



Republic of the Philippines

PHILIPPINE HEALTH INSURANCE CORPORATION

Citystate Centre, 709 Shaw Boulevard, Pasig City

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PhilHealthOfficial X teamphilhealth

TO

HEALTH CARE PROVIDERS, HEALTH CARE PROVIDER NETWORKS, LOCAL GOVERNMENT UNITS, PHILHEALTH REGIONAL OFFICES, LOCAL HEALTH INSURANCE OFFICES, AND ALL OTHERS CONCERNED

SUBJECT

: Supplemental Guidelines for the Health Care Provider

Network (HCPN) Demonstration Program

I. RATIONALE

Republic Act (RA) No. 11223 or the Universal Health Care Act (UHC Act) introduces systems-level reforms that aim to transform the delivery of healthcare in the Philippines. These reforms cover different aspects of the healthcare system, and work to reinforce each other and create an ecosystem that guarantees comprehensive, quality, efficient, and affordable health services.

MASTER COPY C. The Date: 10/16/24 In line with the vision of the UHC Act, PhilHealth has developed innovative healthcare financing policies. Given the complexity and novelty of these reforms, PhilHealth will initially implement the new policy approaches and interventions within integrated sandbox sites. Health Care Provider Network (HCPN) Demonstration Program, a joint initiative with the Department of Health (DOH), identifies sandbox sites that will serve as the testing grounds for the health financing reforms. PhilHealth Circular (PC) No. 2023-0016 (Guiding Principles for Integrated UHC Benefits and Provider Payment Reforms Sandbox Sites) outlines the guiding principles for conducting sandboxes for health financing policies.

To support these reforms, prospective payment and contracting mechanisms shall be employed. PC No. 2024-0011 (PhilHealth Prospective Payment Mechanisms for Health Care Provider Networks) and PC No. 2024-0015 (Guidelines on Contracting Health Care Provider Networks) provide the guidelines for implementing prospective payment mechanisms in HCPN, and contracting HCPN, respectively. By virtue of PhilHealth Board Resolution No. 2932, s. 2024, the set of benefits to be implemented for Calendar Year (CY) 2024 and 2025 has been approved.





II. OBJECTIVES

This PhilHealth Circular aims to supplement the policies for implementation in HCPN demonstration sites.

III. SCOPE

This PhilHealth Circular encompasses the engagement of providers and provider networks, and the benefit packages to be delivered within the HCPN Demonstration Program.

IV. DEFINITION OF TERMS

A. Health Care Provider Network Demonstration Sites (HCPN Demonstration Sites) - possible areas for implementation of healthcare financing reforms of the UHC Act as identified and recognized by the Department of Health (DOH) through the Department Circular (DC) No. 2024-0237 (List of Universal Health Care Integration Sites Qualified for Health Care Provider Network Contracting), as amended by DC No. 2024-0344 (Updated List of Universal Health Care Integrated Sites Qualified for Health Care Provider Network Contracting based on the F.Y. 2024 Local Health Systems Maturity Level Mid-Year Report), and its succeeding revisions/amendments.

V. POLICY STATEMENTS

- A. General Provisions for the Health Care Provider Network (HCPN)
 - 1. To support the implementation of integrated sandbox, also known as the HCPN Demonstration Program, contracting networks shall be allowed to commence during the remaining period of CY 2024. The contract period shall end on December 31, 2024.
 - 2. The HCPN demonstration for CY 2024 shall be facilitated by the following changes:
 - a. The HCPN demonstration sites shall continue to provide primary care services through the enhanced PhilHealth Konsulta Benefit Package, stipulated in PC No. 2024-0013 (Enhancement of the PhilHealth Konsulta Benefit Package). PhilHealth Konsulta Providers shall maintain their current engagement with PhilHealth.
 - b. In addition to PhilHealth Konsulta, Networks in HCPN demonstration sites shall be contracted and paid for the following:
 - b.1 Inpatient services through the All Case Rates-based Global Budget (ACR-GB) as stipulated in PC No. 2024-0004; and



- b.2 Outpatient drug benefit, also known as PhilHealth Guaranteed Access to Medications for Outpatient Treatment (PhilHealth GAMOT) as stipulated in PC No. 2023-0029.
- c. Benefit packages and services not covered under ACR-GB and PhilHealth GAMOT, including but not limited to outpatient benefits (medical services and procedures provided and performed in less than 24 hours) and Z benefits, shall be paid directly to the member facilities of the Network following the existing rules on payment and engagement.
- 3. Networks in HCPN demonstration sites shall be contracted in CY 2025 for the following:
 - a. Inpatient services through the ACR-GB;
 - b. Outpatient Drug Benefit; and
 - c. K+SDG Benefit (PhilHealth Konsulta, Animal Bite Treatment, Outpatient HIV-AIDS Treatment, Outpatient Anti-Tuberculosis Directly Observed Treatment Short Course [DOTS] Benefit, and Outpatient Malaria Treatment Package).

B. Engagement of HCPN

- 1. A separate contracting process shall be conducted for CY 2024 and CY 2025, following the guidelines in PC No. 2024-0015.
- 2. Any extension to the application period shall be published through a PhilHealth Advisory. This extension includes the provisions in the PhilHealth Advisory No. 2024-0042 (Extension of Deadline for Contracting Health Care Provider Networks for CY 2024).
- 3. The template memorandum of agreement (MOA) and service-level agreement (SLA) shall be revised to include only the provisions for ACR-GB and PhilHealth GAMOT for CY 2024 (Annex A.1 and A.2 for Template MOA and SLA Between PhilHealth and Network for CY 2024, respectively). Annexes A.1 and A.2 repeal the Annexes B.1 and B.2 of PC No. 2024-0015.
- 4. The template MOA and SLA shall be updated to reflect the benefits to be implemented during the succeeding years of HCPN demonstration.
- 5. The PhilHealth President and Chief Executive Officer, as authorized by the PhilHealth Board of Directors, shall provide clearance for each MOA and SLA with a Network for HCPN Demonstration Program.



- 6. The first tranche of payment to the Network shall be released within the first 15 calendar days of the contract period. The payment releases to the member health facilities shall be agreed upon with the Network, and shall be reflected on the respective SLA between the Network and member facilities.
- 7. The implementation of this policy shall be subject to the accounting guidelines on prospective payment mechanisms of Commission on Audit, and the functionality of benefit's information systems.
- 8. Contingent to the readiness of the information technology (IT) system to operationalize the benefit, progressive implementation of benefits shall be allowed. The relevant addenda to the SLA shall be executed to reflect any additional benefits.
- 9. Rolling implementation of the HCPN Demonstration Program shall be allowed. Operational details, including but not limited to the contracting period cycle, shall be issued through a PhilHealth Advisory.
- 10. The MOA with an Apex hospital shall no longer be required as a primary prerequisite for CY 2025 Network Contracting.
- 11. The validity of the MOA and SLA between PhilHealth and the Network shall coincide with the outgoing local chief executive's (LCE) term of office on June 30, 2025. A new agreement shall be executed for the incoming LCE beginning July 1, 2025 until December 31, 2025. Reconciliation, in accordance with each benefit guidelines, shall be conducted to determine unutilized funds, if any.
 - a. In case of change in leadership, the newly elected LCE shall enter into a new agreement with PhilHealth.
 - a.1 This new agreement with PhilHealth may be done through execution of new MOA and SLA following the existing contracting process, or through a renewal of MOA, subject to execution of a new SLA.
 - a.2 In the event that the newly elected LCE opts not to continue its Network's previous engagement with PhilHealth, any unutilized fund shall be subjected to PhilHealth Payment Recovery (PPR) Policy.
 - b. If there is no change in leadership, the incumbent LCE may opt to request for renewal of the existing MOA with PhilHealth through a formal written communication, subject to amendment of the existing SLA. Any unutilized fund from the previous agreement shall form part of the succeeding contract amount and shall be stipulated in the new or amended SLA.



c. Further guidelines regarding the above mentioned scenarios shall be issued through a PhilHealth Advisory.

C. Benefits

- 1. PhilHealth Guaranteed Access to Medications for Outpatient Treatment (PhilHealth GAMOT)
 - a. The CY 2024 implementation of PhilHealth GAMOT shall be facilitated by the following modifications:
 - a.1 The 21 medicines and corresponding formulations provided by the enhanced PhilHealth Konsulta shall be removed from the GAMOT Basic Medication List (BML) and GAMOT Application. The beneficiaries shall claim these 21 medicines, as outlined in PC No. 2024-0013, from their respective Konsulta Providers (Annex B.1: List of Medicines Not Included Under PhilHealth GAMOT for CY 2024).
 - a.2 PhilHealth Konsulta Physicians shall be granted access to the GAMOT Application to prescribe the remaining 32 medicines and corresponding formulations covered under PhilHealth GAMOT (Annex B.2: List of GAMOT Basic Medication List for CY 2024, Annex B.3: Summary of Basic Medication List Molecules for CY 2024).
 - a.3 The 32 medicines and corresponding formulations covered under PhilHealth GAMOT shall be included in the computation of the contract amount.
 - a.4 The contract amount shall be calculated to cover only the average sales corresponding to the number of months in the engagement (e.g., three [3] months of October December) (Annex D: GAMOT Contract Amount for CY 2024 Sample Computation).
 - a.5 The full contract amount shall be paid to the Network within the first 15 calendar days of the execution of the contract and the necessary operational adjustments shall be done to facilitate such release of the full contract amount as the sole tranche.
 - b. The GAMOT BML shall be updated to cover all formulations stipulated in PC No. 2024-0013, including the thirteen (13) drug formulations that are not within the Primary Care Formulary (Annex C: Additional 13 Formulations for PhilHealth GAMOT for CY 2025). A PhilHealth Advisory shall be issued to reflect the updated GAMOT BML and the corresponding fixed fee schedule.



