Outpatient Pediatric Special Care. Developing health policies at the primary level of care

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ABSTRACT
In 1978, the World Health Organization defined primary health care as the cornerstone of global health. In recent years, the usual causes of mortality (infections, deficiency diseases) now include chronic and degenerative diseases. It is now possible to provide tertiary care to patients without adequate resources for their survival after discharge. The health system has not been able to adapt to such change and has maintained a fragmented, reactive, disease-centered model.

“Selective primary care” is useful to redirect primary care bases according to the characteristics of the population. A set of specific measures, targeted at a population with specific vulnerabilities, leverages the power of primary health care. Here we describe the “Outpatient Pediatric Special Care” model of care implemented at the Center for Health and Community Action no. 5, in the Autonomous City of Buenos Aires.

Key words: primary health care, child care, public health policies, chronic diseases, follow-up.

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INTRODUCTION

“Health for All by the Year 2000.” This was a well-known utopia, but it paved the way. The Alma-Ata Declaration provided guidance: primary health care (PHC) as the cornerstone of global health.1 The reformulation of certain objectives, centered on equity, social justice, and solidarity as basic new conditions of a novel strategy, was closely related to necessary and expected changes in pediatrics.2

Since then, social, economic, cultural, and epidemiological changes have taken place that significantly modified the transition of health; as a result, new challenges that need to be faced have emerged. The rapid advances in knowledge and technology have helped to reduce mortality in relation to extreme prematurity, complex congenital heart diseases, tumors, etc.; however, the so-called “new morbidities” have then emerged as a result of such survival. These morbidities have turned into chronic conditions, with a major impact on health, and this has also left patients with associated vulnerabilities (housing, financial, cultural conditions, etc.) even more exposed.

This situation was perceived clearly at the Center for Health and Community Action (Centro de Salud y Acción Comunitaria, CeSAC) no. 5. It is worth mentioning some characteristics of the population seen at this facility: it is located in a “critically” vulnerable area;3 most people live below the poverty line and have unmet basic needs (UBNs), with no access to clean running water or sewer systems, a poor environmental sanitation, etc. The CeSAC no. 5 is located in an urban neighborhood in Community 8 of the Autonomous City of Buenos Aires (CABA), with a high rate of several generations of migrant population. Together with other neighborhoods, Community 8 is in the group with the highest socioeconomic exclusion levels in a city that, paradoxically, flaunts the highest per capita income nationwide. In Community 8, the mortality rates and the public health coverage contrast with the
indicators registered in CABA (Table 1). The coexistence of such contrasting scenarios leads to an epidemiological polarization phenomenon, where the first and third stages of epidemiological transition coexist and overlap. The same region accommodates cohorts with a high mortality rate due to infections (many of which are preventable), deficiencies, a high infant mortality rate (IMR), and a low life expectancy at birth (LEB), together with another group with a higher mortality rate due to chronic and degenerative diseases, a low IMR, and a high LEB. In turn, there are tertiary care facilities with sufficient resources to provide care for complex pediatric conditions, capable of reducing neonatal and infant mortality, but which discharge high-risk patients (with various types of sequelae, polypharmacy and/or technology care requirements) into a setting with inadequate socioeconomic and housing resources for their survival. A clear example are extreme preterm infants that survive thanks to a very good perinatal care who, after weeks of receiving complex care, face situations with profound personal and nutritional deficiencies, severe weather conditions, multiple risk factors for acute lower respiratory tract infections (ALRTIs), insufficient support from the Public Health System (PHS), etc. In addition to their high biological vulnerability, they suffer an extensive socioeconomic vulnerability. The PHS, as such, has not been able to adapt to this change and has maintained a fragmented, reactive, disease-centered model.

For years, the pediatricians working at the CeSAC no. 5 have coped with this health problem. This health care approach was split between walk-in appointments assigned in the same day and scheduled appointments. Patient access was determined by the health care team possibilities, order of arrival, and assignment of appointments. The population was forced to adapt to this model. The parents of these patients, who have greater care demands and vulnerabilities, had to “compete” with the rest to ensure care for their children. Their possibilities were usually impaired; their demands were rejected or they were forced to accept off-schedule appointments that could not always be met.

### PLANNING A NEW APPROACH: Selective Primary Care

In 1999, in a health care center located in one of the poorest neighborhoods of CABA, Mario Ripoli, M.D., established a new essential tool in his work: “selective primary care,” which helped to redirect the bases of PHC according to the characteristics of the population. A set of specific measures, targeted at a population with specific vulnerabilities, leveraged the power of PHC. Without leaving primary prevention and the “comprehensive” action model aside, “selective” measures adapted to the prevalent disease were proposed, especially for highly vulnerable groups, together with secondary and tertiary prevention measures. This way, the advantages of PHC complemented the effectiveness of selective models.

