Changes of Estuarine Sedimentation Patterns by Urban Expansion: The Case of Middle Capibaribe Estuary, Northeastern Brazil

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Abstract

The aim of this study is to describe the sedimentary evolution occurred during the last 200-years in the middle Capibaribe Estuary by mean of the sedimentary analysis (magnetic susceptibility, grain size, calcium carbonate, total organic matter—TOM) and geochemical parameters (sedimentation rates, heavy metal concentrations, enrichment and contamination factor) along a core. The core recorded four units and the measured sedimentation rate was 0.52 cm cm⁻¹ y⁻¹. The first unit, dating before 1812, showed environmental characteristics of mangrove with predominance of fine sediments, high total organic matter percentages and heavy metal concentrations probably from natural sources. The second unit, from 1812 to 1937, showed a slight increase in sand percentages and decrease in fine fraction, TOM contents and heavy metals concentrations. These characteristics may be associated with the urban expansion processes and the presence of monoculture of sugar cane occurred in the middle Capibaribe Estuary. The third unit, from 1937 to 2004, showed the highest sand percentages of the core, characterizing a unit exclusively of sand with low fine fractions percentages, total organic matter contents and heavy metals concentrations. This unit represented the intensification of the urban expansion processes of Recife City. The fourth unit showed increases in fine fraction sedimentation, TOM contents and heavy metals concentrations. This new change in sedimentation probably is consequence of rebirth of marginal estuarine banks by mangrove vegetation, due to environmental projects carried out by Recife Prefecture in the early 2000’s. It was not possible to register the anthropic contamination to middle estuary area probably due to the Barreiras Formation influences in the metal concentration records, masking the anthropic contamination inputs in estuarine region. Although, lead and arsenic showing an enrichment level indicating anthropic contamination.
1. Introduction

The estuaries are transition areas between continental and ocean interface and they are important environments in comprehension of the coastal areas development and prevision of future evolving tendencies [1]. The sediments deposits are important because it could preserve the history of sedimentation in coastal environments [2]. Estuarine sedimentation is a consequence of many conditions, such as the sediment source that may be from the river or marine origin [3] [4] [5]. The estuarine regions show a rich fauna and flora and these are important sources of natural resources, and around these estuarine areas located are, in general, in larger cities and industrial complexes [1].

Historically, the coastal areas are attractive to human settlements, and these form huge metropolises, that grew in disorderly way [6]. The urban and industrial development adjacent to estuarine areas may cause environmental imbalance, changing the water column chemical proprieties, sedimentation patterns, mangrove deforestation and typical forest areas, landfill of flood areas to urban expansion, building of dams, contamination of heavy metals and others [7]. The studies with sedimentary columns are used to comprehension the environmental evolution in coastal areas. These sedimentary cores may provide historical records of sedimentation patterns in coastal systems, indicating the natural baselines and changes caused by anthropic modifications during the time [8] [9].

Some parameters are employed to understand the environmental evolution such as sedimentological parameters as grain size, organic matter and calcium carbonate contents in coastal environments, that provide important information for paleoenvironmental reconstructions and may register the historical of the local and regional climatic changes [10]. Variations in magnetic susceptibility have been used for detecting anthropogenic pollution caused by power plants, metallurgical dusts, fly-ashes and urban airborne particulates [11] [12] [13] and heavy metal concentration is employing to identify natural or anthropic sources and contamination levels [14]-[19]. Finally, the use of radionuclides in coastal environments as sediment tracers offers considerable potential for determining sediment sources and sedimentation rates in drainage basins [20].

The sedimentary characteristics in cores may be associated with the development of the urban processes. Anthropic changes around the river, estuary or beach are responsible for a natural imbalance of the coastal systems as well as the interference in transport of sediments and organism cycles [1]. The anthropic induced changes occur in Capibaribe Estuary (8°S/35°W) since the start of colonization processes as such as coastline modification, dredging and landfill activities [21]. Currently, this estuary is being polluted by different sources as well as fish and shrimp farming areas, discharge of industrial and domestic wastes and

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**Keywords**

Sedimentary Evolution, Grain Size, Heavy Metals, Sedimentation Rate
In this way, the aim of this study is to describe the sedimentary evolution occurred in the middle Capibaribe Estuary by means of the sedimentary analysis and geochemical parameters along a sedimentary core, associating with the anthropic change occurred in the estuarine system and infer the background metal values.

