

# Women collectors of recyclables: resistance strategies and medicines during the COVID-19 pandemic<sup>1</sup>

## Catadoras de materiais recicláveis: estratégias de resistência e medicamentos durante a pandemia de covid-19

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### Abstract

The collection of recyclable materials is an occupation with an increasing number of workers, who find in it a livelihood in a context of socioeconomic inequalities and unemployment. This article describes resistance strategies of women collectors who are part of an association in Northeast Brazil, in addition to describing how they managed health problems during the COVID-19 pandemic, especially regarding medicines. A quantitative and an ethnographic approach were used by collecting data on sociodemographic and medicines present in the homes of 13 participants and conducting participant observation and in-depth interviews. Amid the health crisis, the participants mentioned problems such as unemployment, gender relations, violence, and work stigmas. If, on the one hand, the pandemic has further increased vulnerability for certain population segments, on the other hand, it has enhanced collective resistance strategies. The collectors of the association organized to obtain various materials and benefits both for their headquarters and their neighborhood. We identified 58 units of medicines, prescribed and obtained mainly at the Basic Health Unit. Of these, 27.6% had action on the nervous system (analgesics, psycholeptics, and psychoanaleptics) and 17.2% on the cardiovascular system and on the alimentary tract and metabolism. The collectors developed self-attention practices with the medicines highlighting their central role in facing health problems.

**Keywords:** Solid Waste Collectors; COVID-19; Resistance Strategies; Medicines; Health self-attention.

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## Resumo

A catação de materiais recicláveis é uma ocupação com crescente número de trabalhadores, que encontram nela sustento diante das desigualdades socioeconômicas e do desemprego. Este artigo descreve as estratégias de resistência de mulheres catadoras que fazem parte de uma associação no Nordeste brasileiro, além da forma como elas lidaram com problemas de saúde durante a pandemia de covid-19, especialmente no que se refere aos medicamentos. Utilizamos uma abordagem quantitativa e uma etnográfica, coletando dados sociodemográficos de 13 participantes e dos medicamentos presentes em suas casas, e também realizando observação participante e entrevistas aprofundadas. Em meio à crise sanitária, foram mencionados problemas como o desemprego, as relações de gênero, a violência e os estigmas do trabalho. Se por um lado a pandemia vulnerabilizou ainda mais alguns segmentos populacionais, por outro potencializou estratégias coletivas de enfrentamento. As catadoras e a associação se organizaram para obter insumos e melhorias tanto na sede quanto no bairro. Identificamos 58 unidades de medicamentos, prescritos e obtidos principalmente na Unidade Básica de Saúde. Desses, 27,6% tinham ação sobre o sistema nervoso (analgésicos, psicodélicos e psicoanalépticos) e 17,2% sobre o sistema cardiovascular e sobre o trato alimentar e metabolismo. As catadoras desenvolveram práticas de autoatenção com os medicamentos, destacando o papel central deles no enfrentamento dos problemas de saúde.

**Palavras-chave:** Catadores; Covid-19; Estratégias de Resistência; Medicamentos; Autoatenção à Saúde.

## Introduction

In Brazil, collecting recyclable materials became a source of income in large urban centers from the 1950s onwards. The production of waste from the growing Brazilian industrial hub meant that many unemployed people saw waste as a way of earning a living (Rocha; Francischett, 2021).

*“I collect paper. I’m proving how I live!”* is a phrase from the autobiography of Carolina Maria de Jesus (2014, p. 20), a Black woman, a *favela* resident, and collector in São Paulo, who portrays the hard work of these people in terrible socioeconomic conditions, dealing with violence, stigma, and need to work in scavenging.

Excluded populations develop survival strategies to obtain their sustenance, with garbage collection as one of these possibilities. By collecting, separating, and transporting recyclable materials in dumping grounds, landfills, cooperatives, streets, and vacant lots, they begin to occupy new workspaces and build an essential source of income in times of high unemployment rates (Medeiros; Macêdo, 2006; Cruz, 2020; Rocha, 2020).

Worldwide, it is estimated that around 15 million people work in scavenging (Coelho et al., 2018). This activity has been recognized since 2002 in Brazil, appearing in the Brazilian Classification of Occupations as “Workers collecting and selecting recyclable material” (MTE, 2017). According to a 2012 study, 400,000 to 600,000 people worked collecting recyclables in the country at the time, 68.9% men and 31.1% women, with an average age of 39.4 years. Other relevant data from this study are that 66.1% were Black, 38.6% had some contractual employment relationship, and 93.3% lived in urban areas (Ipea, 2013). Furthermore, it is estimated that 75% of professionals who work in solid waste sorting cooperatives are women, mostly Black, excluded from the formal job market, which tends to absorb male labor (Cherfem, 2016; Coelho et al., 2018). It is worth mentioning that female collectors have a lower average income than male in the same role (Ipea, 2013).