Given that “exceptional” circumstances had become usual, it was decided to adopt a new approach: indiscriminately assigning off-schedule appointments was not enough; it was necessary to tackle a complex problem from a broad and proactive perspective, with formal and informal

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**Table 1. Comparison between the Autonomous City of Buenos Aires and Community 8. Demographic, social, and health indicators**

| Indicator | CABA                  | Community 8           |
|-----------|-----------------------|-----------------------|
| Total population | 3 059 122 inhab. | 225 737 inhab. |
| Rate of natural increase | 2.7    | 8.7          |
| Overcrowding | 8.70 %    | 17.20 %      |
| “Critical” overcrowding | 1.30 %  | 3.80 %       |
| Households with income < total basic basket | 25 %    | 54.30 %      |
| Single public health coverage | 20 %    | 49.60 %      |
| IMR per 1000 LBs | 7.2      | 9.7          |
| IMR due to reducible causes | 4       | 6.7          |
| IMR due to hardly reducible causes | 2       | 1.1          |
| NMR | 5.4      | 5.8          |
| PNMR | 1.8      | 3.9          |
| Average age at death - Males (years old) | 73.6    | 66.2         |
| Average age at death - Females (years old) | 81      | 75           |

**Source:** Statistical Yearbook 2016 - General Department of Statistics, Surveys and Censuses, Government of the Autonomous City of Buenos Aires. CABA: Autonomous City of Buenos Aires; IMR: infant mortality rate; LBs: live births; NMR: neonatal mortality rate; PNMR: post-neonatal mortality rate.
resources that would allow to cross the borders existing among and within the different levels of care.

The main objective of such approach was to minimize the rejection of high-risk patients and centralize care by bringing the different levels of care closer. The objectives also included reducing the redundant transit among facilities by rapidly identifying risk situations to take more timely measures.

Health is determined at different levels, so our response needed to be systemic and multidimensional:

- **Social determination:** recognizing socioeconomic characteristics and integrating measures with social services provided by the health care team, maintaining a closer contact with patients and their reality and establishing a greater compliance with follow-up. In addition, collaborating with the application for allowances, benefits or pensions, if applicable.

- **Institutional/health system determination:** due to the scarce organization and the lack of system coordination, an attempt was made to warrant permanent access, in addition to the management of appointments at the different levels of care (specialist consultations, ancillary tests, etc.) by simplifying access to appointments. A complete survey of diseases associated with groups of high IMR by age and cause was carried out to establish their relation to the PHS. The project sought to bring the system closer to patients with chronic conditions or sequelae, who showed a high dependence on the system and greater access difficulties.6,7

- **Biological determination:** working with patients based on their characteristics to reduce their risks (nutritional risks, immunizations, early detection of diseases or sequelae, follow-up of connatal infections, early care of genetic disorders, etc.).

From the beginning, it was observed that this approach could not be handled only by the Department of Pediatrics; instead, it was necessary for the health care team to work in a transdisciplinary manner and to incorporate decisions among the different disciplines of the team (Social Services, Anthropology, Mental Health, Nutrition, Pediatrics, Nursing, etc.). Our “special” patients corresponded to all these disciplines and a network-like operation was necessary to engage, integrate, and protect them.

**METHOD**

The problem was addressed based on the “Langley enhancement model.”9 Problems to be solved were the basis for planning; initial measures based on such planning were carried out; results were analyzed and actions were taken upon them. Such dynamic approach allowed to improve the process on an ongoing basis (plan-do-analyze-act cycle). We required a resilient and innovative response.

**POPULATION, RESOURCES, TOOLS, AND FIRST STEPS**

- **Population**

  The population of Community 8 (CABA) was 187,237 inhabitants (National Household Survey 2010); of them, 59,363 (31.7 %) were children.9 The estimated population of Villa 15 (one of the clusters with the highest population density in the Community) was more than 30,000 inhabitants (due to housing irregularities, such estimation was based on historical demographic data). As per our current records, 9,745 inhabitants are younger than 18 years, including this neighborhood. And there is a number of patients that cannot be estimated that come from surrounding neighborhoods and Greater Buenos Aires, which is adjacent to Community 8.

- **Physical resources**

  The CeSAC no. 5, located in Villa Lugano of Community 8 (CABA), was moved to its new location in 2004. It has a surface area of 1,230 m², 35 medical offices, 2 dental offices, a laboratory sample collection room, an immunization room, a pharmacy, an X-ray room, an ultrasound room, and a teaching room (source: CeSAC no. 5).

- **Human resources**

  In 2015, the CeSAC workforce included 70 employees (both health care providers and non-professional staff), as detailed in Annex 1 (Spanish version).