2. Material and Methods

2.1. Study Area

The Capibaribe Estuary is located in Recife City (8°S/35°W), Pernambuco State (Northeastern Brazil). It is formed by the Capibaribe, Tejipió/Jiquiá, Jordão, Pina and Beberibe rivers. The Capibaribe Estuary has an important ecological function, serving as a reproduction area for mammals, birds, fishes and other organisms and also is the economic and social pole for the Recife Metropolitan Region (RMR) [22] (Figure 1).

The Capibaribe River present 250 km of extension, from its source in Jararacá mountain chain to its mouth, draining 43 municipalities [24] [25], with a drai-
nage basin of approximately 7,700 km$^2$. The river flow on crystalline rocks, and the tertiary and quaternary sediments being drained only in the Capibaribe River lower sector. Tertiary sediments are composed by fluvial sandy deposits, intercalated with muddy sediments. Quaternary sediments form the coastal plain, corresponding to estuarine and shallow marine sediments; fluvial sediments are presents along river margins [26]. The estuary is shallow (<10 m depth).

The climate of the RMR is characterized as humid and warm tropical, As’ type according to Köppen classification, with higher rainfall periods between March and August and dry periods between September and February. The average rainfall is of 2200 mm∙y$^{-1}$ and air annual temperature average of 25.2˚C [26].

The RMR houses 3.7 millions of inhabitants, being approximately 1.5 million in Recife City and, from this number, about 218,000 inhabitants living in the middle Capibaribe Estuary. Recife City has a density population of about 7000 inhabitants/km$^2$ [27]. The surrounding margins of the Capibaribe River present small and degraded mangroves and the Pina Basin houses an extensive mangrove area, with approximately 20 km$^2$, called Mangrove Park with registered *Rizophora mangle*, *Laguncularia racemosa*, *Avicennia germinans* e *Avicennia schaueriana* specimens. Despite its environmental importance, the Mangrove Park is being affected by real estate pressure, pollution and disorderly occupation [23]. This collection of characteristics define the goal of the present research, study the estuarine sedimentary evolution in a densely urbanized tropical area.

According to [24] and [28] the grain size of surface sediments varies from silt to gravelly sand, with a prevalence of fine sediments and sands for all Capibaribe Estuarine System. However, in the middle estuary the authors observed a little change in type of sedimentation with fine fraction predominance. Concerning temporal variations, there is a slight tendency of increasing clays during periods of higher rainfall [28].

### 2.2. Methods

A 178-cm deep core was collected in 27th November 2012, with the aid of a push-core operated by a scuba diving at the coordinates of 8˚02’22.56”S and 34˚55’27.12”W, 0.5 m of water column in a marginal bank. After the core recovery, it was analyzed for magnetic susceptibility in a Bartington MS2C meter. The measurements were made with a 0.1 × 10$^{-5}$ S.I. resolution and readings were taken at 2 cm intervals [13]. After that, he core was opened, described and continuously sub-sampled at intervals of 2 cm. Samples were dried at 50˚C in order to be analyzed for grain size, organic matter and calcium carbonate contents, metal concentrations and determination of sedimentation rates by $^{210}$Pb$\alpha$ activity.

Total calcium carbonate and organic matter were determined according the methods proposed by [29]. The calcium carbonate was determined by weight difference of the sediment prior and after addition of hydrochloric acid (10%). The total organic matter was determined also by weight difference prior and after addition of hydrogen peroxide (10%), until total reaction. Both results are presented as percentages of the bulk sample.
Grain size was determined in a Malvern Mastersizer 2000 laser sedimentometer, after calcium carbonate and organic matter removal of samples. Results allowed the determination of the grain size statistical parameters proposed by [30] and sediment classification by [31], by mean of the Sysgran 3.0 software [32].

The sedimentation rate result followed radionuclide $^{210}\text{Pb}$ determination method described by Saito et al. (2001), that is based in $^{210}\text{Pb}$ half-live decay ($t_{1/2} = 22.3$ years) [34]. The sedimentation rate value was calculated by CIC model (Constant Initial Concentration) [35]. A maximum time span of 200 years was considered for the age determinations [36].