Most of the time, direct work with solid waste occurs in places without structures. Their very disposal can compromise and degrade water resources and soil, triggering diverse and complex

problems, affecting physical health, producing psychological and psychiatric disorders and social disintegration of those who work with scavenging (Coelho et al., 2016, 2018; Filipak et al., 2020). Thus, health problems such as infectious, degenerative, and cardiovascular diseases, anxiety and depression crises, panic syndrome, chemical dependency, and increased violence, among others, are components of the same phenomenon (Siqueira; Moraes, 2009; Santos; Silva, 2011). Furthermore, there are the occupational effects of scavenging, mainly problems related to work accidents (cuts, punctures, burns, etc.), with a high incidence of food poisoning and parasitic diseases (Siqueira; Moraes, 2009).

The year 2020 is marked in world history as the beginning of the SARS-CoV-2/COVID-19 virus pandemic, which generated deserted streets, chaos in public health, and negative consequences for the economy and public policies. Due to the suspension of several economic activities during the lockdown period, there was an impact on the sustainability of many companies and services, which led to the dismissal of many workers, in addition to a decrease in income, an increase in hunger, and the reconfiguration of work in throughout Latin America (Malaver-Fonseca; Serrano Cárdenas; Castro Silva, 2021). For socially vulnerable populations, the consequences were even more significant. Faced with this new reality, many of the unemployed saw collecting recyclable materials as an alternative, given the economic and social difficulties faced during the period. Many groups, collectives, non-governmental organizations (NGOs), and social movements sought mutual support strategies in the first two years of the pandemic. The Juventude na Luta collective, which emerged during the health restrictions due to COVID-19, was made up of young residents of Bons Ares (a peripheral region of one capital in the Brazilian Northeast), who built social actions together with waste collectors from the Bons Ares Waste Collectors Association (ASCABOA). In this article, we describe the local context where they live, their resistance strategies, and how they dealt with their health problems during the pandemic, with a focus on medicines. Research related to waste collectors, such as Galon (2015) and Coelho et al. (2018), investigated these workers' healthcare and how social,

work, and environmental issues influence their life contexts. However, few studies from the perspective of the investigation proposed here, which seeks to understand the use and perceptions of medications from subjects organized in an association.

## Methodology

### Data collection and analysis

The field research occurred in the context of ASCABOA, located in Grande Bons Ares (GBA), in a northeastern capital, from May to October 2021. Female workers affiliated with the Association, aged between 18 and 59, were included. (we excluded women aged 60 or over as they are considered a risk group for COVID-19) Furthermore, their source of income was only collecting recyclable materials (other sources cited were working as a cleaner and a delivery person). During the research, this scenario changed, with men working exclusively at the Association due to unemployment, but the inclusion criterion above remained.

We opted for a mixed method, using a quantitative approach mixed with a qualitative ethnographic one to understand the use and perceptions of medicines by ASCABOA collectors.

For the first approach, the participants' sociodemographic data were collected (number of residents in the house, date of birth, affiliation, education, race/color, religion, and household characteristics), and the medications found in their homes were identified (name, dosage, pharmaceutical form, who prescribed it, where they were purchased, among others). Medication data were organized in an Excel® spreadsheet and analyzed according to frequency and percentage. The ATC/WHO (Anatomical Therapeutic Chemical/World Health Organization International Working Group for Drug Statistics Methodology) classification system was used, with each medicine identified according to its drug, whose ATC/WHO code (fifth level) was obtained from the National List of Essential Medicines (Rename) from 2022 and only then categorized by Anatomical/Pharmacological Group (first ATC/WHO level) and by Therapeutic/Pharmacological Group (second ATC/WHO level).

For the other methodological approach, we chose the ethnographic method, as it allowed us to understand what meanings were attributed to the condition of waste collectors and what impact this work had on their lives and their relationships with medicines. In participant observation, the field diary was used. In-depth interviews were also carried out on a day, time, and place defined by the participants and with a script to facilitate the dialogues (the average time was 60 minutes). The interviews were always recorded when authorized, with subsequent transcription in full.

The research complied with CNS Resolution No. 510/2016, receiving a favorable opinion for its execution by the Human Research Ethics Committee. The names of the locations, institutions, and collectors have been changed to ensure anonymity. All participants were informed about the research and signed the Informed Consent Form.