- **Development and implementation of an outpatient pediatric special care model**

  Initially, participants for inclusion in the health care model were defined. First of all, diagnoses strongly associated with infant mortality were included.10 Secondly, patients with chronic diseases and a high need for contact with the PHS were considered. Risk groups and inclusion criteria were defined (Table 2). The health care team was informed so that every patient that met the criteria was referred to the program officers for their inclusion and, thus, to engage the largest number possible of the target population. A...
database was designed; it included demographic data, diagnoses, date of inclusion, latest visit and next appointment, discharge date, and reason for discharge (discharge, death, transfer, etc.). A specially designed medical record was developed (Annex 2 Spanish version) and incorporated in a separate file available at the department. Every patient susceptible to admission into the model was interviewed, the reasons for inclusion, benefits, and liabilities were explained, and they were asked for their consent to participate. One hundred per cent of patients agreed to participate. (An informed consent was not requested because the project did not imply changes in patient care or the dissemination of information outside the CeSAC). The policy for access to care was defined, and appointments were reorganized so that care was warranted to our patients in 100 % of contacts with the CeSAC. Once included, they were able to access the center any time they requested it by identifying themselves as “Outpatient Pediatric Special Care” (OPSC) patients at the reception desk. Appointments at the second level of care were also coordinated to meet different medical needs (Pediatric Neurology, Preterm Infant Follow-up, Day Hospital, Hospitalization, Ancillary Tests, Special Immunizations, etc.). The entire health care team, including nurses, pharmacists, social workers, general practitioners, nutritionists, administrative staff, etc., had an active participation by alerting on inclusion criteria and changing the approach to these patients. This enabled a rapid adaptation to specific circumstances, such as the provision of drugs and newborn formula, the immediate identification of risk groups for immunization in epidemic outbreaks (e.g., H1N1 flu in 2009), among others.

Once the program organization was completed, the different management levels were informed: CeSAC Management, Department of Pediatrics, and Hospital Santojanni Management. During subsequent visits, the authorities of the Department of Health were also informed, and the project was made public. The response at these levels was favorable.

In March 2004, health care under the OPSC modality started being provided.

**EXPERIENCE AND OUTCOMES**

Between 2005 and 2018, an average of 37 787 total visits/year (± 6348.66) took place at the CeSAC no. 5; 20 464 (± 3191.6) of these corresponded to pediatric visits/year. Pediatric visits accounted for 54.16 % of total visits. Since its launch in March 2004 until January 2019, 946 patients were seen under the OPSC modality: 559 (59.1 %) were males and 387 (40.9 %), females. In turn, these patients were classified into 6 groups (Table 3).

The table also shows the number of patients per group admitted in that period. Figure 1 shows the rate of patient inclusion in the first weeks. In this 15-year period, 201 patients were discharged from

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**Table 2. Inclusion criteria for the Outpatient Pediatric Special Care program (at the time of its launch in 2004)**

| Inclusion criteria                                                                                     |
|--------------------------------------------------------------------------------------------------------|
| • Children born at a gestational age of less than 37 weeks and with a current chronological age younger than 6 years. |
| • Children younger than 1 year and born with a birth weight of less than 2500 g.                        |
| • Children with moderate to severe neurological development disorders, based on clinical criteria.        |
| • Children with congenital infections in the process of diagnosis, treatment or sequelae management.      |
| • Children with genetic disorders or congenital abnormalities that require ongoing follow-up.            |
| • Children with organ transplant, cancer, chronic kidney disease, rheumatic disease, chronic pulmonary disease, cardiovascular disease or requiring neurosurgery. |

**Table 3. Distribution by group at the time of admission to the program (2004-2019)**

| Patients Proportion |
|---------------------|
| Group 1 (clinical conditions) | 246 | 26 % |
| Group 2 (surgical conditions) | 60  | 6.34 % |
| Group 3 (genetic, congenital, and metabolic conditions and skeletal dysplasias) | 113  | 11.95 % |
| Group 4 (neurological and neurodevelopmental disorders) | 267  | 28.22 % |
| Group 5 (cancer and organ transplant) | 13  | 1.37 % |
| Group 6 (risk and preterm newborn infants) | 247  | 26.11 % |
the program: 101 (51.24 %) because they turned 18 years old; of them, 38 were in group 4. Among those who were preterm infants, 93 (46.26 %) were discharged at 6 years old without sequelae and 39 were transferred to group 6. A total of 7 (3.48 %) were dropouts or deceased patients.

There are currently 743 active patients in the OPSC program distributed into the 6 groups mentioned above, as shown in Table 4; examples of diseases receiving follow-up are also described (Annex 3 Spanish version).