The metal concentrations were obtained by energy dispersive X-ray fluorescence (EDXRF). The method is a multi-elemental analysis that is based in exciting of chemical element atoms of samples; the values of element concentrations were calculated according dispersion, detection and mensuration these X-rays [37]. For metal analysis in sedimentary samples were used proximally 1 g of sediment in polyethylene tubes and were sealed with polyethylene film for EDXRF analysis. The results were measured in equipment ED-720 (Shimadzu) with rhodium X-ray tube and Si (Li) detector for characteristic element quantifications. For methodology certification was utilization two reference samples of National Institute of Standard and Technology (NIST) and of International Energy Atomic Agency (IAEA), SRM2709 San Joaquin Soil and IAEA-Soil7 respectively (Table 1). These soil reference were analyzed together the environment sediments and at time of 300 seconds. Cu, Co, Fe, Ga, La, Mn, Ni, Pb, Rb, Sr, Ti, V and Zn elements were used Mo filter and voltage adjusted at 50 kV and Al, Ca, K, and Si elements were measured with Al filter and voltage of 15 kV. The values are obtained and certified for the chemical elements and its respective expanded analytical uncertainties (95% of confidence) for energy dispersive X-ray fluorescence (EDXRF).

To identify the anomalies and evolution of the heavy concentrations, geochemistry standardizations by conservative element were used as tool, to evaluate the level of the anthropogenic contamination [38] [39]. There are two approaches to normalization of metals in sediment, the granulometric and geochemical methods [40]. The geochemical normalization is necessary standardized with alithogenic conservative element [41] [42]. Usually, Aluminum is used as lithogenic conservative element because it is a major constituent of fine-grained alumiosilicates [40] and its concentration is generally not influenced by anthropogenic sources [41] [43].

There are some types of geochemical normalization such as Al-normalized, enrichment factor, geochemical index and contamination factor. In this study will be performed enrichment factor (EF) and contamination factor (CF), and the indexes are based in the trace element concentration in base core, the Earth’s background or the metal analysis reference value.

According to [44], the enrichment factor (EF) is defined as follows Equation (1):

$$EF = \left( \frac{C}{Al} \right)_{sample} / \left( \frac{C}{Al} \right)_{background}$$ (1)
Table 1. Information Value (I.V) and Confidence Interval (C.I) for reference samples SRM 2709 and IAEA Soil 7 (mg·kg⁻¹).

| Element | SRM 2709 | IAEA Soil 7 |
|---------|----------|-------------|
|         | I.V      | C.I         | I.V  | C.I          |
| Al      | 2.6*     | 2.0 - 3.1*  | 47,000 | 44,000 - 5100 |
| As      | <20      | -           | 13.4  | 12.5 - 14.5  |
| Ba      | 398      | 392 - 400   | 159   | 131 - 196    |
| Br      | -        | -           | 7     | 3 - 10       |
| Ca      | 1.5*     | 1.4 - 1.7*  | 163,000 | 157,000 - 174,000 |
| Co      | 12       | 10 - 15     | 1.3   | 8.4 - 10.1   |
| Cu      | 32       | 26 - 40     | 11    | 9 - 13       |
| Fe      | 3.0*     | 2.5 - 3.3   | 25,700 | 25,200 - 26,300 |
| Ga      | -        | -           | 10    | 9 - 13       |
| K       | 0.32*    | 0.26 - 0.37 | 12,100 | 11,300 - 12,700 |
| La      | -        | -           | 28    | 27 - 29      |
| Mg      | 1.4*     | 1.2 - 1.5*  | 11,300 | 11,000 - 11,800 |
| Mn      | 470      | 360 - 600   | 631   | 604 - 650    |
| Ni      | 78       | 65 - 90     | 26    | 21 - 37      |
| P       | 0.07*    | 0.05 - 0.07*| 460   | 460 - 462    |
| Pb      | 13       | 12 - 18     | 60    | 55 - 71      |
| Rb      | -        | -           | 51    | 47 - 56      |
| Si      | <0.01*   | -           | 180,000 | 169,000 - 201,000 |
| Sr      | 101      | 100 - 112   | 108   | 103 - 114    |
| Ti      | 0.038*   | 0.03 - 0.04*| 3000  | 2600 - 3700  |
| V       | 62       | 51 - 70     | 66    | 59 - 73      |
| Zn      | 100      | 87 - 120    | 104   | 101 - 113    |

*concentration values in percentages

where \( C_{\text{sample}} \) is the trace element concentration in the sample, \( C_{\text{background}} \) is the trace element concentration in base core, \( A_{\text{sample}} \) is the aluminum content in the sample, and \( A_{\text{background}} \) is the aluminum content in base core. The EF values were interpreted as the levels of metal pollution as suggested by [45] where EF < 1 indicates no enrichment, <3 is minor, 3 - 5 is moderate, 5 - 10 is moderately severe, 10 - 25 is severe, 25 - 50 is very severe and >50 is extremely severe.