### **The study location**

The GBA is a region comprising five neighborhoods, and the municipality's official data records a population of approximately 225,000 inhabitants in 2021-2022, representing 8.33% of the capital's population. The residents who formed the Juventude na Luta collective and the associates reported that the Association's location was the "*vixe do vixe*" within the GBA, that is, an even more violent space than the rest of the region, already known for the interjection of amazement and fear "*vixe*," which was pronounced by people when they heard someone say they lived there.

As in many more populous Brazilian capitals and cities, the GBA has its origins in the region's disorderly growth, building a social stigma around its inhabitants and producing situations of significant vulnerability and accumulation of social problems, such as violence and structural problems (precarious sanitation, pavement, and lighting conditions, presence of garbage in the streets, among others). It is also worth mentioning the spatial division due to conflicting interests within the locality itself, which is reflected in the movement of residents from one community to another, impacting differentiated access to public facilities, such as healthcare units.

The collectors mentioned a Basic Health Unit (UBS) and two private pharmacies close to ASCABOA (about 1 km), the leading health equipment.

## **Results and discussion**

### **Women collectors, stigmas, and resistance strategies**

During 2021, there was an increase in the number of members: from 24 (20 women and four men) to 36 members (30 women and six men). Thirteen women, aged between 23 and 59, participated in the study, six of whom were Black, five were mixed race, one was Indigenous, and one was White. Just over half were married (seven), and the rest were divided between single (five) and divorced (one). The number of children was between one and five, with two saying they did not have children. The level of education varied between not reading/not writing (two), incomplete (four) and complete (two) primary education, incomplete (three) and complete (two) secondary education. Most professed the Christian faith (five evangelicals and four Catholics), one identified herself as an Umbanda follower, and three did not want to declare their religion. All 13 residences had electricity and running water. During the research period, six houses received access to the public sewage network, achieved via sanitation works from the Association's articulation with other social movements; the others had septic tanks (five) or open sewers (two). Most (nine) of the collectors lived in their own homes, coming from urban occupations regulated by the state; three were in a borrowed house, and one was renting.

Amid the health crisis caused by the pandemic, participants pointed to issues and problems that had a direct impact on their lives, such as unemployment, gender relations, local violence, and the stigmas of scavenging work.

In family relationships, the central role of these women was evident, being a point of convergence in their stories. Maintaining the family's subsistence and raising their children and grandchildren were their shared needs. These women, especially those who lived in a single-parent situation, emphasized the relevance of scavenging in their lives, even more so in times of unemployment and the absence of a father figure:

*Because I was unemployed. I had two children to support, and I like to depend on God first and foremost. And I raise this little girl there, who is my granddaughter, and at the same time, she has no father, no mother; she is an orphan. Everybody knows it. So, instead of going to houses asking for money to feed them, I took a bag from the recycling bin. It was when Alessandra started here at the Association. I started in a bag and ended in a car. (Iara)*

Vanessa spoke of her troubled relationship with her husband, at times citing issues related to psychological violence when he belittled her situation as an unemployed woman. She portrayed with great hurt the fact that her husband required earning and, at the same time, boycotted her when collecting:

*Even now, he keeps saying, "Ah, you're working." I had a cart without a wheel, and he didn't want to help me fix it. We had another good cart, and he sold it, leaving that crap for me. It's much smaller, and I must bend over to adjust it. I'm looking forward to returning to formal work to get away from this man. (Vanessa)*

All participants, except one, stated that they had worked as domestic workers at some point in their lives. Iara reported that scavenging was a salvation amid unemployment caused by the insecurity of her job as a domestic worker, which she left when she had psychological problems after her son's death.

Conflicts between "parallel powers" (an expression used by the community) were intense during the research period, generating a territorial division that prevented free movement between neighborhoods, which limited both the scavenging work in different regions and participation in the Association. In some reports, the impacts of urban violence were mentioned by women who lived in places outside the perimeter of the ASCABOA headquarters, speaking primarily of the difficulty in accessing health services and medicines:

*I buy these two at the pharmacy [referring to captopril and omeprazole]. Because I believe that the healthcare unit has blood pressure medication, but like us, many people from where I live don't have access to it at the healthcare unit. Then, getting treatment and looking for a physician to improve our*

*health becomes more complicated. There is no way. So, we have to do everything possible and impossible because scheduling an appointment takes time, and it also doesn't work because we go and are blocked, we go, and something might happen [referring to the limits imposed by the "parallel powers"] So I prefer not to go. (Sara)*

