2. The product may be made available on the market after the notification of start of manufacture or import has been filed.

Paragraph 3. The notification of the start of manufacture or import of the product does not make it approved by the health authority.

Paragraph 4. The communication of start of manufacture or import is valid indefinitely.

Article 27. After receiving the communication of start of manufacture or import, the competent health authority may, at its discretion, carry out a health inspection in the manufacturing or storage units of the food or packaging.

Section II

Amendment to the communication of start of manufacture or import

Article 28. When there is a change in the information provided in the communication of start of manufacture or import, a request to amend the communication must be made by filing a new form contained in Annex XI of Normative Instruction – IN No. 281 of 22 February 2024, duly completed.

Paragraph 1. The request referred to in the caption of this article results in the automatic cancellation of the initial communication.

Paragraph 2. The availability of the altered product on the market may begin after filing the form referred to in the caption of this article.

Section III

Cancellation of the communication of start of manufacture or import

Article 29. The communication of start of manufacture or import of a product shall be cancelled when the holder requests its cancellation, in accordance with item V of Article 5 of this Resolution.

Sole paragraph. The cancellation referred to in the caption of this article shall be carried out without prejudice to other actions or measures provided for in health legislation.

CHAPTER VI

FINAL AND TRANSITIONAL PROVISIONS

Article 30. The deadline is set until 1 September 2026 to request adaptation of products classified in the categories listed in Annex I of Normative Instruction – IN No. 281 of 22 February 2024, which have already been granted marketing authorization on the date this Resolution comes into effect.

Paragraph 1. The products referred to in the caption of this article whose marketing authorization expires by 1 September 2026 may request their revalidation within 60 (sixty) days before the expiration date.

Paragraph 2. The products referred to in the caption of this article that are manufactured up to the date of publication of the final decision on the request for adaptation may be made available on the market until the end of their validity periods.

Paragraph 3. The adaptation of the products referred to in the caption of this article must be carried out by means of a specific post-marketing authorization petition protocol.

Paragraph 4. Failure to comply with the deadline established in the caption of this article for the request for adaptation of the product shall result in the cancellation of the marketing authorization.

Article 31. The deadline is set until 1 September 2025 to apply for marketing authorization of diet therapy formulas for inborn errors of metabolism that have been the subject of a communication of start of manufacture or import to the competent health authority up to the date this Resolution enters into force.

Sole paragraph. The products referred to in the caption of this article that are manufactured up to the date of publication of the final decision on the application for marketing authorization may be made available on the market until the end of their validity periods.

Article 32. The deadline is set until 1 September 2025 for the notification of weight control foods and dietary supplements that have been the subject of a communication of start of manufacture or import to the competent health authority up to the date this Resolution enters into force.

Sole paragraph. The products referred to in the caption of this article that are manufactured up to the date of notification may be made available on the market until the end of their validity periods.

Article 33. Notification of products included in the food and packaging categories listed below that have been granted marketing authorization up to the effective date of this Resolution must be filed by the expiration of their marketing authorization:

I – desalinated, potable, and bottled seawater;

II – foods with functional or health claims;

III – transitional foods for infant feeding;

IV – cereals for infant feeding;

V – resin, precursor article, or final packaging of food-grade PET-PCR; and

VI – food supplements containing probiotics or enzymes.

Paragraph 1. The notification referred to in the caption of this article must be filed simultaneously with the request for cancellation of the marketing authorization.

Paragraph 2. Requests for marketing authorization and post-marketing authorization changes for the products listed in the caption of this article that are pending a decision shall be closed.

Paragraph 3. The products referred to in the caption of this article that are manufactured during the validity of the marketing authorization may be made available on the market until the end of their validity periods.

Paragraph 4. The products referred to in the caption of this article shall not be subject to post-marketing authorization changes.

Paragraph 5. The marketing authorizations of the products referred to in the caption of this article shall not be revalidated after the date this Resolution enters into force.

Article 34. Item 4.3 of the Annex to Ordinance SVS/MS No. 27 of 18 March 1996 shall come into force with the following wording:

"4.3. Ceramic, glass, or metal packaging and equipment that are enameled or vitrified on the side in contact with food must comply with the limits specified in points 5.1.7 and 5.2.4 of this technical regulation." (new wording)

Article 35. Resolution – RES No. 18 of 30 April 1999 shall come into force with the addition of the following items 3.6, 3.7, and 3.8:

"3.6 Scientific proof of the claim of functional and/or health properties referred to in item 3.4 of this Resolution must be requested prior to its use, by means of a specific petition protocol, containing the information required in item 4 of this Resolution. Additional information may be requested when necessary and justified by Anvisa.

3.7 The decision on the petition referred to in item 3.6 of this Resolution shall be published in the Federal Official Gazette, by means of a Specific Resolution (RE, in Portuguese).