The contamination level in sediments by metal is expressed in terms of a contamination factor (CF). According [46] theses values are calculated as follows Equation (2):

\[
CF = \frac{C_{\text{metal}}}{C_{\text{background}}}
\]

where, \( C_{\text{metal}} \) is metal concentration in sample and \( C_{\text{background}} \) is the Earth's background or reference value of the metal analysis. The CF value is divided into four classes: CF < 1 refers to low contamination; 1 ≤ CF < 3 means moderate contamination; 3 ≤ CF ≤ 6 indicates considerable contamination and CF > 6 indicates very high contamination [47].

3. Results

The sedimentation rate for the core was 0.52 cm·y⁻¹, and according to the sedi-
mentary and geochemistry characteristics were recorded four units. The first unit (106 - 178 cm) refers to period before at 1812, this unit shows mean magnetic susceptibility values of $22.5 \times 10^{-5}$ SI, grain size varied from fine silt to very fine sand with predominance of coarse silt, low sand percentages (average 43.6%) and high silt and clay percentages (averages of 47.0% and 9.4%, respectively). The calcium carbonate percentages and TOM contents showed higher percentages with average of 11.8% and 8.5%, respectively, although the samples are siliciclastic [48]. The Mn, Fe, Co, Ni, Cu and Ga elements show low concentrations and Zn, Pb, Ti, V, Mg and Al high contents, when compared with background values of [49] (Table 2).

The Unit 2 (42 - 106 cm), refers to the period from 1812 to 1937, shows average MS values are $23.0 \times 10^{-5}$ SI. This interval is represented by sandy characteristics, grain size varying from medium silt to very fine sand with very fine sand predominance. This unit registered a slightly increase in sand percentages (average of 61.6%) and light decrease in silt and clay percentages (averages of 31.7% and 6.8%, respectively) when compared with the Unit 1. The calcium carbonate percentages and TOM contents show averages of 10.7% and 8.7%, respectively.

**Table 2.** Average concentrations of parameters analyzed to core of middle Capibaribe Estuary. SM: Susceptibility Magnetic; CaCO$_3$: Calcium Carbonate; TOM: Total Organic Matter.

| Element | Unit 1 | Unit 2 | Unit 3 | Unit 4 | Turekian & Wedepohl (1961) [49] |
|---------|--------|--------|--------|--------|-------------------------------|
| MS**    | 22.5 ± 4.6 | 23.0 ± 3.0 | 28.5 ± 3.6 | 12.5 ± 2.5 | - |
| CaCO$_3$* | 11.8 ± 3.7 | 10.7 ± 2.4 | 6.9 ± 2.9 | 9.9 ± 2.1 | - |
| TOM*   | 8.5 ± 3.2 | 8.7 ± 2.4 | 6.6 ± 2.9 | 7.1 ± 0.6 | - |
| Sand* | 43.6 ± 15.6 | 61.6 ± 10.4 | 86.5 ± 6.8 | 48.1 ± 3.8 | - |
| Silt* | 47.0 ± 12.5 | 31.7 ± 8.4 | 11.4 ± 5.8 | 45.4 ± 4.2 | - |
| Clay* | 9.4 ± 3.7 | 6.8 ± 2.4 | 2.1 ± 1.0 | 6.5 ± 0.3 | - |
| Mn | 438.4 ± 100.3 | 349.4 ± 35.8 | 320.7 ± 50.9 | 313.7 ± 18.0 | 850 |
| Fe* | 3.9 ± 0.7 | 3.8 ± 0.5 | 2.8 ± 0.6 | 3.1 ± 0.4 | 4.7 |
| Co | 15.8 ± 2.9 | 16.7 ± 2.8 | 11.7 ± 2.4 | 13.1 ± 2.3 | 19 |
| Ni | 28.9 ± 3.6 | 23.1 ± 2.4 | 19.6 ± 2.6 | 277.2 ± 260.1 | 68 |
| Cu | 50.8 ± 9.1 | 29.9 ± 14.2 | 22.9 ± 9.8 | 36.8 ± 22.0 | 45 |
| Zn | 157.0 ± 33.0 | 156.9 ± 27.2 | 116.7 ± 28.6 | 138.7 ± 22.2 | 95 |
| Ga | 10.9 ± 1.0 | 10.8 ± 0.6 | 9.5 ± 0.7 | 9.7 ± 0.4 | 19 |
| As | 288.4 ± 74.4 | 224.6 ± 33.4 | 118.8 ± 44.5 | 120.6 ± 15.3 | 13 |
| Pb | 90.1 ± 18.1 | 74.5 ± 9.0 | 49.2 ± 10.7 | 48.9 ± 2.3 | 20 |
| Ti | 4685.0 ± 398.7 | 4770.7 ± 305.6 | 4264.3 ± 326.2 | 4257.4 ± 176.3 | 4600 |
| V | 290.4 ± 24.7 | 298.1 ± 24.0 | 267.4 ± 29.3 | 255.0 ± 21.6 | 130 |
| Mg* | 0.7 ± 1.1 | 0.6 ± 0.1 | 0.5 ± 0.1 | 0.5 ± 0.1 | 1.5 |
| Al* | 10.8 ± 1.5 | 11.1 ± 0.6 | 8.9 ± 1.3 | 8.8 ± 0.1 | 8.0 |