The fact that they work with recyclable materials and live in a place historically known as violent and with structural problems highlighted social stigmas:

*He said, "bunch of garbage men messing around with the trash." So I said, "No sir, we're not going through the trash, we're wearing gloves, we're wearing masks, we're just the way we are supposed to be. When I get home, I wash my hands, too; I wash everything properly." Then he threatened to call the child protection council because I was with the boy. Then, this pandemic started, and I had no one to leave him with. There was no daycare; there was nothing. (Fernanda)*

Some women spoke about the expectation that scavenging would open doors and be another source of income, becoming a new possibility for the reinvention of the poorest segments and social emancipation (Velloso, 2005):

*And then, I started to see the work of the collectors as something different, something I hadn't seen before. I started to see that for many people, what is garbage becomes money for us. On the street, it's trash; in the Association, it's money. And I started to see it like this: I said, "No, I'm going to recycle." In a way, we help the environment because we are removing from the environment all those materials that will take years to degrade, and then we sell them and still earn. (Renata)*

*Then I saw how important it was because people said, "Ah, this can give you extra income." Then I stopped to think, "Yes, really." I had no income, and today, I'm setting up a salon for myself; I'll already have two incomes. (Meire)*

The organization, via the Association, was a central point in the union and articulation of

these women's resistance. Cooperative/associated collectors have more possibilities to negotiate their sales and guarantee continuity of supply, adequate physical space, and better working conditions (Grimberg; Blauth, 1998). In addition to improving productivity and participation in social movements and forums, their selective collection service consolidates their participation within institutions (Siqueira; Moraes, 2009).

ASCABOA was organized in 2012, bringing together collectors from different neighborhoods in the GBA. In 2015, with the support of some institutions and in conjunction with community movements in the region, a place was provided that served as a warehouse and headquarters. Even though it only had a warehouse of around 24 m<sup>2</sup> and only one bathroom, this improved the quality of the work, valuing the construction of trade networks for recyclable materials in the region itself. However, workers did not have manual transport vehicles, making it impossible to carry materials over long distances. At first, there were just six collectors, as Alessandra reported. This number increased over the years after the construction of the first headquarters and intensified during the COVID-19 pandemic, as mentioned above. As highlighted, the consequences of the pandemic were reflected in several sectors beyond public health, directly impacting less favored populations and increasing unemployment and hunger numbers (Malaver-Fonseca; Serrano Cárdenas; Castro Silva, 2021). However, the political organization of these women during the pandemic period was fundamental for the construction of collective strategies, including the physical change of the headquarters. The territorial dispute between the "parallel powers" led to the end of the concession of the site provided by the Residents Association, and ASCABOA was transferred to a region with better access for the majority of members. Despite the rental cost, the property brought advantages such as a larger plot of land, a terrace where materials were separated and weighed, an administrative room (consisting of a bathroom, a living room, and a kitchen), and a shed where a machine press was installed. In addition,

two houses were provided to female collectors who did not have their residences.

Throughout the study, the importance of organizing these women around the Association was perceived. Alessandra was the oldest collector among the participants and was present from the first political articulations for the cooperative's emergence. Renata, in turn, joined in 2021. It was clear that the feeling of belonging that emerged builds a bond of identity and resistance in the face of adversities in the local context.

Women occupied a prominent position in ASCABOA, coordinating the organizational aspects, political articulations, the construction of support networks, and the mobilization of members. The few associated men performed manual duties, repairing equipment and maintaining infrastructure.

Operationally, the Association was organized by distributing the material collected and the items received through donations. Sorting and weighing were carried out at headquarters by the collectors themselves, with part of this process being the responsibility of those who had young children and could not leave them alone at home.

During the pandemic, ASCABOA joined other waste collector associations in articulating the achievement of *Auxílio Catador*<sup>2</sup> and basic sanitation in the neighborhoods where the members lived. It also collaborated with other social movements, such as the Juventude na Luta collective mentioned above, aiming to participate in projects and obtain basic food baskets, personal hygiene materials, personal protective equipment (PPE), and health education about COVID-19, among others.

### **"So the easiest thing I think is to self-medicate": home pharmacies and self-attention practices**

A total of 58 medicines were observed in 12 of the 13 residences, with a minimum of one in one home and a maximum of 11 in another. Regarding the pharmaceutical form, 63.8% (n=37) were tablets, 15.5% (n=9) were liquids for oral use, 12.1% (n=7) were capsules, 3.4% (n=2) were dragees, 3.4% (n=2) were ointments, and 1.7% (n=1) were ophthalmic solutions.