- c. Free-standing facilities (e.g., retail drug stores/pharmacies, FOURmula One Plus Botika ng Bayan [BNB] outlets of the DOH) and infirmaries with licensed pharmacies having less than three (3) years but at least one (1) year of operations shall be exempted from 3-year operations requirement and may apply as GAMOT providers. The contract amount shall be estimated based on the annual sales for those facilities with 1 year of operations, and average annual sales for those facilities with 2 years of operations.
- d. The GAMOT contract amount shall be computed by multiplying all the medication's fixed fee schedule with its average volume of sales for the most recent three years, as applicable. Annex D: GAMOT Contract Amount for CY 2024 Sample Computation shall repeal the Annex H of PC 2023-0029.
- e. The prospective payment to the Network for GAMOT shall be released in tranches and shall be guided by the following rules:
 - e.1 The first tranche shall be 60% of the contract amount and shall be released within the first 15 calendar days of the contract period.
 - e.2. The second tranche shall be 30% of the contract amount and can be requested upon 60% consumption of the first tranche.
 - e.3. The third and last tranche shall be 10% of the contract amount and can be requested upon 80% consumption of the first and second tranches.
- f. Accredited health facilities that are applying to be GAMOT providers as an ancillary or additional service (e.g., hospital pharmacy) shall not be charged an accreditation fee. In case of payment already made, the paid accreditation fee for GAMOT shall be deducted from the accreditation fee for the next application.
- g. Considering the short time of implementation for CY 2024, the accreditation fee of applicant free-standing GAMOT providers and infirmaries with licensed pharmacies for CY 2024 shall be waived and any payment made earlier shall be considered as advanced payment for CY 2025 accreditation.
- 2. All Case Rates-Based Global Budget (ACR-GB)
 - a. The CY 2024 implementation of ACR-GB shall be facilitated by the following modifications:
 - a.1 The contract amount shall be estimated based on the applicable months in the SLA (e.g., October, November, and December) of the immediately preceding three (3)



- complete years excluding CY 2020 and 2021 (i.e., 2019, 2022, 2023).
- a.2 Level 1-3 hospitals as ACR-GB facilities shall have a soft budget cap and shall be eligible to request for supplementary budget (10% of the contract amount), if the GB is fully utilized during the contract period or the occurrence of fortuitous events with official declaration (i.e., state of calamity, state of emergency, or any analogous certification) during the contract period.
- a.3 The full contract amount shall be paid to the Network within the first 15 calendar days of the execution of the contract and the necessary operational adjustments shall be done to facilitate such release of the full contract amount as the sole tranche.
- b. The ACR-GB shall be calculated following the previous three-year average, as stipulated in PC No. 2024-0004 (Implementation of an All Case Rates-Based Global Budget in Health Care Provider Network Demonstration Sites). Paid claims admitted during the preceding three (3) complete years, excluding CY 2020 and CY 2021, shall be used for computing the contract amount.
 - b.1 With the implementation of adjusted case rates as stipulated in PC No. 2024-0012 (Rules for Adjusting Case Rates [Revision 1]), the necessary modifications on the contract amount computation (Annex E: ACR-GB Contract Amount Sample Computation) shall be carried out.
 - b.1.1 To determine the claims amount of each case, the count (volume) of paid claims per case (including primary and secondary cases) shall be multiplied to the corresponding current case rate; and
 - b.1.2 To determine the contract amount, the claims amount of all applicable cases shall be summed and shall be divided by three (3).
- c. In the event that the case rates are adjusted during the contract period, the contract amount shall be re-computed to accommodate the new rates from the date of effectivity of case rates adjustment until the end of the contract period (Annex F: ACR-GB Case Rates Adjustment Sample Computation). Supplemental SLA shall be executed to formalize these adjustments (Annex G: ACR-GB Case Rates Adjustment Frontloading Schedule).



- d. Existing rules and procedures on claims filing shall apply. The utilization of ACR-GB shall be the aggregate of good claims for patients admitted within the contract period, and filed or refiled within the contract period and up to 60 days from its termination; provided that the claims are filed or refiled within 60 days from the date of discharge and return-to-hospital (RTH), respectively.
 - d.1 When assessed as good claims, claims for patients admitted within the contract period which are filed within the contract period and up to 60 days thereafter shall be deductible from the ACR-GB.
 - d.2 When assessed as good claims, those claims for patients admitted within the contract period but filed beyond the 60-day period from the expiration of the contract shall be reimbursed based on existing rules on the All Case Rates, and not deductible from the ACR-GB.
 - d.3 When assessed as good claims, RTH claims for patients admitted within the contract period that are refiled within the contract period and up to 60 days thereafter shall be deductible from the ACR-GB.
 - d.4 RTH claims for patients admitted within the contract period that are not refiled within the contract period and up to 60 days thereafter shall be subject to reimbursement when assessed as good claims. RTH claims reassessed as denied claims shall not be deductible from the ACR-GB.
 - d.5 Denied claims for patients admitted within the contract period which are reassessed as good claims upon motion for reconsideration or appeal within the contract period and 60 days thereafter shall be deductible from the ACR-GB.
 - d.6 Denied claims for patients admitted within the contract period which are not reassessed as good claims, upon motion for reconsideration or appeal, within the contract period and up to 60 days thereafter, shall be subject to reimbursement based on existing rules on the All Case Rates, and not be deductible from the ACR-GB.
 - d.7 The processing of claims from the ACR-GB facilities shall be prioritized in time for reconciliation.
- e. Any unutilized funds before reconciliation shall be credited to the first tranche payment for the succeeding contract period, provided that the Network maintains its engagement.



Otherwise, the PhilHealth Payment Recovery (PPR) Policy shall apply.

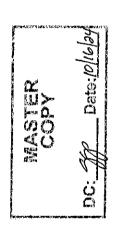
- e.1 Reconciliation for the applicable contract period shall be conducted within three (3) months from the date of its expiration, as stipulated in PC No. 2024-0004.
- e.2 Upon reconciliation, any difference in the amount of unutilized funds shall be credited to the applicable tranche payment of the succeeding contract period and front loaded within 15 days from reconciliation (Annex H: Sample Reconciliation of ACR-GB).
- D. Annexes (posted at the official website of PhilHealth at www.philhealth.gov.ph)
 - Annex A.1: Template MOA Between PhilHealth and Network for CY 2024
 - 2. Annex A.2: Template SLA Between PhilHealth and Network for CY 2024
 - 3. Annex B.1: List of Medicines Not Included Under PhilHealth GAMOT for CY 2024
 - 4. Annex B.2: GAMOT Basic Medication List for CY 2024
 - 5. Annex B.3: Summary of Basic Medication List Molecules for CY 2024
 - 6. Annex C: Additional 13 Formulations for PhilHealth GAMOT for CY 2025
 - 7. Annex D: GAMOT Contract Amount for CY 2024 Sample Computation
 - 8. Annex E: ACR-GB Contract Amount Sample Computation
 - 9. Annex F: ACR-GB Case Rates Adjustment Sample Computation
 - 10. Annex G: ACR-GB Case Rates Adjustment Frontloading Schedule
 - 11. Annex H: Sample Reconciliation of ACR-GB

VI. PENALTY CLAUSE

Any violation of this PhilHealth Circular and all existing related PhilHealth Circulars, shall be dealt with and penalized in accordance with the pertinent provisions of Republic Act No. 7875, as amended by Republic Act Nos. 9241 and 10606 (National Health Insurance Act of 2013) and Republic Act No. 11223 (Universal Health Care Act), and their respective Implementing Rules and Regulations, and other pertinent laws and rules.

VII. SEPARABILITY CLAUSE

If, for any reason, any part of this PhilHealth Circular is declared invalid or unconstitutional, any part or provision not affected thereby shall remain in full force and effect.