*values in percentage; other elements in mg·kg$^{-1}$; **$10^{-5}$ S.I.
The Unit 2 showed the same patterns in heavy metal concentration of the Unit 1, when compared with background values of [49] (see Table 2).

The Unit 3 (4 - 42 cm), refers to the period from 1937 to 2004, shows average MS values of $28.5 \times 10^{-5}$ SI. This interval presents sandy characteristics, however is slightly different from Unit 2, mainly due to the mean diameter. This unit showed grain size varying from very fine sand to medium sand with predominance of fine sand. Sand, silt and clay percentages records averages of 86.5%, 11.4% and 2.1%, respectively. The calcium carbonate percentages and TOM contents show averages of 6.9% and 6.6% respectively. The heavy metal concentration showed lower levels, when compared with Units 1 and 2. However, the Zinc, Arsenic and Aluminum concentrations show indexes above the background values from [49] (see Table 2).

Lastly, the Unit 4 (0 - 4 cm), refers to the period from 2004 to 2012, showing the lowest averages of MS values of units with $12.5 \times 10^{-5}$ SI. This unit presents a silty character, with grain size classified as coarse silt. This unit shows sand, silt and clay percentages averages of 48%, 45% and 6%, respectively. The calcium carbonate percentages and TOM contents record averages of 10% and 7%, respectively. This unit show a lightly decrease in heavy metal concentration when compared with Unit 1, showing concentrations below of [49] (see Table 2).

The vertical variation records MS, TOM, calcium carbonate, sand, silt and clay percentage and heavy metal concentration peaks along of core are represented by the Figure 2 and Figure 3.

The enrichment factor (EF) [44] shows no enrichment values to Mn, Fe, Co, Ni, Cu, Ga, Ti, Mg with EF < 1. Zn, Pb and V show values of lower enrichment with EF < 3, and As shows a moderate to moderately severe enrichment factor values, with higher enrichment ($5 \leq EF < 25$) (Figure 4).

The contamination factor [46] values show lower contamination to Mn, Fe, Co, Cu, Ga with CF < 1. Ni, Zn, Ti, V and Al show moderately contamination with $1 \leq CF \leq 3$. The Pb and As show considerable contamination ($3 \leq CF \leq 6$) and higher one (CF > 6), respectively (Figure 5).

4. Discussion

The sedimentary characteristics in cores may be associated with the development of the urban processes. Anthropic changes around the river, estuary or beach are responsible for a natural imbalance of the coastal systems as well as the interference in transport of sediments and organism cycles [1]. The units registered in the core demonstrate influences of different events in sedimentation patterns to middle Capibaribe Estuary.

The Unit 1 recorded the finer sedimentation, TOM and metals, with some peaks of sandy sedimentation at 130, 134 and 142 cm of the core-deep. This unit shows probably the estuarine mangrove environment characteristics, with variation in the hydrodynamic energy represented by the sand peaks. Other evidence that corresponds the mangrove environment is associated with high TOM contents (from 2.1% to 15.6%) [50] [51]. Calcium carbonate peaks registered in this
Figure 2. Vertical variation [31] facies diagram classification, mean diameter, magnetic susceptibility, calcium carbonate and total organic matter contents.
Figure 3. Vertical variation of metal concentrations in sedimentary core.
**Figure 4.** Vertical variation of enrichment factor (EF) of the sedimentary core.

**Figure 5.** Vertical variation of contaminant factor (CF) of the sedimentary core.
unit may be associated with high productivity of organisms with carbonate shells present in mangroves [52] [53]. Probably, the metal concentration values are from natural sources, associated with the local/regional inputs from the Barreiras Formation [54] [55] [56] [57].