<sup>2</sup> The *Auxílio Catador* Program, approved in 2020 by state law, transferred a financial amount directly to collectors through a personalized bank card, seeking, in return for this support, an increase in activities related to reuse, recycling, and treatment of waste solids.

Storage took place in the bedroom (four homes), kitchen (three homes), and living room (three homes). In one of them, the medicines were in the bathroom, and one of the participants reported that she carried them in the bag she used daily. For seven medications, it was impossible to identify the expiration date, and two expired. It is worth mentioning that in the house with the largest number of medications, one was expired, and six had no expiration date.

When asked who recommended the medicines, 60.3% (n=35) of the items were prescribed by the neighborhood UBS physician (28 prescriptions) or by a psychiatrist (seven prescriptions). Indications made by pharmacists were reported for 19% (n=11) of the medications, by UBS nurses for 8.6% (n=5), and for information found on Google, 5.2% (n=3). The other medications were recommended by family members (sister recommended it once, mother recommended it twice, and the collector herself self-medicated in one case).

The majority of medications were obtained in public health establishments: 48.3% (n=28) at the UBS and 6.9% (n=4) at the Psychosocial Care

Center (CAPS). One of the private pharmacies in the neighborhood played an essential role in this acquisition, having been observed in 37.9% (n=22) of the items, which demonstrates that even though a physician from UBS or CAPS prescribed the majority of these medicines, some of them were obtained in the private sector. The remaining medications were received through donations (n=2), as a free sample provided by the physician (n=1), or even by the mother (n=1).

In the home pharmacies, a majority percentage of medicines related, according to the first ATC/WHO level (Anatomical/Pharmacological Group), to the nervous system (n=16; 27.6%), followed by drugs acting on the cardiovascular system (n=10; 17.2%), in the alimentary tract and metabolism (n=10; 17.2%), in the blood and blood-forming organs and anti-infectives for systemic use (for each group n=4; 6.9%), among others (Table 1). The medicines identified as “not included” (n=4; 6.9%) were items that did not have an ATC/WHO code, as they were combinations of drugs such as the compound Dipyrone + Orphenadrine + Caffeine.

**Table 1 – Anatomical Therapeutic Chemical/World Health Organization International Working Group for Drug Statistics Methodology (ATC/WHO) classification of medicines found in the homes of women collectors of recyclables, Grande Bons Ares, 2021**

ATC/WHO FIRST LEVEL	n	%	ATC/WHO SECOND LEVEL	n	%
A Alimentary tract and metabolism	10	17.2	A02 Drugs for acid related disorders	3	5.2
			A03 Drugs for functional gastrointestinal disorders	2	3.4
			A06 Drugs for constipation	1	1.7
			A10 Drugs used in Diabetes	2	3.4
			A11 Vitamins	1	1.7
			A12 Mineral supplements	1	1.7
B Blood and blood-forming organs	4	6.9	B01 Antithrombotic agents	2	3.4
			B03 Antianemic preparations	2	3.4
C Cardiovascular system	10	17.2	C03 Diuretic	2	3.4
			C07 Beta blocking agents	2	3.4
			C09 Agents acting on the Renin-Angiotensin system	4	6.9
			C10 Lipid modifying agents	2	3.4
D Dermatologicals	1	1.7	Do6 Antibiotics and Chemotherapeutics for dermatological use	1	1.7

continues...

**Table 1 – Continuation.**

ATC/WHO FIRST LEVEL	n	%	ATC/WHO SECOND LEVEL	n	%
H Systemic hormonal preparations, excl. sex hormones and insulins	2	3.4	Ho2 Corticosteroids for systemic use	1	1.7
			Ho3 Thyroid therapy	1	1.7
J Antiinfectives for systemic use	4	6.9	Jo1 Antibacterials for systemic use	4	6.9
M Musculo-Skeletal system	2	3.4	Mo1 Antiinflammatory and Antirheumatic products	2	3.4
			No2 Analgesics	8	13.8
			No3 Antiepileptics	1	1.7
			No5 Psycholeptics	4	6.9
N Nervous system	16	27.6	No6 Psychoanaleptics	3	5.2
			Ro3 Drugs for obstructive airway diseases	1	1.7
			Ro6 Antihistamines for systemic use	2	3.4
S Sensory organs	2	3.4	So1 Ophthalmologicals	2	3.4
Not included	4	6.9	Not included	4	6.9
TOTAL	58	100.0		58	100.0

The most frequently identified therapeutic/ pharmacological groups (ATC/WHO, second level) were: analgesics (n=8; 13.8%), agents acting on the renin-angiotensin system (n=4; 6.9%), antibacterials for systemic use (n=4; 6.9%), psycholeptics (n=4; 6.9%), and psychoanaleptics (n=3; 5.2%), as shown in Table 1. As stated previously, four (6.9%) medicines were not classified because they were not on the ATC/WHO list.