VIII. REPEALING CLAUSE

Policy statements of corporate issuances, including but not limited to the following, that are inconsistent with any provisions of this PhilHealth Circular are hereby amended, modified, or repealed accordingly:

- A. PC No. 2023-0012: Omnibus Guidelines on the Accreditation of Health Facilities (HFs) to the National Health Insurance Program
 - 1. Policy Statement (V) C.7
- B. PC No. 2023-0029: Implementing Guidelines For The PhilHealth Outpatient Drug Benefit Package
 - 1. Policy Statements (V) E.2.a.6.1.1 and E.2.a.6.2.2
 - 2. Policy Statements (V) G.3, G.4.c, G.4.c.2, and G.4.d.2
 - 3. Annexes A.1, F.1, F.2, and H
- C. PC No. 2024-0004: Implementation of an All Case Rates-Based Global Budget in Health Care Provider Network (HCPN) Demonstration Sites
 - 1. Policy Statement (V) C.2.a
 - 2. Policy Statements (V) F.6, F.11, F.12, F.13, and F.14
 - 3. Annexes E and G
- D. PC No. 2024-0011: PhilHealth Prospective Payment Mechanisms (3PM) For Health Care Provider Networks
 - 1. Policy Statement (V) B.2.c
 - 2. Transitory Clause (VI) A
- E. PC No. 2024-0015: Guidelines Contracting of Health Care Provider Networks (HCPN)
 - 1. Policy Statement (V) C.2.a.2
 - 2. Transitory Clauses (VII) A.1.b, and A.1.c
 - 3. Transitory Clause (VII) B
 - 4. Annexes B.1 and B.2

IX. DATE OF EFFECTIVITY

This PhilHealth Circular shall be published in a newspaper of general circulation and shall take effect immediately upon publication. Thereafter, this PhilHealth Circular shall be deposited with the Office of the National Administrative Register (ONAR) at the University of the Philippines Law Center.

EMMANUEL R. LEDESMA JR. President and Chief Executive Officer

Date signed: 10/15/24

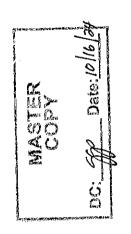
Supplemental Guidelines for the Health Care Provider Network (HCPN) Demonstration Program

Annex A.1: Template MOA Between PhilHealth and Network for CY 2024

MEMORANDUM OF AGREEMENT

KNOW ALL MEN BY THESE PRESENTS:

This Memorandum of Agreement entered into by and between the following parties:



PHILIPPINE HEALTH INSURANCE CORPORATION (PHILHEALTH), a government-owned and controlled corporation duly created by virtue of Republic Act No. 7875 as amended by RA No 9241, RA No. 10606 and RA No. 11223 or the "Universal Health Care Act", as amended, with office address at Citystate Centre, 709 Shaw Boulevard, Oranbo, Pasig City, represented herein by its President and Chief Executive Officer, EMMANUEL R. LEDESMA, JR., hereinafter called "PHILHEALTH"

-and-

[PUBLIC HEALTH CARE PROVIDER NETWORK NAME], an/a (LGU/Province/City-owned network), authorized through a Sanggunian Panlalawigan/Panlungsod Resolution No. ______ issued on [Date] and existing under the laws of the Republic of the Philippines, with principal address at [Address] represented herein by Governor/City Mayor, [Hon. NAME], hereinafter called "NETWORK".

(each a "Party", and collectively the "Parties").

WITNESSETH THAT:

WHEREAS, Republic Act No. 11223 or the Universal Health Care (UHC) Act mandates health reforms that aim to deliver equitable, accessible, and quality health services, and empowers the Philippine Health Insurance Corporation (PHILHEALTH) to implement health financing reforms;

WHEREAS, in response to the need to implement reforms stipulated in UHC Act, **PHILHEALTH** endeavors to contract health care provider networks (HCPN) for the delivery of individual-based health services and to shift to prospective payment mechanisms to finance HCPN;

WHEREAS, the UHC Act mandates integrated local health systems to pool and manage, through a Special Health Fund (SHF), all resources intended for health services to finance population-based and individual-based health services, health system operating costs, capital investments, and remuneration of additional health workers and incentives for all health workers;

WHEREAS, PHILHEALTH has published PhilHealth Circular (PC) No. 2024-0015 outlining the guidelines on contracting HCPN for the delivery of individual-based health services, and PC No. 2024-0011 for implementing the

prospective provider payment for HCPN, which form an integral part of this Agreement;

WHEREAS, **PHILHEALTH** enjoins provinces, highly urbanized cities, independent component cities, and private facilities to integrate their resources and services in a HCPN and to initially implement an **HCPN** in a sandbox, referred to as the HCPN demonstration site; and

WHEREAS, the HCPN, also referred to as NETWORK, is a group of health care providers, hereinafter referred to as MEMBER FACILITIES, governed by a Local Health Board, organized to deliver health care services in a coordinated and integrated manner;

NOW THEREFORE, for and in consideration of the foregoing premises, and of the mutual covenants and stipulations hereinafter set forth, the **Parties** hereby agree, and by these presents, bind themselves, to wit:

I. AGREEMENT

- a. The **NETWORK** agrees to actively participate as an HCPN demonstration site.
- b. The **NETWORK** agrees to maintain compliance with the requirements as prescribed by **PHILHEALTH** to implement benefits and its corresponding payments.
- c. The **NETWORK** shall provide comprehensive and continuous care, from primary to tertiary levels, to PhilHealth beneficiaries through an integrated and coordinated service delivery rendered by its component health care providers and facilities, hereinafter called **MEMBER FACILITIES**.
- d. **PHILHEALTH** and **NETWORK** shall enter into a service-level agreement (SLA) for the delivery of individual health services by the **NETWORK**. The same SLA shall indicate the volume of services and contract amounts to be paid by **PHILHEALTH** to the **NETWORK**. The same SLA specifies the benefits to be implemented by the **NETWORK**.
- e. The **NETWORK** shall enter into SLA with each **MEMBER FACILITY** to formalize engagement between them. The same SLA shall specify the type and volume of services, corresponding contract amounts, and its payment arrangement and schedule.
- f. **PHILHEALTH** agrees to provide the **NETWORK** with regular updates and orientation on **PHILHEALTH** policies and guidelines, address concerns, and issue clarifications as needed to facilitate benefit implementation.
- g. **PHILHEALTH** monitor and evaluate outcomes of the HCPN demonstration to inform future policy and program development.



II. DURATION AND TERMINATION

a.	This Agreement shall	be	effective	upon	signing	of	both	Parties	and	shall
	remain valid from		to)			_			

- b. Either party may terminate this Agreement in the event of a material breach, provided that the breaching party is given an opportunity to rectify the breach within [Number] days of receiving notice.
- c. The Agreement shall remain binding until terminated pursuant to the termination provisions of this Agreement.

III. MISCELLANEOUS PROVISIONS

- a. Both **Parties** agree to maintain the confidentiality of information related to the implementation of this Agreement.
- b. This Agreement shall be governed and construed in accordance with the laws of the Republic of the Philippines. All PhilHealth circulars and issuances shall form an integral part of this Agreement. Venue of all actions arising from this Agreement shall be brought exclusively to the jurisdiction of the appropriate courts of the Philippines, without prejudice to the settlement of dispute through amicable settlement or alternative dispute resolution mechanisms under existing laws.
- c. No amendment or modification of any of the terms and conditions of this Agreement shall be valid unless evidenced by a written agreement executed by the authorized representatives of both **Parties**.
- d. If any part of this Agreement is declared unenforceable or void, the rest of the Agreement shall nevertheless remain in full force and effect.
- e. Both Parties acknowledge that this Agreement constitutes the entries agreement between them and shall completely supersede all other prior understandings, previous communications or contracts, oral or written, between the **Parties** relating to the subject matter hereof.

N WITNESS WHEREOF, the parties have set their hands this day of, 20					
PHILIPPINE HEALTH INSURANCE CORPORATION	[NAME OF THE NETWORK]				
By:	By:				
President & CEO	Authorized Representative of the NETWORK				



PHILHEALTH	NETWORK
ACKNOW	LEDGEMENT
REPUBLIC OF THE PHILIPPINES) S.S.
BEFORE ME, this da following persons, showing to me their respectively.	y of, appeared the respective competent proofs of identity to wit:
NAME ISSUED	GOV'T I.D. NOS. DATE /PLACE
EMMANUEL R. LEDESMA, JR. PCEO, PhilHealth	
Authorized Representative, NETWORK	
and (Name of NETWORK) consacknowledgment page. The Parties ack	who executed the foregoing Memorandum of HEALTH INSURANCE CORPORATION sisting of pages including this nowledged before me that the same is their nat of the corporate entities they respectively
WITNESS MY HAND AND SEAL on	the date and at the place above written.
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"BAWAT PILIPINO, MIYEMBRO BAWAT MIYEMBRO, PROTEKTADO KALUSUGAN NG LAHAT, SEGURADO"

SERVICE-LEVEL AGREEMENT FOR HEALTH CARE PROVIDER NETWORK

This	Agreement for Health Care Provide	ler Network (HCPN) is n	nade and entered into
this_	day of2024,	in Pasig City, Metro Mar	nila, by and between:
	PHILIPPINE HEALTH	INSURANCE	CORPORATION
)ate: 10/16/34	(PHILHEALTH), a governme created by virtue of Republic Act 10606 and RA No. 11223 or the "office address at Citystate Centrepresented herein by its EMMANUEL R. LEDESMA,	ent-owned and controll No. 7875 as amended by Universal Health Care A re, 709 Shaw Boulevard President and Chief JR., hereinafter called "F	led corporation duly RA No 9241, RA No. ct", as amended, with , Oranbo, Pasig City, Executive Officer.
No.		-and-	

[PUBLIC HEALTH CARE PROVIDER NETWORK NAME], an/a (LGU/Province/City-owned network), authorized through a Sanggunian Panlalawigan/Panlungsod Resolution No. ______ issued on [Date] and existing under the laws of the Republic of the Philippines, with principal address at [Address] represented herein by Governor/City Mayor, [Hon. NAME], hereinafter called "NETWORK".