3.8 Anvisa's favorable opinion on the petition referred to in item 3.6 of this Resolution does not exempt foods with claims of functional and/or health properties from meeting the other requirements necessary for their regularization." (new wording)

Article 36. Items 3.3, 3.5, 3.6, 3.7, 3.8, and 3.9 of the Annex to Collegiate Board Resolution – RDC No. 20 of 26 March 2008 shall come into force with the following wording:

"3.3. Food-grade PET-PCR precursor articles and packaging must be regularized with the Competent Health Authority, following the procedures established in Collegiate Board Resolution – RDC No. 843 of 22 February 2024, and Normative Instruction – IN No. 281 of 22 February 2024, and must declare whether they are single-use or returnable multilayer or monolayer precursor articles or packaging containing food-grade PET-PCR, as applicable.

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3.5. Food producers may only use PET-PCR precursor articles or packaging food-grade PET-PCR that are duly regularized by the Competent Health Authority, observing the specified foods and the conditions stipulated in the respective regularization, based on the special authorizations for use defined in item 2.6.

3.6. Establishments producing packaging or precursor articles of food-grade PET-PCR must be authorized by the Competent Health Authority and must request the regularization of these packaging or their precursor articles, following the established procedures.

3.7. For an establishment that produces packaging or their precursor articles of food-grade PET-PCR to be authorized, the following must be available:

- Written procedures and their application records on Good Manufacturing Practices for consultation by the Competent Health Authority;

- Records of origin and composition or characterization of food-grade PET-PCR and virgin PET, with documentation confirming this;

- Equipment suitable for packaging and processing food-grade PET-PCR;

- Process control procedures to produce packaging or their precursor articles of food-grade PET-PCR that allow their traceability;

- Personnel for the operation of all equipment and for process control specifically trained for this purpose; and

- A quality assurance system that prevents contamination with other sources of recycled material for applications other than food grade.

3.8. Establishments authorized to produce packaging or precursor articles of food grade PET-PCR must use for this purpose, in addition to virgin PET resin, only food grade PET-PCR obtained through a physical or chemical recycling technology regularized with the Competent Health Authority and analyzed by its recognized Reference Laboratory.

3.9. Establishments authorized to produce food-grade PET-PCR packaging or precursor articles must obtain food-grade PET-PCR from a producer authorized by the Competent Health Authority and use it to manufacture packaging or precursor articles intended to contain only the specified foods and only under the conditions stipulated in the regularization with the Competent Health Authority, based on the special authorizations for use defined in item 2.6" (new wording)

Article 37. Article 21 of Collegiate Board Resolution – RDC No. 243 of 26 July 2018, shall come into force with the following wording:

"Article 21. The regularization of dietary supplements, as provided for in Collegiate Board Resolution – RDC No. 843 of 22 February 2024, and Normative Instruction – IN No. 281 of 22 February 2024, shall be linked to the prior decision on the petition for safety and efficacy assessment, in accordance with the procedure provided for in Article 20 of this Resolution." (new wording)

Article 38. Articles 5 and 6 of Collegiate Board Resolution – RDC No. 241 of 26 July 2018, shall come into force with the following wording:

"Article 5. Proof of safety and health benefits of probiotics does not exempt foods containing probiotics from meeting the other requirements necessary for their regularization established in Collegiate Board Resolution – RDC No. 843 of 22 February 2024, and Normative Instruction – IN No. 281 of 22 February 2024.

Article 6. The regularization of foods containing probiotics shall be linked to the prior decision on the petition for safety and efficacy assessment, in accordance with the procedure set forth in Article 4 of this Resolution." (new wording)

Article 39. Failure to comply with this Resolution constitutes a health violation, subjecting violators to the penalties of Law No. 6,437 of 20 August 1977, and other applicable provisions.

Article 40. The following provisions are hereby revoked:

I – Resolution – RES No. 19 of 30 April 1999;

II – Collegiate Board Resolution – RDC No. 22 of 15 March 2000;

III – Collegiate Board Resolution – RDC No. 23 of 15 March 2000;

IV – Collegiate Board Resolution – RDC No. 27 of 6 August 2010;

V – Collegiate Board Resolution – RDC No. 240 of 26 July 2018;

VI – item 10 of the Annex to Ordinance SVS/MS No. 34 of 13 January 1998;

VII – item 10 of the Annex to Ordinance SVS/MS No. 36 of 13 January 1998;

VIII – items 10, 11, and 12 of the Annex to Resolution – RES No. 105 of 19 May 1999;

IX – the sole paragraph of Article 11 of Collegiate Board Resolution – RDC No. 243 of 26 July 2018;

X – Article 23 and Annex I of Collegiate Board Resolution – RDC No. 460 of 21 December 2020; and

XI – Article 12 and Annex III of Collegiate Board Resolution – RDC No. 818 of 28 September 2023.

Article 41. This Resolution enters into force on 1 September 2024.

ANTONIO BARRA TORRES

Director-President