The drugs most frequently present in residences were the analgesics dipyron and paracetamol, with four units of each (6.9%), followed by omeprazole (treatment of gastric or duodenal ulcers) and losartan (antihypertensive), both with three units (5.2%). One or two units represented the frequency of other medications. It is essential to highlight medications with psychoactive action, classified in the nervous system together with analgesics: antipsychotics (haloperidol; lithium; risperidone), antidepressants (venlafaxine; escitalopram; nortriptyline), anxiolytics (diazepam), and antiepileptics (clonazepam).

It is worth mentioning that we did not identify medicines from the so-called “early treatment” of COVID-19 (or “COVID kit”) in the residences, such

as ivermectin, chloroquine, and hydroxychloroquine, nor were they mentioned by the participants, which is a paradox compared to what was observed in the country (Melo et al., 2021; Hentschke-Lopes et al., 2022).

Collectors of recyclable materials reported varied situations that can be understood under self-attention, both in its broad and strict sense (Menéndez, 2005). In the broadest sense, they involve “[...] all forms of self-attention used to ensure the biosocial reproduction of subjects and groups at the level of microgroups and, especially, of the domestic group” (Menéndez, 2005, p. 55). Thus, the strategies for coping with the pandemic (described in the previous topic), going to the *forró* to drink beer and dance, and the children’s day party promoted by the Association, among other activities, are part of this self-attention perspective. In the strict sense, it also occurs within the scope of the family and social groups, i.e., it is based on the collective. It is characterized by intentionality in the health-disease-care process without the direct intervention of specialists, whether from biomedicine or other forms of health care. In the processes of illness, subjects seek solutions with

health-restoring-oriented praxis. This search does not imply excluding or privileging a form of health care but often occurs in the articulation between different practices. Furthermore, according to Menéndez (2005), praxis is centered on the subject and their social group, which synthesizes, articulate, mix, or juxtapose the different forms of attention, reconstituting and organizing part of them in self-attention activities.

Biomedicine considers self-attention only to be self-medication with medicines. However, for Menéndez (2005), it is only part of it: self-medication is using medicines and all other substances, such as herbs, alcohol, marijuana, etc., as well as activities of many types, such as cupping, massages, and poultices. According to him, it is given by the intention of using any substance, treatment, or carrying out activities that, according to its users, enable better performance at work, in sport, in sexual life, etc.

Among the women collectors, self-attention practices involved using medicines and other preparations. Luzia was a reference in the neighborhood and for the Association's collectors in recommending medicinal plants and teas. Other preparations mentioned were home-made syrups and home-bottled medicinal preparations bought from the "sisters" of the church they attended. These solutions were not only practices based on African religions or Indigenous medicine, as Luzia explained in one of the conversations, but they also constituted alternatives to the difficulty in accessing health services.

The use of medications, whether through self-medication or prescription, was reported in these women's routines. There was a common practice, for example, of using anti-inflammatories for pain caused by work. Vitória described that for these purposes, she used teas recommended by Luzia, paracetamol, and Dorflex®, which were recommended by other associates: *"I drink this [tea] to relieve back pain. There are days when we carry much weight, and as I am in the screening area, sometimes it is too much. The girls give me paracetamol and Dorflex when it doesn't go away. Most always have it in their pocket"* (Vitória).

Meire spoke about her health condition, relating it to the medicines she used and medical diagnoses.

She presented herself as a very sick woman: *"I can't live without my medicine. I'm all messed up, you know?"*. These problems portrayed by Meire were high blood pressure, fatigue attacks (related to shortness of breath), varicose veins, and profound anemia. All of them had, at some point in her life, a medical diagnosis, but her understanding of treatments went beyond the physician-patient relationship.