As seen when comparing the distribution by group at the time of admission and that of active patients, many were transferred from a group of high incidence (group 6) to a group of high prevalence (group 4) due to prematurity-related morbidity, which has a high impact on our population.

Table 4. Distribution of active patients as of January 2019

| Patients | Proportion | Age (mean ± SD) |
|----------|------------|-----------------|
| Total    | 743        | 100 %           | 9.7 years old (± 4.52) |
| Group 1  | 221        | 29.74 %         | 10.85 years old (± 4.16) |
| Group 2  | 60         | 8.08 %          | 9.32 years old (± 4.44) |
| Group 3  | 99         | 13.32 %         | 9.84 years old (± 4.35) |
| Group 4  | 268        | 36.07 %         | 10.52 years old (± 4.22) |
| Group 5  | 10         | 1.35 %          | 10 years old (± 5.44) |
| Group 6  | 85         | 11.44 %         | 4.26 years old (± 1.48) |

SD: standard deviation.

The definition of criteria to assess results was attempted from the beginning. Treatment compliance, variations in the Z-score for weight or height, etc. were not used due to the heterogeneity in terms of disease and population. The positive indicators of this model were its generalized acceptance for adoption, the very high rate of compliance with the national immunization schedule, the lack of significant rejections, and that there were no formal claims about the program quality or compliance in the studied period.

A NEW GENERATION OF PHYSICIANS.
TOWARDS A NEW PARADIGM

The teaching experience at the CeSAC no. 5 is extensive. It hosts graduate students who are taking Pediatrics as part of the Clinical course of OPSC: Outpatient Pediatric Special Care.
the School of Medicine and rotating interns from the Annual Rotating Internship (Universidad de Buenos Aires, UBA), as well as pediatric residents and attending physicians from different national and CABA hospitals, workers’ unions health insurance organizations and private hospitals.\textsuperscript{11,12}

The objective of such rotating internships is to learn, on the field, the integrated concept of the health-disease dyad as a dynamic process conditioned by social, environmental, cultural, and family factors. Emphasis is placed on preventive and health promotion aspects. These guidelines are defined in Section 40 of the Health Law of CABA, Ordinance 40997/86, as amended, the syllabus of Pediatrics from the graduate program, and the reference framework of the pediatric residency program.\textsuperscript{13-16}

Rotating interns participate in patient care as part of the OPSC program, and this allows them to understand the care of high-risk patients in previously unknown situations, thus redefining the role of PHC, not only as a simple form of medicine but also as a relevant, integrated part of the PHS. This includes specific health care experience and strategic thinking in PHC.\textsuperscript{17-19}

The teaching experience with residents and students created awareness of the necessary reconciliation between the theory of prevalent problems and the complex reality of our population.

**CURRENT SITUATION**

**Strengths, weaknesses, opportunities and threats analysis**

The progress of the model was analyzed in 2 assessment stages: 2005 and 2013. This allowed to adjust inclusion criteria, clear databases, and improve the health care approach. The current situation was assessed based on a strengths, weaknesses, opportunities, threats (SWOT) analysis (Table 5).

**DISCUSSION**

**Programs, plans or policies**

In recent decades, several plans and programs have been implemented aimed at solving different health problems. Most of them were imposed from central levels in a vertical and parallel manner, leaving community involvement aside and with a lack of coordination among them.\textsuperscript{20} Organizational fragmentation, with coexistent sub-systems that were not coordinated or integrated, increased costs due to duplications and transactions, which resulted in a varied quality in terms of provision.\textsuperscript{21} This was due to the absence of a rational, planned, integrated, well-executed, patient-centered NHS with community involvement. As an answer to these problems, the concept was modified to the establishment of health networks.\textsuperscript{22}

| **Table 5. Current situation of the Outpatient Pediatric Special Care program. SWOT analysis** |
| --- |
| **INTERNAL analysis** | **EXTERNAL analysis** |
| **Strengths** | **Opportunities** |
| • Human resources with a high level of training. | • Extended coverage and accessibility (null rejections). |
| • Positive attitude towards a challenge with a greatly cohesive staff. | • Adequate social acceptance. |
| • Comprehensive care, transdisciplinary approach. | • Community involvement. |
| • Knowledge of the population. | • Multiplying effect of other measures and policies. |
| • Building resources and adequate equipment. | • Rapid recognition and identification of the target population. |
| • Affordable. | • Likely synergy with integrated health care networks (redes integradas de servicios de salud, RISS). |