According to [58], the Poço da Panela, the neighborhood area located in the middle estuary, was a small village founded in 1758 and its accesses to village were made in tracks inside the Atlantic forest. The urban development in the Poço da Panela started 1812, with open of streets and hills. In this time, river bath was the medical recommendation to cure of diseases, and the Poço da Panela was chosen as a local exclusive for high society of Recife.

Surrounding the middle Capibaribe Estuary, there were big farms of sugar cane plantation as Casa Forte and Madalena mills. The monoculture of sugar cane was the main economy product of the colonial Pernambuco province [59] [60]. Because of medical recommendation, the urbanization processes of Poço da Panela was intensified by the building of summer houses at margins of Capibaribe River, changing urban space in this estuary area as registered in the Unit 2.

The increasing in sand percentages and decreasing in silt and clay contents is associated in Unit 2 probably are related with mangrove deforestation specifically in the margins of Capibaribe River caused by the urban expansion of Recife City westward. Furthermore, mangrove deforestation associated to the sugar cane mills installations also may have been the greater contributor in the increase of sand percentage. The monoculture of cane sugar was responsible by to deforest the majority part of Atlantic Forest located around of Capibarbe estuary, facilitating the erosion and increasing the sediment transport from continental and margins to the river causing silting processes [61] [62]. The urban processes intensification occurred during 19th and 20th centuries, mainly with arrival of Englishman and German, that were employees of the utility companies and preferred to live in this area [58] [59] (Figure 6).

*Figure 6.* Poço da Panela Village in 1847. The Capibaribe River margins were without mangrove vegetation (Unknown Author).
In this unit (Unit 2) shows a decrease in averages of metal concentrations when compared with Unit 1 (see Table 2). Probably this is associated with greater the sand percentages in sedimentation and decrease in the clay percentages [54] [63]. [64] affirms that clays adsorbs four times more organic matter and metals from sands, and two times more from silts. [56] also affirmed that increases in the metal concentrations are associated with high depositions of finer sediments or inputs in the metal concentrations of natural and/or anthropic sources. How Recife City shows a small industrialization processes, this peaks observed in Unit 2 are possibly inputs of natural material from Barreiras Sedimentary formation present around of all estuarine area [21] [57].

The Unit 3 is essentially sandy and probably records the intensification of urban development of middle Capibaribe Estuary in mid-19th century. The increase of the buildings together with the population growth were responsible by the cause of the most changes in around the middle estuary, such as landfill of flood areas, the intensification of mangrove deforestation and canalization of some rivers that compound the hydrographic basin of Capibaribe River estuarine portion [21] [58]. During the 19th century, the urbanization turns its back to the river, and thus, the Capibaribe River passed to be the city’s dumping area, receiving all industries and domestics wastes [65] [66]. During this period the Recife City did not show a development of industrial plants. Therefore, the decrease in metal concentrations recorded in Unit 3 may be associated the greater sedimentation of sand (averages of 86.5%) and decrease in silt and clay percentages and TOM (11.4%, 2.1% and 6.6%, respectively).

Other important fact is that the Recife’s prefecture made a reforestation project of mangrove vegetation in Capibaribe River margins at the mid-90s and in the early 20th century. In this period was registered an increase in mangrove vegetation of approximately 14.2 km² (analyzing the period from 1999 to 2006) and 12.6 km² (analyzing period from 1997 to 2007) according [67] [68] to middle Capibaribe Estuary, respectively.

This project of Recife’s prefecture showed great results as observed by [67] [68], being observed in sedimentary core of middle estuary represented by Unit 4. This unit showed increases in silt and clay percentages (averages of 45.4% and 6.5%, respectively) when compared with unit 3, and a layer highly sandy (averages of 11.4% and 2.1%, respectively). In Unit 4, the increases in fine sedimentation are associated possibly with progradation of the mangrove areas located in margins of Capibaribe Estuary as well as the TOM contents increasing and metal concentrations (see Figure 2).

According to the graphic of enrichment factor (EF) and contamination factor (CF), the majority of metals do not show high levels of enrichment and contamination (Figure 5 and Figure 6). Only Pb and As showed enrichment and contamination levels. According [38] [45] [69], rich rocks in heavy metals in its composition may mask the enrichment and contamination level. In the case of Capibaribe Estuary, the Barreiras Formation is a greater contributor of metal source [57].
[70] [71] [72] affirmed that the strong correlations between metals with Fe and/or Al possibly represent same metal source. In case of middle estuary, the metal concentrations to estuarine system may be from Barreiras Formation such as main source. The anthropic sources also may be registered in this metal concentrations, however this anthropic contribution do not show clearly along of sedimentary core. The correlation values between the metals demonstrated this same metal source for middle Capibaribe Estuary, confirming the Barreiras Formation influences in sedimentation (Table 3).