WITNESSETH THAT:

I. GENERAL PROVISIONS

KNOW ALL MEN BY THESE PRESENTS:

- 1. **PHILHEALTH** and the **contracted NETWORK** herein mutually and voluntarily agree to provide eligible PhilHealth members and their dependents primary, secondary, and tertiary care services, including stand-alone facilities that provide specific and/or specialized care.
- 2. All existing PhilHealth Circular/s for Health Care Provider Network and all other applicable PhilHealth Circulars are deemed read into this Agreement. Subsequent issuances pertaining to the subject matter shall also form part of this Agreement.

II. RESPONSIBILITIES OF THE NETWORK

1. For the entire duration of this Agreement, the **NETWORK** shall render the delivery of the following benefits, in accordance with the expected targets, and other terms of delivery.



Services	Benefits	References
Secondary and	All Case Rates-based	PC No. 2024-0004
Tertiary Care	Global Budget	(Subject: Guidelines for the
Services	(ACR-GB)	Implementation of an
		All-Case Rates-Based
		Global Budget in Health
		Care Provider Network
		Demonstration Sites) or
		revision
Specialized Care	Outpatient Drug Benefit	PC No. 2023-0029
Services	(ODB) or PhilHealth	(Subject: Implementing
	Guaranteed and	Guidelines for the
	Accessible Medications	Outpatient Drug Benefit
	for Outpatient	Package) or revision
	Treatment (GAMOT)	

- 2. The **NETWORK** agrees to fully comply with pertinent laws, policies and miscellaneous provisions and fully cooperate during regular surveys, administrative investigations, and domiciliary visitations.
- 3. The **NETWORK** agrees to maintain an information system, including updated registries of beneficiaries and health workers, and to ensure the truthfulness, legitimacy, and accuracy of all data submitted to **PHILHEALTH**.
- 4. The **NETWORK** shall maintain the minimum component health, and shall execute a Memorandum of Agreement with relevant health facilities closest to its geographical jurisdiction for referral of patients when the minimum number of component health facilities are not met due to either of the conditions stipulated in PhilHealth Circular No. 2024-0015 and its succeeding revisions.
- 5. The **NETWORK** guarantees that the following facilities, called "**MEMBER FACILITIES**" are capable of delivering the services expected from them.

Type of Facility	Services	License/ Certificate	PhilHealth Accredited (Y/N)	Head of Facility
:				
-				
	Type of Facility	Type of Facility Services	Type of Facility Services License/Certificate	Facility Services License/ Accredited

DC: MASTER Date: 10|16|24

6. GAMOT

6.1 The **NETWORK**, through its **MEMBER FACILITIES**, dispenses medications under the PhilHealth GAMOT for the first nine thousand pesos (PHP 9,000.00) per individual per calendar year.

6.2 After full utilization of the benefit coverage, the **NETWORK** shall charge to the beneficiaries the medications at prices following the fixed fee schedule or with store prices, whichever is lower or preferred by the beneficiary.

7. ACR-GB

- 7.1 The **NETWORK**, through its **MEMBER FACILITIES**, shall not charge PhilHealth members for health services provided in basic accommodation.
- 7.2 The **NETWORK**, through its **PUBLIC MEMBER FACILITIES**, shall only charge PhilHealth members an amount not exceeding the value of case rate for health services provided in non-basic accommodation.
- 7.3 The **NETWORK** agrees to reach the patient volume of **[number]**, considering the historical patient volume of each hospital:

Name of hospital	Level	Target patient volume
		949
		-
	Total	

7.4 The **NETWORK** may request for an additional 10% of their Global Budget as the supplementary budget for a **MEMBER FACILITY** that fulfills one of the conditions: a) full utilization of the Global Budget within the contract period, b) the occurrence of fortuitous events during the contract period.

III. RESPONSIBILITIES OF PHILHEALTH



- 1. **PHILHEALTH** agrees to provide the **NETWORK** with regular updates and orientation on **PHILHEALTH** policies and guidelines, address concerns, and issue clarifications as needed to facilitate benefit implementation.
- 2. **PHILHEALTH** agrees to assist the **NETWORK** to ensure interoperability and connectivity with **PHILHEALTH** databases to support innovations and electronic medical record initiatives.
- 3. **PHILHEALTH** shall conduct scheduled and/or random on-site or virtual validation visits to observe the actual implementation. **PHILHEALTH** shall provide the results of the monitoring and assessment to the **NETWORK.**

IV. CONDITIONS AND SCHEDULE OF PAYMENT

- The PHILHEALTH only pays the NETWORK once the necessary information system for payment and benefit utilization are already in place.
- 2. The **PHILHEALTH** pays the agreed contract amount of [Amount in words] (Php _______) stipulated in this Contract, including the corresponding schedule of payment and conditions for payment releases stated in applicable PhilHealth Circular/s.

Services	Contract Amount	
a. ACR-GB	[Amount in words] (Php)
b. GAMOT	[Amount in words] (Php	
TOTAL	[Amount in words] (Php)

3. The **NETWORK** shall ensure the timely frontloading of PhilHealth payment amounts to the **MEMBER FACILITIES**, following the payment arrangement and schedule stipulated in the SLA between the **NETWORK** and **MEMBER FACILITIES**, subject to due reconciliation to determine unutilized funds.

A. ACR-GB

Name of the facility	Contract Amount	**
	[Amount in words] (Php)

B. GAMOT

Name of the facility	Contract Amount	
	[Amount in words] (Php	_)

- 4. The **NETWORK** assumes full responsibility of the management of prospective payment released by PhilHealth, subject to pertinent government budgeting, accounting and auditing rules and regulations.
- 5. The **NETWORK** shall submit payment utilization reports, financial reports, and other monitoring reports to **PHILHEALTH** in accordance with the provisions of relevant **PHILHEALTH** Circulars.
- 6. The **NETWORK** agrees to fully comply with mechanisms set by **PHILHEALTH** to recover unutilized funds after due reconciliation.

V. JURISDICTION AND VENUE

The parties hereby consent and agree that jurisdiction and venue for any claim or cause of action arising under or related to this Contract or any order shall be properly and exclusively filed in the applicable courts of Pasig City only, without prejudice to resorting to Alternative Dispute Resolution mechanisms under existing laws.



VI. PENALTIES FOR BREACH OF CONTRACT

- Breach of any of the terms and conditions provided herein shall cause the cancellation of this Agreement and the guilty party shall be liable for damages without prejudice to availing other legal remedies.
- 2. Fabrication of reports shall be penalized in line with the provisions of R.A. 7875 as amended by R.A. 9241, R.A. 10606 and R.A. 11223 and its Implementing Rules and Regulations and/or other applicable laws.

VII. SEPARABILITY CLAUSE

If any provision of this Agreement is declared null and void, invalid or contrary to law by a court of competent jurisdiction, all other provisions not affected shall remain in full force and effect, and binding upon the Parties.

VIII. REVISION OR AMENDMENT

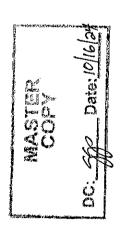
Any revision or amendment to this Agreement shall require a written consent from both parties.

IX. EFFECTIVITY

This Agreement shall commence and shall remain valid from	upon the signing of the PCEO of PhilHealth to
IN WITNESS WHEREOF, the part 2024.	ties have set their hands this day of
PHILIPPINE HEALTH INSURANCE CORPORATION	[NAME OF THE NETWORK]
By:	By:
President & CEO	Authorized Representative of the NETWORK
Signed PHILHEALTH	in the presence of:
PHILHEALTH	NETWORK

ACKNOWLEDGEMENT

REPUBLIC OF THE PHI	. *	
) S.S.	
BEFORE ME, this following persons, showing t	day ofo me their respective compe	, appeared the tent proofs of identity to wit
NAME	GOV'T I.D. NOS.	DATE /PLACE ISSUED
EMMANUEL R. LEDESM PCEO, PhilHealth	IA, JR.	
Head, NETWORK		
known to me to be the same HEALTH CARE PROVIDE INSURANCE CORPORA pages including to before me that the same is corporate entities they respectively.	R NETWORK between the TION and <i>(Name of 1)</i> his acknowledgment page. their free and voluntary a	PHILIPPINE HEALTH NETWORK) consisting of The Parties acknowledged
WITNESS MY HAND AN	D SEAL on the date and at	the place above written.
Doc. No Page No Book No Series of 202 .		