| **Weaknesses** | **Threats** |
| --- | --- |
| • Limited coordination with other levels of care. | • Rigidity at other levels of care to accept the new policy and the program. |
| • Lack of projection to other health centers. | • Lack of drive in relation to the health administration. |
| • Limited hours with specially designated staff. | • Difficult relationship with the different care and health management levels due to the program’s local nature. |
| • Administrative limitations for data and statistics entry. | • Reduced availability of hours for care provision due to the administrative tasks inherent to the program. |
In the past decade, the World Health Organization (WHO) has disseminated a new concept in this regard. Turning current systems into integrated health care networks (redes integradas de servicios de salud, RISS), understood as an “integrated network of equal and comprehensive health care services for a specific population willing to account for its clinical and economic outcomes and the health status of the population it serves.” This concept surpasses that of health networks because it focuses on a comprehensive functioning and is based on PHC systems that improve access, equity, quality, and effectiveness to provide the population with favorable responses. A practical example to understand the idea is the “health care model for people with chronic diseases (HCMPCD),” which has established a national strategy to address a new epidemiological reality in a comprehensive manner based on PHC and in coordination with the different levels of care. This constitutes a new paradigm for Argentina, where the health care system works, most of the time, contrary to how a network functions.

Based on this experience, the tendency is believed to go towards the replacement of autonomous programs by actual policies and an effective and rational organization that governs health.

The OPSC program was created in the same line with RISS and, therefore, as in the case of the HCMPCD model, it may be implemented as a policy within the system, boosted by the information tools provided by the new proposal. The OPSC is a health policy. It establishes a targeted health care model based on the risk for mortality and provides a wider framework for the epidemiological surveillance of pediatric health in each geographic unit. The CeSAC functions as the gateway and also as the main coordinator of patient care.

New challenges
Health care dynamics should go along with social and epidemiological dynamics. In this changing reality, resources should be implemented in addition to technological advances. The integration of electronic medical records, telemedicine, teaching manikins, etc. may help to develop a better PHC.

The information should focus on patients and their setting, and on identifying risk groups, generating priorities, establishing alerts and reminders that activate actions among the different disciplines for their interaction, developing recommendations, and promoting communication among the health system actors.

Weaknesses
The lack of an adequate information support and the possibility of keeping it updated hindered the presentation of more detailed statistical analyses. Future assessments are required for their development.

CONCLUSIONS
Recognizing children at risk, identifying them, supervising their care, ensuring universal access and an adequate coordination with other disciplines and the different levels of care seems to be the most direct path towards the reduction of infant mortality, an enhanced health promotion and protection, and, therefore, a more equitable society. The success of this mission lies in an organization with a consistent care and a clear knowledge of the health reality of our children.

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### ANEXO 1

**Recursos humanos disponibles en el Centro de Salud y Acción Comunitaria N.° 5**

| Personal             | Cantidad | Condición       |
|----------------------|----------|-----------------|
| Personal administrativo | 7        |                 |
| Técnica en Farmacia | 1        |                 |
| Enfermería           | 8        |                 |
| Antropóloga          | 1        |                 |
| Farmacéuticas       | 2        |                 |
| Nutricionistas      | 2        |                 |
| Odontología          | 2        |                 |
| Anatomía Patológica | 1        |                 |
| Obstetricia          | 2        | Personal estable|
| Psiquiatría          | 1        |                 |
| Psicología           | 7        |                 |
| Trabajadora social  | 5        |                 |
| Psicopedagoga       | 1        |                 |
| Médicos pediatras   | 13       |                 |
| Médicos clínicos    | 3        |                 |
| Médicos generalistas| 6        |                 |
| Médicos tocoginecólogos | 6     |                 |
| Médicos ecografistas| 2        |                 |
| Residentes de Pediatría | 25     | Rotaciones trimestrales |
| Otros residentes    | 26       | Rotaciones de formato variado |

*Fuente: CeSAC N.° 5 (2014).*
ANEXO 2

HISTORIA CLÍNICA - CONSULTORIO DE ALTO RIESGO       H. C. No.: 

Nombre y apellido: 
Fecha del 1.º control: / / 

Domicilio: 
Fecha de nacimiento: 
Lugar de nacimiento: 

Cobertura médica: 
Nº. de afiliado: 

Pediatra de cabecera: 
Centro: 

Antecedentes perinatológicos: 

Gesta [] Para [] Cesáreas [] Abortos [] 

Embarazo controlado: sí/no 
Desde (fecha): / / 
Cantidad de controles: 

Lugar de control: 

Antecedentes: Hipertensión [] Diabetes [] Tabaco [] Fármacos/tóxicos [] 
(Maternos) Traumatismos [] Cirugías [] Infecciones [] ...................... 

Otros: ......................... 

Antitetánica: [] Cantidad de dosis: ......... Duración del embarazo: sem. 