[73] affirm that soils and rocks of Pernambuco State are composed by high concentrations of Arsenic in their composition. The strong correlations between As and Al ($r > 0.7$) indicate increases of natural metal concentrations, derived of erosion or lixiviation of Tertiary rocks, of the Barreiras Formation, that compounds the low lands of the drainage system.

Correlations between metals and calcium carbonate implies that it could be related to the origin of carbonate that it diagenetic. These formations occur in zones of shallow sub-bottom layers, presence of metabolizable organic matter, oxide and hydroxide (principally iron and manganese), oxygen and sulfate ions concentrations dissolved in pore water [74] [75] associated to the bacterial action [76]. [16] [77] [78] affirm that the carbonate diagenetic formation may oc-

|       | MS | CaCO$_3$ | TOM | Sand | Silt | Clay | Mn | Fe | Co | Ni | Cu | Zn | Ga | As | Pb | Ti | V | Mg | Al |
|-------|----|----------|-----|------|------|------|----|----|----|----|----|----|----|----|----|----|----|----|----|
| SM    | 1.0|          |     |      |      |      |    |    |    |    |    |    |    |    |    |    |    |    |    |
| CaCO$_3$ | -0.5| 1.0      |     |      |      |      |    |    |    |    |    |    |    |    |    |    |    |    |    |
| TOM   | -0.1| 0.3      | 1.0 |      |      |      |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Sand  | 0.4 | -0.5     | -0.3| 1.0 |      |      |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Silt  | -0.4| 0.5      | 0.3 | -1.0| 1.0 |      |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Clay  | -0.4| 0.5      | 0.2 | -0.9| 0.9 | 1.0 |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Mn    | 0.1 | 0.1      | 0.5 | -0.4| 0.4 | 0.3 | 1.0|    |    |    |    |    |    |    |    |    |    |    |    |
| Fe    | -0.5| 0.6      | 0.5 | -0.6| 0.6 | 0.6 | 0.4| 1.0|    |    |    |    |    |    |    |    |    |    |    |
| Co    | -0.4| 0.6      | 0.4 | -0.6| 0.5 | 0.6 | 0.3| 0.9| 1.0|    |    |    |    |    |    |    |    |    |    |
| Ni    | -0.3| 0.1      | 0.0 | -0.1| 0.2 | 0.0 | 0.0| 0.1| 1.0|    |    |    |    |    |    |    |    |    |    |
| Cu    | -0.1| 0.1      | 0.1 | -0.3| 0.4 | 0.2 | 0.3| 0.1| 0.0| 0.2| 1.0|    |    |    |    |    |    |    |    |
| Zn    | -0.6| 0.6      | 0.4 | -0.6| 0.6 | 0.6 | 0.2| 0.9| 0.9| 0.0| 0.1| 0.9| 1.0|    |    |    |    |    |    |
| As    | -0.5| 0.6      | 0.4 | -0.7| 0.7 | 0.7 | 0.5| 0.9| 0.8| 0.0| 0.4| 0.9| 0.8| 1.0|    |    |    |    |    |
| Pb    | -0.5| 0.5      | 0.4 | -0.7| 0.7 | 0.7 | 0.5| 0.9| 0.8| 0.0| 0.4| 0.8| 0.8| 1.0| 1.0|    |    |    |    |
| Ti    | -0.4| 0.6      | 0.4 | -0.5| 0.5 | 0.6 | 0.3| 0.9| 0.9| 0.0| 0.1| 0.8| 0.9| 0.7| 0.7| 1.0|    |    |    |
| V     | -0.3| 0.5      | 0.4 | -0.4| 0.4 | 0.5 | 0.3| 0.9| 0.8| 0.0| 0.0| 0.8| 0.9| 0.7| 0.7| 0.9| 1.0|    |    |
| Mg    | 0.2 | 0.0      | 0.1 | -0.3| 0.3 | 0.2 | 0.5| 0.0| 0.0| -0.1| 0.5| 0.0| 0.0| 0.2| 0.3| -0.1| 0.0| 1.0|    |
| Al    | -0.4| 0.5      | 0.3 | -0.5| 0.4 | 0.6 | 0.1| 0.8| 0.8| -0.1| 0.0| 0.7| 0.9| 0.7| 0.7| 0.9| 0.8| 0.1| 1.0|

-bold values (p < 0.001; n = 87).
cur in any layer during sedimentation processes, depending on the available organic matter. This relation may be observed in correlation values between Iron and calcium carbonate and total organic matter ($r = 0.6$) (see Table 3).