Since going to the UBS with severe headaches and a blood pressure reading of around 16/10 mmHg, when the physician diagnosed her with high blood pressure, Meire understood herself as having hypertension. Another physician from the same UBS recommended non-pharmacological treatment practices and removed the antihypertensive medication. However, the collector contradicted the medical discharge and other recommendations, questioning the contradictory conduct and the relationship with the healthcare team. When asked about the captopril present in her home pharmacy, she said:

*I took captopril at the healthcare unit. My captopril. But when the physician removed it, he said I didn't need it, but I knew I needed it. Because a physician already said that I needed continuous use, and another physician arrived there, the physician even left. Everything for him was "You're going to go for a walk because you're morbidly obese" and no medicine. That's it; he took away all the medicines I had to use, at least in my case. Then I self-medicate. Am I in need? I'm not going to the unit. Because to go and ask for medicine, you have to go to a GP, and without saying that, it is difficult to get a consultation, so the easiest thing, I think, is to self-medicate. (Meire)*

According to Conrad (1985), individuals develop strategies based on perceptions about diagnosis and treatment, adapting them to their realities. The author called this mechanism self-regulation, which can involve adjusting doses or abandoning treatment, for example, by not complying with medical instructions.

The change of physician at the healthcare unit and the possible guidance on doing physical exercises to take care of excess weight were not effective from a biomedical point of view. Meire

created ways to deal with her hypertensive condition, measuring her blood pressure, understanding the characteristic symptoms when her blood pressure was altered, and seeking advice from family members with experience of hypertension:

*Because I've been to the healthcare unit several times with a terrible headache, and I got my blood pressure measured, and my blood pressure was 18/8, you know? Then I bought my pressure measuring device, and it's always fluctuating. My normal blood pressure is 10/6, 10/7, normal. Then, when it reaches 13, they say it's normal, but I already feel a headache as my blood pressure is always low. Then I take my medicine, you know? I take my captopril and take medicine for my headache.[...]. (Meire)*

*My mother recommended that I take AAS [acetylsalicylic acid] too. Sometimes, when I have a headache, I take AAS, too. In addition to thinning the blood, it prevents stroke. That's what my mother always tells me: a headache and high blood pressure are enough to cause a stroke. She already had it. AAS is for me to take every day, but there are days when I don't remember, and I take it when I remember. (Meire)*

Like Meire, four other collectors, aged between 28 and 59, were diagnosed with high blood pressure, giving different accounts of the origin of the problem. In the same local context, paradoxically, the physician-patient relationship had other implications for the practice of self-attention by these women. Alessandra and Patrícia lived around 20 m from the UBS. The other two collectors who lived with hypertension were Sara and Fernanda. Both, like Meire, had in common that they lived close to each other and further away from the UBS.

The lesser or greater proximity to the healthcare unit reflected in the way these women dealt with their hypertension. Alessandra and Patrícia reported no problems acquiring the medicines. They stated that they strictly followed the instructions given

for use, praising the good service and how well they were treated at the UBS.

On the other hand, Sara, Fernanda, and Meire discussed different situations involving compliance with treatment. In these cases, difficulty accessing medicines led them to construct other narratives. Fernanda, talking about the use of captopril: *"I prefer to buy it right away at the pharmacy. There are many things that are not available at the healthcare unit, and I buy it right there"* (Fernanda). Sara built her relationship with the medication based on her neighborhood and the way her fellow members of the Association took their medication for high blood pressure:

*I use captopril because of my blood pressure. When I have a headache, I feel nauseous; then I know that my blood pressure is high, so I take two tablets. [...] I don't take it every day. Because I don't want to be dependent on medicine. So when I see that the blood pressure is high, I take it. (Sara)*

Iara reported having a chronic problem, which she called "phobia," which she described as sadness, difficulty sleeping, agitation, fear of leaving the house, rapid heartbeat, and, sometimes, even fainting. She was cared for by CAPS and received all her medication through the Unified Health System (SUS). According to her, the physician prescribed diazepam 10 mg to help her sleep, venlafaxine 75 mg, lithium carbonate<sup>3</sup> 300 mg to *"stop feeling sad"* (Iara), and liquid haloperidol to stop *"jumping like a cricket"* (Iara) due to her agitation and fear of leaving home. At the beginning of the conversation, her report of stronger crises seemed to link her problems to the death of her teenage son, as mentioned above. However, when asked how the "phobia" started, she said that it was not caused by the event above but instead due to sexual violence suffered in her childhood, building a process of chronic illness that affected her mental health:

*It wasn't with my boy's death. I already had a phobia, and it wasn't because of my boy's death. Yes, I already had it before I had a child, but I didn't*

<sup>3</sup> While lithium is classified by ATC/WHO and Rename as an antipsychotic when associated with an antidepressant, it receives another code from ATC/WHO. The lithium leaflet indicates its use together with an antidepressant in the case of severe recurrent depression.