Serologías del embarazo: 

VDRL ( ) / / 

Toxopl. ( ) / / 

Hepat. B ( ) / / 

HIV ( ) / / 

Chagas ( ) / / 

Otras: ......................... ( ) / / .............................. ( ) / /
PARTO:

Comienzo: espontáneo □ inducido □ Lugar de nacim.: htal. ..................................  
Finalización: vaginal □ forcipal □ cesárea □  Causas: ...........................................

Presentación: cefálica □ podálica □ otras: RN ( ) EG 

Ruptura de membranas _______ Liq. amniótico _________

Peso al nacer: _________ g  Talla al nacer: _________ cm  P. craneal: _________ cm

Apgar: /  Edad gestacional: ......... semanas

Alta:  Caida del cordón:  FEI:

Complicaciones:

Familgrama y antecedentes familiares:

![Familgrama](image_url)

Enfermedades prevalentes

1-

2-

3-

4-

Antecedentes personales patológicos:
INMUNIZACIONES

| Edad         | BCG | Cuádruple | Doble | Sabin/Salk | Triple viral | Hepatitis B | Otras |
|--------------|-----|-----------|-------|------------|--------------|-------------|-------|
| Al nacer     |     |           |       |            |              |             |       |
| 2 meses      |     |           |       |            |              |             |       |
| 4 meses      |     |           |       |            |              |             |       |
| 6 meses      |     |           |       |            |              |             |       |
| 12 meses     |     |           |       |            |              |             |       |
| 18 meses     |     |           |       |            |              |             |       |
| Ingr. escol. |     |           |       |            |              |             |       |
| 16 años      |     |           |       |            |              |             |       |

Alimentación:

**Lactancia exclusiva**: □ Duración: ...........  Lactancia + complem. □ Tipo:.................. Cantidad:..................

**Alimentación artificial**: □ Tipo: .................. Cantidad: .................. Fe □ Dosis ........mg/kg/día Desde:........

**Alimentación complementaria. Edad de inicio:** ..................  Vit. ACD □ Desde: ..................

Tipo de preparaciones: ........................................................................................................................................

Lugar:  Casa □  Guardería □ ..................  Comedor barrial. □ ..................  Caja: □

ANTROPOMETRÍA

| Menores de 1 año | Mayor de 1 año |
|------------------|----------------|
| Fecha | Edad | Peso | PIE | V.g/día | Talla | T/E | P.cef | Fecha | Edad | Peso | PIE | Talla | T/E | Adec. P/T | Vel. P | Vel. T |
|------------------|----------------|
|                  |                |

CONSULTORIO DE ALTO RIESGO

**Criterio de ingreso:**

**Fecha de ingreso al programa:**
### Derivado por el siguiente:

| PATOLOGÍA RESPIRATORIA: | SÍ - NO |
|-------------------------|---------|
| **Diagnóstico:**       | □ SDRt  |
|                         | □ EMH   |
|                         | □ NMN   |
|                         | □ SALAM |
|                         | □ HPPRN|
| **Otros:**             |         |
| **Tratamiento:**       | □ Halo: días |
|                         | □ CPAP: días |
|                         | □ ARM: Máx. MAP: días |
|                         | □ Maduración pulmonar: |
|                         | □ Surfactante: dosis: |
| **Complicaciones:**    | □ NMTx |
|                         | □ Est. laringea |
|                         | □ Atelectasia |
|                         | □ Otros: |
| **Displasia broncopulmonar:** |         |
| **Tratamiento:**       | □ O₂: |
| **Fármacos:**          |         |
| **Apneas:**            | □ Centrales |
|                         | □ Obstructivas |
|                         | □ Mixtas □ Secundarias |
| **Comentarios:**       |         |

### HIPERBILIRRUBINEMIA: SÍ – NO

| Incompatibilidad: |         |
|                   | □ Tipo: |
|                   | □ ABO  |
|                   | □ Rh   |
|                   | □ Subgrupo |
| **Bilirrubina:** | Total/indirecta |
| **Máxima:**      | mg     |
| **Horas de vida:** |         |
| **Luminoterapia:** | □ Días: |
| **Exanguinotransfusión:** | □ Cantidad: |
| **Comentarios:** |         |