Annex B.1: List of Medicines Not Included Under PhilHealth GAMOT for CY 2024 (Already Provided through PhilHealth Konsulta)

		Molecule	Preparation
	1	Amlodipine	5 mg (as Besilate/ Camsylate) Tablet
	2	Amlodipine	10 mg (as Besilate/Camsylate) Tablet
	3	Amoxicillin	250 mg (as Trihydrate) Capsule
	4	Amoxicillin	500 mg (as Trihydrate) Capsule
	5	Amoxicillin	100 mg/mL (as Trihydrate) Granules/powder for drops in 15 mL
	************	A ***	250 mg/5 mL (as Trihydrate) Granules/powder for
	6	Amoxicillin	suspension in 60 mL
j	7	Amoxicillin + Clavulanic acid	500 mg (as Trihydrate) + 125 mg (as Potassium
İ	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		clavulanate) Tablet
	8	Amoxicillin + Clavulanic acid	875 mg (as Trihydrate) + 125 mg (as Potassium clavulanate) Tablet
	e e dechar des experie assesse sans	RT S S A RESEARCH AND STOCK AND A STOCK STOCK SECURITY OF THE STOCK STOCK STOCK SECURITY AND A STOCK STOCK SECURITY AS A STOCK SECURITY AS A STOCK SECURITY AS A STOCK SECURITY AS A STOCK STOCK SECURITY AS A STOCK STOCK SECURITY AS A STO	200 mg (as Trihydrate) + 28.5 mg (as Potassium
١	9	Amoxicillin + Clavulanic acid	clavulanate) per 5 mL, Granules/powder for suspension in
	9 10 11 12 13 14 15 16 17 18		70 mL
			400 mg (as Trihydrate) + 57 mg (as Potassium
	10	Amoxicillin + Clavulanic acid	clavulanate) per 5 mL, Granules/powder for suspension in
	P. P. St. of Street, was grown throughous suggested		70 mL
	11	Amoxicillin + Clavulanic acid	600 mg (as Trihydrate) + 42.9 mg (as Potassium clavulanate) per 5 mL, Granules/powder for suspension
Î	12	$\mathbf{Aspirin}$	80 mg Tablet
	13	Aspirin	100 mg Tablet
ľ	Charles the State of the State	Chlorphenamine	4 mg Tablets
ĥ	. 0. 12.0 . 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1	Chlorphenamine	2.5 mg / 5 mL syrup / 60 mL
۱.	Verbero v est a marro nome a	Ciprofloxacin	250 mg (as Hydrochloride) Tablet
-	. were green and a sough	Ciprofloxacin	500 mg (as Hydrochloride) Tablet
1	· • • • • · • · • · • · • · • · • · • ·	Clarithromycin	3.30 A 20 A
	ARA SA AL SANDO PROGRAMA		250 mg Tablet (Base)
,	2-06-22-020-024-2-00-0	Clarithromycin	500 mg Tablet (Base)
	20	Clarithromycin	125 mg/5 mL Granules/Powder for suspension in 50 mL
	21	Clarithromycin	250 mg/5 mL Granules/Powder for suspension in 50 mL
	22	Enalapril	5 mg (as Maleate) Tablet
125	23	Enalapril	20 mg (as Maleate) Tablet
	and an analysis of the second	A MARIANA A MARIAN A	100 mcg (as Propionate) + 50 mcg (as Xinafoate) x 28
1	24	Fluticasone + Salmeterol	doses, DPI with appropriate accompanying dispenser
1	***************************************		100 mcg (as Propionate) + 50 mcg (as Xinafoate) x 60
	25	Fluticasone + Salmeterol	doses, DPI with appropriate accompanying dispenser
			250 mcg (as Propionate) + 50 mcg (as Xinafoate) x 28
$\!$	26	Fluticasone + Salmeterol	doses, DPI with appropriate accompanying dispenser
1		a and a second	250 mcg (as Propionate) + 50 mcg (as Xinafoate) x 60
-	27	Fluticasone + Salmeterol	doses, DPI with appropriate accompanying dispenser
	~0		500 mcg (as Propionate) + 50 mcg (as Xinafoate) x 28
1	28	Fluticasone + Salmeterol	doses, DPI with appropriate accompanying dispenser
	00	Elutiangona Calmataval	500 mcg (as Propionate) + 50 mcg (as Xinafoate) x 60
Į	29	Fluticasone + Salmeterol	doses, DPI with appropriate accompanying dispenser

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MASTER COPTER	DC: 37 Date: pliblay

		Molecule	Preparation			
	30	Gliclazide	30 mg Modified Release (MR) tablet			
	31	Gliclazide	80 mg Tablet			
	32	Hydrochlorothiazide	12.5 mg Tablet			
	33	Hydrochlorothiazide	25 mg Tablet			
	34	Ipratropium Bromide + Salbutamol	500 mcg (as Bromide anhydrous) + 2.5 mg (as Base) in 2.5 mL Unit dose (For nebulization)			
	35	Losartan	50 mg (as Potassium) Tablet			
	36	Losartan	100 mg (as Potassium) Tablet			
i	37	Metformin	500 mg (as Hydrochloride) Tablet/Film-coated tablet			
j	38	Metformin	850 mg (as Hydrochloride) Tablet			
	39	Metoprolol	50 mg (as Tartrate) Tablet			
ĺ	40	Metoprolol	100 mg (as Tartrate) Tablet			
Ì	41	Nitrofurantoin	50 mg Capsule as Macrocrystals			
	42	Nitrofurantoin	100 mg Capsule as Macrocrystals			
Î	***************************************		(L of water) composed of:			
ŀ		and unanappear	Sodium chloride = 2.6 g			
***		and other than the state of the	Trisodium citrate dihydrate = 2.9 g			
		TOTAL TO A MARKET	Potassium chloride = 1.5 g			
			Glucose anhydrous = 13.5 g			
ŀ		of management of the state of t	Total weight = 20.5 g			
1		OC Date of Academic Academic	(mmol/L) composed of:			
			Sodium = 75			
		PROPERTY AND ADDRESS OF THE PROPERTY AND ADDRESS OF THE PROPERTY ADDRESS OF TH	Chloride = 65			
ľ		Co. Markon	Potassium = 20			
1		To the second se	Citrate = 10			
-1		No delication	Glucose anhydrous = 75 Total osmolarity = 245			
		Actions and the second	rotal osmolarity = 245			
۱		Oral Rehydration Salts (ORS	(WHO recommended)			
ļ.,	43	75-replacement)	(N.B.: Reconstitute with clean potable water)			
ļ	44	Paracetamol	300 mg Tablet			
	45	Paracetamol	500 mg Tablet			
	46	Paracetamol	100 mg/mL Drops in 15 mL (Alcohol-free)			
1			120 mg/5 mL (125 mg/5 mL) Syrup/Suspension			
ļ.,	******	Paracetamol	(Alcohol-free) in 60 ml			
	The state of the s		250 mg/5 mL Syrup/Suspension (Alcohol-free) in 60 mL			
49 Paracetamol		Same a complete production and a community of the communi	125 mg Suppository			
6.2		Paracetamol	250 mg Suppository			
	************	Prednisone	5 mg Tablet			
	THE REAL PROPERTY.	Prednisone	10 mg Tablet			
	AAAA AA TAA YYYYY	Prednisone	20 mg Tablet			
, A.	54	Prednisone	10 mg/5 mL Suspension in 60 mL			
			The street control of			

	Molecule	Preparation
55	Salbutamol (as Sulfate) + Ipratropium	Resp. Soln. (for nebulization): 500 micrograms ipratropium (as bromide anhydrous) + 2.5 mg salbutamol (as base) x 2.5 mL (unit dose)
56	Salbutamol	100 mcg/dose x 200 (as Sulfate)
57	Salbutamol	1 mg/mL (as Sulfate) Respiratory solution in 2.5 mL Unit dose (For nebulization)
58	Salbutamol	2 mg/mL (as Sulfate) Respiratory solution in 2.5 mL Unit dose (For nebulization)
59	Salbutamol	2 mg/5 mL (as Sulfate) Syrup in 60 mL
60	Simvastatin	20 mg Tablet
61	Simvastatin	40 mg Tablet
62	Sulfamethoxazole + Trimethoprim	400 mg + 80 mg Tablet/Capsule
63	Sulfamethoxazole + Trimethoprim	800 mg (as Sulfate) + 160 mg Tablet
64	Sulfamethoxazole + Trimethoprim	200 mg + 40 mg/5 mL Suspension in 70 mL
65	Sulfamethoxazole + Trimethoprim	200 mg + 40 mg/5 mL Suspension in 120 mL
66	Sulfamethoxazole + Trimethoprim	400 mg + 80 mg/5 mL Suspension in 60 mL