*know what a phobia was; I knew what fear was. I went to other places; I didn't feel well; I got to certain places and sat down. Once, I went to work, arrived at the terminal, and crouched down [I squatted] so I wouldn't fall in the middle of the terminal. Then I asked for a hand from one person and another to help me get up because I was squatting. Then people thought [I was asking for something]: "No, I don't have, sorry." I wasn't asking for money! I was asking for a hand. I couldn't go home alone. I called my daughter; I couldn't even hold the cell phone. I really started very young. It wasn't work, but many things happen in people's lives that sometimes people keep quiet about to avoid feeling ashamed. At that time, it was embarrassing. Not anymore today. The person being tempted, grabbed, tried to rape me twice. How old was I? Eleven to twelve years old.* (Iara)

This suffering, which lasted for more than 40 years, had a history of mental illness that began with physical violence, which impacted her life, causing her to develop a stigma for having chronic psychiatric problems:

*In the beginning, what I talked to him about, I was embarrassed to say that I was undergoing treatment at CAPS. I don't like talking about life; we don't have to talk about our problems that no one will solve for each other. Then, over time, I started saying: "I'm going for a consultation tomorrow." Then he: "Consultation for what?" "No, I'm going to CAPS." "Do what at CAPS?" "No, I do a follow-up." "That's it, you know? But there's that fear, like, people are afraid of that person; they say the person is a bit of a fool.* (Iara)

Iara also expressed a relationship of dependence on the psychotropic drugs she used: *"They are the source that calms me down. By taking them, I don't go around jumping around like cricket, I can sleep, I don't feel a strange presence watching me, and I can even leave the house. When I'm in crisis, I can't go out to collect"* (Iara).

Renata narrated her story of mental illness, attributing the factor that triggered a panic attack to the COVID-19 pandemic, leading her to seek psychiatric help:

*When it was 2020, the pandemic started, and I had COVID, and then I became a little psychologically ill. I panicked because I thought I was going to die from the disease, and so on, and I started experiencing bouts of shortness of breath and everything. But everything was always very correct. Every day I was at the healthcare unit, I checked the blood pressure and the oxygen, and I was fine, you know? But psychologically, I was very sick; I woke up at dawn as if I were dying from lack of air.* (Renata)

For Renata, Rivotril® (clonazepam<sup>4</sup>) and Lexapro® (escitalopram) improved her sleep, describing them as her *"buckets of happiness."* The life story of these two women highlights the psychoactive medicines' central role that has been addressed in other studies (Silveira, 2000; Whyte; Van Der Geest; Hardon, 2002; Maluf; Tornquist, 2010), revealing a clear relationship between gender, mental health, and suffering.

## Final considerations

The growth in the number of recyclable material collectors follows the increase in unemployment rates, which causes these socially excluded populations to develop survival strategies to obtain their livelihood and to deal with their illnesses, especially during the COVID-19 pandemic. This study identified that the gender perspective was very relevant in the Association, which mainly comprised female workers responsible for administration and organization and acted politically.

The literature records the collection of materials as a work activity with a risk of physical and mental illness, subjecting workers to unhealthy situations and exposure to harmful materials. However, the socioeconomic conditions they face are also factors that influence their health.

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<sup>4</sup> While clonazepam is considered an antiepileptic/anticonvulsant in the ATC/WHO and Rename classification, it has indications for treating anxiety and mood disorders in adults, which appears to be the case with Renata.

Identifying where each participant lived, observing their routines, describing their contexts and perceptions about health and illness, and listening to their stories allowed us to understand that stigmatization due to the territory where they live and their work and gender relations impact their illnesses and treatments.

As they lived in the capital region with the worst human development indices, being part of a territory with conflicts between “parallel powers,” building a more significant link with public healthcare services was difficult. Reports of illnesses linked to gender-based violence were identified in the research, as well as difficulties in maintaining a single-parent family.

The relationship between these women and medicines showed that self-attention through self-medication has become an essential strategy for dealing with adverse situations present in their daily lives. Even though physicians prescribed most of the medicines found in the homes, each of the participants manipulated their use according to their knowledge and previous experiences. The use of psychoactive medicines stood out, showing a process of medicalization of life and psychological and emotional suffering.

To face many difficulties, active participation in the Association began to provide more possibilities for improving working and housing conditions, in addition to building a bond of identity and resistance in the face of adversities in the local context, directly reverberating in improvements in living conditions observed during the COVID-19 pandemic.

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### Authors' contributions

Coutinho Júnior contributed to the conception, data collection and analysis, and article review and writing. Diehl contributed to the conception, data analysis, article review, and writing.

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