### TRASTORNOS HEMATOLÓGICOS: SÍ – NO

| Policitemia: |         |
| Hto. máx.:  |         |
| Hemodilución: | □ Cantidad: |
| **Otras patologías:** | □ CID |
| Plaquetopenia | □ Trombosis |
| Hemorragias | Otras: |
| **Tratamientos:** |         |
| **ANEMIA:** | □ | Hto./Hb. mínimo: | Edad: | Transfusiones: □ | Cant.: |
|-----------------------------|-----|----------------|-------|----------------|-------|
| Hto/Hb. alta:              |     |                |       |                |       |

| **EPO:** | □ | Semanas: |
|-----------------|-----|---------|

| **PATOLOGÍA CARDIOVASCULAR:** | SÍ – NO |
|-------------------------------|---------|

| **Trastornos:** | □ Shock | □ Bradicardias | □ Taquiarritmias | □ Hipertensión | □ Insuf. cardíaca |
|-----------------|---------|----------------|-----------------|----------------|-------------------|
| Otros:          |         |                |                 |                |                   |

| **Fármacos:** | □ Indometacina | □ Drogas vasoactivas | □ Diuréticos | □ Indom./ibuprofeno |
|----------------|----------------|----------------------|-------------|---------------------|

| **Cuáles:** |
|-------------|

| **Complic. transit.:** | Ductus □ | Cierre: espont. - farmacológico – quirúrgico |
|------------------------|---------|-------------------------------------|

| **HPPRN:** | □ |
|-------------|---|

| **Tipo de tratamiento:** | Halo – ARM – ARF AF – NO inhal. – Surfact. |
|--------------------------|-------------------------------------------|

| **Cardiopatía congénita:** | □ Cuál: |
|---------------------------|---------|

| **Ecocardiografías:** | / / . Resultado: |
|-----------------------|------------------|
|                       | / / . Resultado: |
|                       | / / . Resultado: |

| **INFECCIONES:** | SÍ – NO |
|------------------|---------|

| **Tipo:** | □ Connatal | □ Intranasocomial | Cuál: |
|-----------|------------|-------------------|-------|

| **TORCHS:** | □ Toxo. – | □ Rubéola – | □ Chagas – | □ Sífilis – | □ HIV | □ |
|-------------|-----------|------------|-----------|------------|------|---|

| Sospechada/confirmdada |
|-------------------------|

| **Sepsis:** | □ | Temprana – Tardía |
|--------------|---|-------------------|

| **Germen:** | |
|-------------|---|

| **Tratamiento:** |
|------------------|

| **Comentarios:** |
|------------------|

| **ALTERACIONES DIGESTIVAS:** | SÍ-NO |
|-----------------------------|-------|
### ALTERACIONES METABÓLICAS: SÍ – NO

- [ ] Hipoglucemia
- [ ] Hiperglucemia
- [ ] Hiponatremia
- [ ] Hipernatremia
- [ ] Hipocalcemia
- [ ] Hipokalemia
- [ ] Hiperkalemia
- [ ] Acidosis metabólica
- [ ] Hiperamoniemia
- [ ] Otras:

**Comentarios:**

### ALTERACIONES NEUROLÓGICAS: SÍ – NO

- [ ] Trast. succión degluc.
- [ ] Hipotonía
- [ ] Hipertonia
- [ ] Convulsiones
- [ ] Parálisis
- [ ] Paresias
- [ ] Otras:

**Hidrocefalia:**

- [ ] DVP

**HIC:**

- [ ] Grado
- [ ] Leucomalacia periventric.

**Medicación:**

**Ecogr. cerebral:**

- [ ] / / . Resultado:

- [ ] / / . Resultado:

- [ ] / / . Resultado:

- [ ] / / . Resultado:

**Ex. neurol. a las 40 semanas:**

- Normal
- Dudos
- Anormal

**Comentario:**
### ALTERACIONES AUDITIVAS: SÍ-NO

**Evaluación auditiva prealta:** sí-no  **Método:** Fecha:  **Resultado:**

- **Fact. de riesgo:**
  - [ ] Familiar sordo – [ ] TORCHS – [ ] Malf. craneofaciales – [ ] Menos de 1500 g – [ ] Hiperbilirrubinemia c/exanguinotransfusión –
  - [ ] Fcos. ototóxicos – [ ] Meningitis bacteriana – [ ] Apgar 1° entre 0 y 4 – [ ] Apgar 5° entre 0 y 6
  - [ ] ARM más de 5 días –
  - [ ] Síndromes genéticos c/hipoacusia

**OEA**  fecha:  **Resultado:**

**PEAT**  fecha  **Resultado:**

**Comentarios:**

### ALTERACIONES VISUALES: SÍ-NO

| Malformaciones | Cuál: | Cataratas: | Ref. |
|----------------|-------|------------|------|
| rojo: | |

**Ojo izq. ROP:**  Grado  Plus:  Láser  Coment.: |

**Ojo der. ROP:**  Grado  Plus:  Láser  Coment.: |

**Controles:**
- 1.°  /  /  . Result.: |
- 2.°  /  /  . Result.: |
- 3.°  /  /  . Result.: |
- 4.°  /  /  . Result.: |
- 5.°  /  /  . Result.: |

**Fecha del próximo control:**  /  /  .

**Alta oftalmológica:**  /  /  .

**Comentarios:**
DATOS DEL EGRESO HOSPITALARIO

Fecha en que cumple 40 semanas:  /  /  .