Annex B.2: GAMOT Basic Medication List for CY 2024

		Molecule				
	1	Atenolol	Preparation 50 mg Toblet			
	2	Atenolol	50 mg Tablet 100 mg Tablet			
	3	Atorvastatin	10 mg (as Calcium) Tablet			
	4	Atorvastatin	20 mg (as Calcium) Tablet			
	5	Atorvastatin	40 mg (as Calcium) Tablet			
	6	Atorvastatin	80 mg (as Calcium) Tablet			
	7	Azithromycin	NOTE OF THE PROPERTY AND A SECOND PROPERTY A			
	8	Azithromycin	250 mg (as Base*/dihydrate) Capsule			
	9	Azithromycin	250 mg (as Base*/dihydrate/ monohydrate) Tablet			
	У.	AZIUIIOIIIyem	500 mg (as Base*/dihydrate/ monohydrate) Tablet			
	10	Azithromycin	200 mg/5 mL (as Base*/dihydrate/ monohydrate)			
	10	FAZILIII OIII y CIII	Powder for suspension in 15 mL			
	11	Azithromyein	200 mg/5 mL (as Base*/dihydrate/ monohydrate) Powder for suspension in 60 mL			
	12	Captopril	25 mg Tablet			
i	13	Cefixime	12111110			
	13 14	Cefixime	200 mg Capsule 20 mg/mL in 10 mL (drops)			
	15	Cefixime				
	15 16	Cefuroxime	100 mg/5 mL Granules for Suspension in 60 mL			
	10	CETUI OXIIIIE	500 mg (as Axetil) Tablet			
	17	Cefuroxime	125 mg/5 mL (as Axetil) Granules for Suspension in 70 mL			
		COLUMN OXAMIC	# CONTROL OF THE PROPERTY OF T			
	18	Cefuroxime	250 mg/5 mL Granules for Suspension in 50 mL and 120 mL Bottle			
l	19	Celecoxib				
	20	Celecoxib	100 mg Capsule 200 mg Capsule			
1	21	Celecoxib	400 mg Capsule			
l	22	Clindamycin	***************************************			
ľ	23	Clindamycin	150 mg (as Hydrochloride) Capsule			
		CHINA	300 mg (as Hydrochloride) Capsule			
1	24	Clindamycin	75 mg/5 mL (as Palmitate hydrochloride) Granules for suspension in 60 mL			
ŀ	25	Clotrimazole	1% Cream in 10 g Aluminum collapsible tube			
ł	26	Cloxacillin	500 mg (as Sodium) Capsule			
Ì	27	Cloxacillin	CONTROL OF THE PROPERTY OF THE			
,	-/ 28	Diltiazem	250 mg/5 mL (as Sodium) Powder for solution in 60 mL 60 mg (as Hydrochloride) Tablet			
ŀ		10 11 11 11 11 11 11 11 11 11 11 11 11 1	Websites 1997 1997 1997 1997 1997 1997 1997 199			
	29	Diltiazem	60 mg (as Hydrochloride) Modified Release (MR) capsule			
	7-7		TOTAL CONTRACTOR AND A STATE OF THE STATE OF			
011010	30	Diltiazem	120 mg (as Hydrochloride) Modified Release (MR) capsule			
	ľ		180 mg (as Hydrochloride) Modified Release (MR)			
30.55	31	Diltiazem	capsule			
3	1 ~-		120 mg (as Hydrochloride) Modified Release (MR)			
_	32	Diltiazem	tablet			
7	¥ <u>~</u>	the format of the section and the section is a fig. of the transfer frame of the section is the respective to the section is the section in the section in the section is the section in the section in the section in t	180 mg (as Hydrochloride) Modified Release (MR)			
ľ	33	Diltiazem	tablet			
ŀ	34	Diltiazem	30 mg (as Hydrochloride) Tablet			
5	35	Diphenhydramine	25 mg (as Hydrochloride) Capsule			
***	36	Diphenhydramine	50 mg (as Hydrochloride) Capsule			
J.,,	TOTAL	alamente a managan da la companya d	. G (un ray ar control out of the			

	Molecule	Promaration			
37		Preparation			
38	TALLY TO A SECOND CONTROL OF THE SAME AND	12.5 mg/ 5mL (as Hydrochloride) Syrup in 30 mL			
, .	The state of the s				
39					
40	COLA A SERVA A BAB COMMENT Y YES TO COLA A CARRIER A REGION OF THE TOP TOP TO COLAR AS A CARRIER OF THE PROPERTY OF THE COLAR AS A CARRIER OF THE PROPERTY OF THE COLAR AS A CARRIER OF THE COLAR AS A C	ANDERS OF THE THE COLD AND A DESCRIPTION OF THE COLD AND A DESCRIP			
41	to related from a construction of the first state of the construction of the first state of the construction of the constructi	The state of the s			
42	A CAMPAG AND COMMITTEE OF THE SECTION AND ADMITTACK AND CONTRACTION ASSESSMENT AND ADMITTACK	**************************************			
43	от выправления в при в при на выправления на выправления и при на выправнительной в при на выправления в при н В при на выправнительной в при на выправления в при на выправнительной в принентической в при на выправнительной в принентичения в при на выправнительной в принентичения в принентичения в при	500 mg (as Stearate) Tablet			
44	Erythromycin	200 mg/5 mL (as Ethyl for suspension in 60 mL			
	remain avera	60 mg (elemental iron) + 400 mcg			
45		Tablet/Capsule/Film-coated tablet			
46	A CONTROL OF THE PROPERTY OF T	100 mg Capsule			
47	Gabapentin	300 mg Capsule			
48	Ibuprofen	200 mg Capsule			
49	Ibuprofen				
50	Ibuprofen	are really as a mean of the contract of the co			
51	Ipratropium Bromide				
2-1966/10/06/01/0	·····································				
	70.000.00	(N.B.: The elemental iron content of a ferrous salt			
		20 mg (as Maleate) Tablet 0.5% Ophthalmic ointment in 3.5 g Tube 500 mg (as Stearate) Tablet 200 mg/5 mL (as Ethyl for suspension in 60 mL 60 mg (elemental iron) + 400 mcg Tablet/Capsule/Film-coated tablet 100 mg Capsule 300 mg Capsule 200 mg Capsule 400 mg Capsule 100 mg/5 mL Syrup/Suspension in 60 mL 250 mcg/mL (as Bromide) Respiratory solution in 2 Unit dose (For nebulization) (equiv. to 60 mg elemental iron) Tablet (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous fumarate = 33% Ferrous gluconate = 12% Ferrous sulfate, hydrated = 20% Ferrous sulfate, desiccated = 32%) (equiv. to 15 mg elemental iron per 0.6 mL) Solution 15 mL (Drops) (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous fumarate = 33% Ferrous gluconate = 12% Ferrous sulfate, hydrated = 20% Ferrous sulfate, desiccated = 32%)			
	100000000000000000000000000000000000000	12.5 mg/ 5mL (as Hydrochloride) Syrup in 6o mL 100 mg (as Hyclate) Capsule 5 mg (as Maleate) Tablet 20 mg (as Maleate) Tablet 0.5% Ophthalmic ointment in 3.5 g Tube 500 mg (as Stearate) Tablet 200 mg/5 mL (as Ethyl for suspension in 60 mL 60 mg (elemental iron) + 400 mcg 13 Tablet/Capsule/Film-coated tablet 100 mg Capsule 300 mg Capsule 200 mg Capsule 400 mg Capsule 100 mg/5 mL Syrup/Suspension in 60 mL 250 mcg/mL (as Bromide) Respiratory solution in Unit dose (For nebulization) (equiv. to 60 mg elemental iron) Tablet (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous fumarate = 33% Ferrous lactate = 19% Ferrous sulfate, hydrated = 20% Ferrous sulfate, desiccated = 32%) (equiv. to 15 mg elemental iron per 0.6 mL) Solutio 15 mL (Drops) (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous fumarate = 33% Ferrous flactate = 19% Ferrous fundate = 12% Ferrous flactate = 19% Ferrous fundate = 12% Ferrous fundate = 20% Ferrous sulfate, desiccated = 32%) (equiv. to 15 mg elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous sulfate, desiccated = 32%) (equiv. to 15 mg elemental iron per 0.6 mL) Solutio 30 mL (Drops) (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous fumarate = 33% Ferrous fundate = 33% Ferrous fundate = 12% Ferrous fundate = 33% Ferrous fu			
	TO A COLUMN STATE OF THE STATE	100 mg/5 mL Syrup/Suspension in 60 mL 250 mcg/mL (as Bromide) Respiratory solution in 2 Unit dose (For nebulization) (equiv. to 60 mg elemental iron) Tablet (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous fumarate = 33% Ferrous gluconate = 12% Ferrous lactate = 19% Ferrous sulfate, hydrated = 20% Ferrous sulfate, desiccated = 32%) (equiv. to 15 mg elemental iron per 0.6 mL) Solution 15 mL (Drops) (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous fumarate = 33%			
		500 mg (as Stearate) Tablet 200 mg/5 mL (as Ethyl for suspension in 60 mL 60 mg (elemental iron) + 400 mcg Tablet/Capsule/Film-coated tablet 100 mg Capsule 300 mg Capsule 200 mg Capsule 400 mg Capsule 100 mg/5 mL Syrup/Suspension in 60 mL 250 mcg/mL (as Bromide) Respiratory solution in 2 mL Unit dose (For nebulization) (equiv. to 60 mg elemental iron) Tablet (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous flumarate = 33% Ferrous gluconate = 12% Ferrous sulfate, hydrated = 20% Ferrous sulfate, desiccated = 32%) (equiv. to 15 mg elemental iron per 0.6 mL) Solution in 15 mL (Drops) (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous flumarate = 33% Ferrous gluconate = 12% Ferrous lactate = 19% Ferrous sulfate, hydrated = 20% Ferrous sulfate, hydrated = 20% Ferrous flumarate = 33% Ferrous flumarate = 33% Ferrous flate, desiccated = 32%) (equiv. to 15 mg elemental iron per 0.6 mL) Solution in 30 mL (Drops) (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous sulfate, desiccated = 32%) (equiv. to 15 mg elemental iron per 0.6 mL) Solution in 30 mL (Drops) (N.B.: The elemental iron content of a ferrous salt depends on the type of preparation as follows: Ferrous flumarate = 33% Ferrous gluconate = 12% Ferrous lactate = 19% Ferrous lactate = 19% Ferrous sulfate, hydrated = 20% Ferrous lactate = 19% Ferrous sulfate, hydrated = 20% Ferrous lactate = 19% Ferrous sulfate, hydrated = 20%			
52	Iron Ferrous				
	7 A A SOC A A SOC				
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		depends on the type of preparation as follows:			
	Security Sec	Ferrous fumarate = 33%			
	TO A	Ferrous gluconate = 12%			
	NO COMPANY AND ADDRESS AND ADD				
53	Iron Ferrous	Ferrous sulfate, desiccated = 32%)			
	THE PROPERTY OF THE PROPERTY O	(equiv. to 15 mg elemental iron per 0.6 mL) Solution in			
	C	, · · · · · · · · · · · · · · · · · · ·			
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54	Iron Ferrous				
	PRINCIPALITY				
4	Assessables	· (· · · · · · · · · · · · · · · · · ·			
	St. O. Commonweal				
	Inon Formor-	Ferrous fumarate = 33%			
55	Iron Ferrous	Ferrous gluconate = 12%			