Arsenic and Lead enrichment in Brazilian coastal environments were reported in several works as [79]-[84]. High levels of As (i.e. above the nationally threshold of 70 mg.kg$^{-1}$) were found in shelf, beach and nearshore sands, lagoonal/swamp and mangrove sediments of Espírito Santo, Bahia [80] [81], Rio de Janeiro [82] and Rio Grande do Sul states [84]. A significant positive correlation between As and calcium carbonate ($r=0.6$) could indicate that the calcareous bioclasts participate in metalloid retention and its accumulation in coastal sediments as observed by [81]. This affinity with carbonates added to the As inputs from: (a) the eroded material from continental rocks and sediments [80] local sources (Borborema Crystalline Complex and the Barreiras Group ferrous sandstones); (b) the groundwater aquifer [82]; (c) the atmospheric fallout [82], could implies in the high values observed in the studied core (> 120.0 mg.kg$^{-1}$). Similarly as observed for the Paraíba do Sul delta region [82] and Patos Lagoon estuarine sediments [84] the Arsenic is retained mainly by iron hydroxides in upper and probably by sulfides in lower layers due to diagenetic processes. In fact the high significant correlation of As with iron ($r = 0.9$) and Pb (1.0) reinforce the local enrichment from post depositional processes and atmospheric fallout, also indicated by the Pb enrichment, especially in the upper units of the core (1 and 2).

[85] observed finer sedimentation and TOM peaks associated with occurrence of floods registered in Recife to the lower Capibaribe Estuary. In case of middle estuary is not possible to identify finer sedimentation associated with flood registers. It is believed that the suspended sediments by rains transport is deposited mainly in the middle estuarine area, registering a prevalence of fine sedimentation.

5. Conclusions

The modification of urban space changed the sedimentation patterns in the middle Capibaribe Estuary. The core registered four units with different sedimentary and geochemistry characteristics. First unit (Unit 1), represent ages before 1812 year, showed mangrove environment characteristics with predominance of fine sediments, high total organic matter contents and heavy metal concentrations possibly from natural sources.

The second unit (Unit 2), from 1812 to 1937, showed a small influence of sand characteristics with slightly decreases in the fine fraction percentages, TOM contents and heavy metals concentrations. This increase in sand percentages may be associated with the beginning of urban expansion processes in the middle estuary, together with intensification of monoculture of sugar cane. These two processes were responsible by majority of the Atlantic Forest deforesting, increasing the lixiviation and the transport of sandy sediments from the margins to the river.
Third unit (Unit 3), from 1937 to 2004, showed the highest sand percentages of the sedimentary core, characterizing an exclusive sandy unit with low fine fractions percentages, total organic matter contents and heavy metals concentrations. This unit represents the intensification of urban processes expansion of Recife City. Increases in landfills of flood areas, deforesting of mangrove areas together with intensification of building processes were the main factors that contributed in the increase of sand sedimentation in estuarine system.

The fourth unit (Unit 4) showed increases in finer sedimentation, in TOM contents and heavy metals concentrations. The environmental improvement projects for Capibaribe River, in late 20th and early 21st centuries, were responsible by the reforesting of mangrove marginal areas of the estuary. These projects showed great results in increase of mangrove areas and these progradations could be observed in the higher mud contents in this unit of the sedimentary core.

Although all modifications that occurred since the 18th century, the Capibaribe River was not possible to register clearly the anthropic contamination to middle estuary area. The metal concentration records may have been also influenced by natural sources, which are rich in heavy metal concentrations and surround the estuarine area. And this way, the anthropic influences could have been masked by this geological character. Although as observed by the presented results, and in similar studies, that the enrichment of As and Pb could be directly related to these elements retention and accumulation by diagenetic processes, associated to several local sources as: the carbonate sediments, the local rocks and continental sediments, the groundwater aquifer and the atmospheric fallout.

Lastly, this research may be useful in the comparison with similar sedimentary studies in densely urban estuarine areas. As well as to subsidize the determination of local metals reference/background levels in future studies employing the data collection presented in this paper.

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