| Fecha de egreso: | Edad cronológica: | Edad corregida: |
|------------------|-------------------|-----------------|
| Al alta: Peso:   | g ( )             |                 |
| cm ( )           | Talla: cm ( )     | Per. craneal:   |
| cm ( )           |                   |                 |

Alimentación al alta: Pecho ☐ Mixta ☐ Artificial ☐ Cuál: ml/kg/día

Medicación al alta:

1) dosis cada h
2) dosis cada h
3) dosis cada h
4) dosis cada h
5) dosis cada h

Otros tratamientos:

Laboratorio al alta: Hto: Hb: Retic: Na+: K+:

Calcio: Fósforo: FAL: Got: Gpt: Prot.

tot.: Alb: Otr: Otros:

Diagnósticos al egreso: 1)
2)
3)
4)
| Estudios pendientes: |
|---------------------|
| I)                  |
| II)                 |
| III)                |

| Consideraciones iniciales del seguimiento y plan de estudios: |
|-------------------------------------------------------------|
| -                                                            |
| -                                                            |
| -                                                            |
| -                                                            |
| -                                                            |
| -                                                            |

Fecha: / / .  Firma y sello
EVALUACIÓN

Nombre y apellido: H. C.: 

FECHA / / Edad cronológica: Edad corregida: 

- PESO: Pc ( ) - TALLA Pc ( ) P. C. Pc ( ) 

Estado de las vacunas: Desarrollo neuromadurativo: 

Alimentación: Pecho [ ] Mixta [ ] Fórmula [ ][ ] Tipo: 

Cantidad: ml. Cada: h. Total: ml/kg/día. Semisólidos Sólidos 

Suplementos: [ ] Fe: [ ] Fólico: [ ] ACD: [ ] Vit. D: [ ] Calcio: [ ] Fósforo: [ ] Otros: 

¿Recibió información preventiva para la edad (accidentes, estimulación, etc.)? [ ] 

Examen físico (datos +): 

Diagnóstico: 

Trae resultados de 

Solicito lo siguiente para la próxima visita: 

- 
- 
- 

Tratamiento e indicaciones: 

1) 
2) 
3) 
4) 
5) 
6) 

Citado el día / / . 

Firma y sello
ANEXO 3

**Ejemplos de patologías atendidas en Cuidados Especiales Ambulatorios Pediátricos**

- **Grupo 1**: Bronquitis obstructiva recurrente, enfermedad pulmonar obstructiva crónica posviral, bronquiectasias, abceso pulmonar, malformación adenoidea quística, tetralogía de Fallot, comunicación interventricular, ductus arterioso persistente, comunicación interauricular, miocardiopatía dilatada, retinitis pigmentaria, cataratas congénita, estrabismo grave, ambliopía, artritis reumatoidea juvenil, espondilitis anquilosante, focomelia, enfermedad celiaca, refluo gastroesofágico grave, hipotiroidismo congénito y autoinmune, vitíligo, fiebre reumática, síndrome urémico hemolítico, síndrome nefrítico, síndrome nefrótico, refluo vesicouretal, agenesia renal, ureterocele.

- **Grupo 2**: Onfalocele, higroma quístico, fistula traqueoesofágica, fisura labio-alvéolo-palatina, atresia duodenal, membrana yeyunal, gastosquisis, atresia de vías biliares, síndrome de Alagille, cavernoma de la vena porta, enfermedad de Hirschprung.

- **Grupo 3**: Down, Klinefelter, Stickler, Williams, Sotos, neurofibromatosis 1, Prader-Willi, fragilidad del cromosoma X, Poland, Sturge-Weber, asociación VATER, alcoholismo fetal, osteogénesis imperfecta, displasia cefalocraneal, discrepancia de miembros, craneosinostosis (Apert, Pfeiffer, trigonocefaalia), mielomeningocele, lipoma de raquis, Dandy-Walker, artrogríasisis, diabetes tipo I y II, hipercolesterolemia familiar.

- **Grupo 4**: Hipoacusia neurosensorial, encefalopatía crónica no evolutiva, displéjia espástica, epilepsia, hidrocefalia, distrofia de Duchenne, retraso madurativo, retraso mental, trastornos de aprendizaje, trastorno generalizado del desarrollo, parálisis cerebral.

- **Grupo 5**: Leucemia linfoblástica aguda, retinoblastoma, linfoma, craneofaringioma, tumor de órbita, tumor de tálamo.

- **Grupo 6**: Prematuroz extrema, bajo peso y muy bajo peso al nacer, infecciones congénitas (sífilis, toxoplasmosis, citomegalovirus, chagas, HIV, rubéola), retinopatía del prematuro.