155 65	Molecule					
		Preparation 2				
	1999	Ferrous lactate = 19%				
	NET CALLANA	Ferrous sulfate, hydrated = 20% Ferrous sulfate, desiccated = 32%)				
56	isosorbide Dinitrate	\$87 (10 f 10				
	\$	10 mg (as Dinitrate) Tablet				
57	TROSOLDING DIMITAGE	20 mg (as Dinitrate) Tablet				
58	Isosorbide Dinitrate	20 mg (as Dinitrate) Modified Release (MR) tablet/Capsule				
		THE RESIDENCE AND ASSOCIATION OF THE PROPERTY				
59		5 mg (as Dinitrate) Sublingual (SL) tablet				
60	Isosorbide Mononitrate	30 mg (as 5-Mononitrate) Modified Release (MR)				
	TAGNODIUC MUMOMMA PROPRESSION OF THE PROPRESSION OF	tablet/Capsule				
61	Isosorbide Mononitrate	60 mg (as 5-Mononitrate) Modified Release (MR)				
62	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	tablet/Capsule				
Address of the Asperture of the	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	50 mg losartan + 12.5 mg hydrochlorothiazide Tablet				
63	opport of the same temporal contract	250 mg Tablet /Capsule				
64		500 mg Tablet /Capsule				
65	**************************************	250 mg Tablet				
66	**************************************	250 mg Tablet				
67	TOTAL CONTROL OF THE STATE OF T	500 mg Tablet				
68	TO THE STATE OF TH	125 mg/5 mL (as Base) Suspension in 60 mL				
69	varant como roma roma en vara en vara en vara en mana anama competan persona como como como a secución en vara	4 mg (as Sodium) Chewable tablet				
70	CONTROL CONTROL AND A VARIANCE OF THE CONTROL AND A MARKATANA AND A A A A A A A A A A A A A A A A	5 mg (as Sodium) Chewable tablet				
71	**************************************	10 mg (as Sodium) Tablet				
72	TO THE STATE OF TH	275 mg (as Sodium) Tablet				
73	4. 2.72, ATTENDADO - 100	550 mg (as Sodium) Tablet				
74	ann francisco mentendententententententententententententente	75 mg (as Phosphate) Capsule				
75	O TOTAL OF THE STATE A A STATE AND A STATE OF THE STATE O	10 mg (as Calcium) Tablet				
76	YYE: # 10 10 10 10 10 10 10 10 10 10 10 10 10	20 mg (as Calcium) Tablet				
	Tamsulosin	200 mcg (as Hydrochloride) Capsule				
78		0.3% Ophthalmic drop solution in 5 mL Bottle				
79	Tobramycin	0.3% Ophthalmic ointment in 3.5 g Tube				
		0.3% tobramycin + 0.1% dexamethasone Ophthalmic				
80	Tobramycin + Dexamethasone	drop suspension in 5 mL Bottle				
		0.3% tobramycin + 0.1% dexamethasone Ophthalmic				
81	Tobramycin + Dexamethasone	ointment in 3.5 g Tube				
82	***************************************	300 mg Tablet				
83		600 mg Tablet				
84	where for an ever a construction and the property of the construction and the construction and a construction of the construct	300 mg/5 mL Syrup in 60 mL				
85	Vitex Negundo	300 mg/5 mL Syrup in 120 mL				
	To the state of th	(equiv. to 10 mg elemental zinc) (as Gluconate)				
86	Zinc	Chewable tablet				
	33.00	(equiv. to 10 mg elemental zinc per mL) (as Sulfate				
87	Zinc	monohydrate) Drops in 15 mL				
	The second secon	(equiv. to 20 mg elemental zinc per 5 mL) (as Sulfate				
88	Zinc	monohydrate) Syrup in 60 mL				
,		70 mg/5 mL (equiv. to 10 mg elemental zinc) (as				
89	Zinc	Gluconate) Syrup in 60 mL				
	The state of the s	70 mg/5 mL (equiv. to 10 mg elemental zinc) (as				
90	Zinc	Gluconate) Syrup in 120 mL				

Annex B.3: Summary of Basic Medication List Molecules for CY 2024

	Basic Medication List	Communication of the second of
Anti-infectious 1. Azithromycin 2. Cefixime 3. Cefuroxime 4. Clindamycin 5. Clotrimazole 6. Cloxacillin 7. Doxycycline 8. Erythromycin 9. Metronidazole 10. Oseltamivir 11. Tobramycin Anti-asthma and COPD 12. Montelukast	Supportive/Other Therapy 13. Celecoxib 14. Diphenhydramine 15. Elemental Iron 16. Folic acid + Iron Ferrous 17. Ibuprofen 18. Mefenamic Acid 19. Naproxen 20. Vitex Negundo (Lagundi) 21. Zinc Anti-dyslipidemia 22. Atorvastatin	Anti-hypertensive and Cardiology 24. Atenolol 25. Captopril 26. Diltiazem 27. Enalapril 28. Isosorbide Dinitrate 29. Isosorbide Mononitrate 30. Methyldopa 31. Tamsulosin Nervous System 32. Gabapentin
	23. Rosuvastatin	



Annex C: Additional 13 Formulations for PhilHealth GAMOT for CY 2025

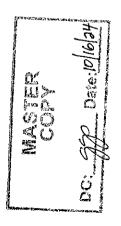
2 S. E.	Molecule	Preparation			
1	Aspirin	300 mg Tablet			
2	Aspirin	325 mg Tablet			
3	Enalapril + Hydrochlorothiazide	20 mg enalapril + 12.5 mg hydrochlorothiazide tablet			
4	Fluticasone + Salmeterol	Metered-Dose Inhaler (MDI): 50 mcg (as Propionate) + 25 mcg (as Xinafoate) x 120 actuations (with dose counter)			
5	Fluticasone + Salmeterol	Metered-Dose Inhaler (MDI): 125 mcg (as Propionate) + 25 mcg (as Xinafoate) x 120 actuations (with dose counter)			
6	Fluticasone + Salmeterol	Metered-Dose Inhaler (MDI): 250 mcg (as Propionate) + 25 mcg (as Xinafoate) x 120 actuations (with dose counter)			
7	Gliclazide	60 mg Modified Release (MR) tablet			
8	Paracetamol	120 mg/5 mL (125 mg/5 mL) Syrup/Suspension (Alcohol-free) in 30 ml			
9	Paracetamol	120 mg/5 mL (125 mg/5 mL) Syrup/Suspension (Alcohol-free) in 120 ml			
10	Paracetamol	250 mg/5 mL Syrup/Suspension (Alcohol-free) in 30 ml			
Paracetamol ml Dry Powder Inhaler (DPI):		250 mg/5 mL Syrup/Suspension (Alcohol-free) in 120 ml			
		200 mcg/dose with appropriate accompanying			
13	Ipratropium Bromide + Salbutamol	Metered-Dose Inhaler (MDI): 20 mcg ipratropium + 100 mcg salbutamol x 200 doses x 10 mL			



Annex D: GAMOT Contract Amount for CY 2024 Sample Computation

Step 1	 A. Determine the following: The calendar year and months in which the GAMOT will be implemented The number of months to be covered in SLA B. Obtain the sales/dispensing records of the health facility within the immediately complete years three (3) years (or applicable complete years for facilities with less than 3 years but at least 1 year of operations) 	Sample: Facility North Star Pharmacy Implementation year Implementation Implemen				
Step 2	List and identify annual historical sales included in GAMOT*^	es (or dispensing) volume of drugs				
Step 3	Compute for the average annual sales volume by adding the annual sales volume from the the applicable complete years and dividing the sum by number of years (i.e., 3, 2)					
Step 4	Compute the monthly average sales volume by dividing the annual sales by 12					
Step 5	Multiply the monthly average sales volume by the number of months covered in SLA (e.g. three [3] for October – December)					
Step 6	Multiply the computed sales volume by contract amount for each molecule and		nine the			
Step 7	Add the contract amount of all applicable molecules and preparations to determine the contract amount for a GAMOT Provider (Round up the contract amount to the nearest thousands)					

^{*32} molecules (90 preparations) covered under GAMOT for CY 2024 ^53 molecules (169 preparations) covered under GAMOT for CY 2025



Step 2					Step 3	Step 4	Step 5	St	ер 6
Molecule+	Preparations+	CY 2021	CY 2022	CY 2023	Annual Average Sales Volume	Monthly Average Sales Volume	3 Months Sales Volume	Fixed Fee	Contract Amount per Molecule
Atenolol	50 mg Tablet	307	326	.345	326.00	27.17	81.50	₱8.01	₱652.82
Atenolol	100 mg Tablet	205	254	366	275.00	22.92	68.75	₱11.55	₱794.06
Atorvastatin	10 mg (as Calcium) Tablet	790	780	812	794.00	66.17	198.50	₱6.00	₱1,191.0o
Atorvastatin	20 mg (as Calcium) Tablet	799	609	900	769.33	64.11	192.33	₱ 12.00	₱2,308.00
Atorvastatin	40 mg (as Calcium) Tablet	2 34	546	790	523.33	43.61	130.83	₱16.00	₱2,093.33
Atorvastatin	80 mg (as Calcium) Tablet	765	468	789	674.00	56.17	168.50	₱25.00	₱4,212.50
Azithromycin	250 mg (as Base*/dihydrate) Capsule	800	678	234	570.67	47.56	142.67	₱98.79	₱ 14,094.04
Azithromycin	250 mg (as Base*/dihydrate/ monohydrate) Tablet	234	456	678	456.00	38.00	114.00	₱99.79	₱11,376.06
Azithromycin	500 mg (as Base*/dihydrate/ monohydrate) Tablet	400	456	676	510.67	42,56	127.67	₱53.50	₱6,830.17
Azithromycin	200 mg/5 mL (as Base*/dihydrate/ monohydrate) Powder for suspension in 15 mL	300	678	679	552.33	46.03	138.08	₱270.00	₱37,282.50
Azithromycin	200 mg/5 mL (as Base*/dihydrate/ monohydrate) Powder for suspension in 60 mL	200	345	670	405.00	33.75	101.25	₱381.74	₱38,651.18
Step 7 Contract Amount						485.66 000.00)			

+Partial list for sample computation purpose only

Note: The annual sales volume will be multiplied by the fixed fee to determine the contract amount for each molecule/preparations for CY 2025.



Annex E: ACR-GB Contract Amount Sample Computation

Step 1 Determine the following: Health facility which will implement the ACR and its service level The calendar year in which the ACR-GB will be implemented Extract the ACR claims of the health facility within the immediately complete years three (3) years, except for the operational years of 2020 and 2021 (date of admission as the reckoning date)	Facility Service level Implementation year Years of historical claims needed for estimation Note: CY 2024 is n	2019
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STEP 2		Step 3		
Determine case from t	Determine the total count of paid claims per case from the applicable three (3) years		count of paid claims per ponding current case rate laims amount per case	
Code	Total Count of Paid Claims per Case (CY 2023, CY 2022, CY 2019)	Current Case Rate	Total Claims Amount per Case (CY 2023, CY 2022, CY 2019)	
44950	64	31,200.00	1,996,800.00	
44960	55	31,200.00	1,716,000.00	
58120	79	14,300.00	1,129,700.00	
59409	82	12,610.00	1,034,020.00	
59513	169	24,700.00	4,174,300.00	
59514	222	24,700.00	5,483,400.00	
A01.0	40	13,000.00	520,000.00	
E11.9	44	5,200.00	228,800.00	
H81.1	41	7,410.00	303,810.00	
J06.9	70	5,200.00	364,000.00	
K29.1	188	7,930.00	1,490,840.00	
060.0	47	8,840.00	415,480.00	
T00.9	139	5,200.00	722,800.00	
Step 4	Determine the contract amount by adding claims amount of all applicable cases, and dividing the sum by three (3) (Round up the contract amount to the nearest thousands)	Contract Amount	19,579,950.00 ÷ 3 = 6,526,650.00 (6,527,000.00)	
Step 5	Multiply by 10% to determine the supplementary budget (This step is only done for facilities eligible for supplementary budget [e.g., L1-L3 hospitals]) (Round up the contract amount to the nearest thousands)	Supplementary Budget	652,665.00 (653,000.00)	
Step 6	Add the contract amount and supplementary budget to determine the maximum contract amount	Maximum Contract Amount	7,180,000.00	

Annex F: ACR-GB Case Rates Adjustment Sample Computation

Step 1	Determine the following: A. Cases with rate adjustment B. Period covered by the case rates adjustment, from the date of effectivity (as stipulated in relevant PhilHealth issuance) to the end of contract period C. Number of months covered by the case rates adjustment	Sample: Facility Service level Period covered by case rates adjustment Number of months covered by case rates adjustment Cases with rate adjustment	Filipino Hospital L1 hospital Date of effectivity - end of contract July - December, 2024 6 months 59409 59513 59514 E11.9 K29.1 O60.0 Too.9			
Step 2 Step 3	Multiply the average monthly count of paid claims per case by the number of months (from the effectivity of adjustment to the end contract period) to					
Step 4	Determine the difference between the old case rates and new case rates after the case rates adjustment.					
Step 5	Multiply the total count of paid claims per case during the period of adjustment with the difference between the old case rates and new case rates to determine the adjustment per case.					
Step 6	Add the adjustment for all applicable cases to determine the total adjustment to the contract amount (Round up to the nearest thousands)					



Step 2		Step 3	Step 4	Step 5		
Code	Total Count of Paid Claims per Case (CY 2023, CY 2022, CY 2019)	Average Annual Count of Paid Claims per Case	Average Monthly Count of Paid Claims per Case	Total Count of Paid Claims per Case during the period of rate adjustment	Difference between old case rates and new case rates	Adjustment per Case
44950	64	21.33	N/A	N/A	N/A	N/A
44960	55	18.33	N/A	N/A	N/A	N/A
58120	79	26.33	N/A	N/A	N/A	N/A
59409	82	27.33	2.28	13.67	3,783.00	51,701.00
59513	169	56.33	4.69	28.17	7,410.00	208,715.00
59514	222	74.00	6.17	37.00	7,410.00	274,170.00
A01.0	40	13.33	N/A	N/A	N/A	N/A
E11.9	44	14.67	1.22	7.33	1,560.00	11,440.00
H81.1	41	13.67	N/A	N/A	N/A	N/A
J06.9	70	23.33	N/A	N/A	N/A	N/A
K29.1	188	62.67	5.22	31.33	2,379.00	74,542.00
O60.0	47	15.67	1.31	7.83	2,652.00	20,774.00
T00.9	139	46.33	3.86	23.17	1,560.00	36,140.00
Step 6			Total	l Adjustment	625,781.00 (626,000.00)	



Annex G: ACR-GB Case Rates Adjustment Frontloading Schedule

Table G.1 Frontloading Schedule of Case Rate Adjustment

Tranches	Percentage of ACR-GB	Percentage of Total Case Rate Adjustment		
Tranche 1	60%	60%		
Tranche 2	30%	30%	90%	
Tranche 3	10%	10%	10%	100%
		Scenario A: Case rates adjustment happening before the frontloading of 1st tranche payment	Scenario B: Case rates adjustment happening before the frontloading of 2nd tranche payment	Scenario C: Case rates adjustment happening before the frontloading of 3rd tranche payment

Table G.2 Sample Frontloading of Case Rate Adjustment

Tranches	Amount of	Amount of Case Rate Adjustment^			
ACR-GB*		Scenario A	Scenario B	Scenario C	
Tranche 1	3,916,200.00	375,600.00			
Tranche 2	1,958,100.00	187,800.00	563,400.00	•	
Tranche 3	652,700.00	62,600.00	62,600.00	626,000.00	

Examples provided in Annex E^* and F^* .



Annex H: Sample Reconciliation of ACR-GB

Particulars Description		Sample Computation		
Unutilized fund before reconciliation (a)	Remaining/ unutilized fund at the end of contract period and prior to reconciliation	Given: 150,000.00	= 150,000.00	
1st tranche amount (b)	Computed 1st tranche based on SLA (contract amount x .60)	Example: 1,000,000.00 (contract amount) x .6	= 600,000.00	
Frontloaded 1st tranche amount before reconciliation (c)	The amount remaining after the unutilized fund is deducted from the 1st tranche amount. (b - a)	Example: 600,000.00 - 150,000.00	= 450,000.00	
Unutilized fund after reconciliation (d) (within 90 days from the expiration of the contract)		Given: 50,000.00	=50,000.00	
Remaining Amount to be front loaded for the 1st tranche (e)	The amount to be front loaded to complete the 1st tranche amount (a-d)	Example: 150,000.00 - 50,000.00	=100,000.00	

