**Original Research** 

# Determining Salinity and Ion Soil Using Satellite Image Processing

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## Abstract

Arid and semi-arid zones frequently present salinity problems in soils. The agriculture of the municipality of Ahome, Sinaloa has an agricultural region where its soils are characterized by problems of salinity and sodicity – conditions that reduce production. Salinity can be detected by implementing remote sensing techniques; there are ways to enhance the detection of satellite salinity through the use of diverse quantitative models, using the spectral signature of each of the components of the study area through algorithms named indices. For this study we used the normalized differential salinity index (NDSI) from a Landsat OLI image for the southern area of the city, which is related to the electrical conductivity (EC) of the soils (R = 0.90). At the same time, it is related to some anions and cations. As a result, it is possible to determine since the NDSI, the anion Cl– and Cations Na<sup>+</sup>, Ca<sup>++</sup>, and Mg<sup>++</sup>. We found a relationship between EC - Cl– (R = 0.94), EC - Na<sup>+</sup> (R = 0.84), EC - Ca<sup>++</sup> (R = 0.85), and EC-Mg<sup>++</sup> (R = 0.86). The electrical conductivity in the field and laboratory, anions, cations, and NDSI index were filtered with the Kalman filter obtaining better fitter, eliminating dispersivity in the variable relations.

Keywords: NDSI, salinity, electric conductivity, anions, and cations

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## Introduction

It is estimated that between 20 and 30% of soils worldwide are affected in varying degrees by salinity [1], causing environmental problems, mainly in agriculture [2]. The main factors that cause salinity in soils are the weathering of minerals and the high levels of the phreatic level with high concentrations of salt and surface water [3].

Arid regions are prone to accumulate soluble salts in close vicinity to the surface. There are different forms of salinization of soils, they are classified as primary and secondary; primary salinization consists of the development of salts through natural processes, geological deposits, and physical and chemical erosion that cause the movement of ions Ca <sup>++</sup>, Mg <sup>++</sup>, Na<sup>+</sup> [3, 4].

On the other hand, secondary salinization is caused by human interference in production processes, inadequate irrigation practices, and poor drainage conditions [3, 4], which promote the enrichment of soluble salts and the decrease in the productivity of agricultural lands with irrigation systems [5, 6]. Salinization of soils should be addressed, so information on its scope and magnitude is required for better planning and implementation of effective recovery programs [8].

Salinity can be detected by implementing remote sensing techniques [9]. The salinity process is not exclusive to the surface, but to the whole profile, causing a limitation in the use of optical sensors [10]. However, there are ways to potentiate salinity detection by satellite through the use of diverse quantitative models, using the spectral signature of each of the components of the study area [10-13]. Wu et al. [14] used images of the Landsat and SPOT satellites in Iraq with which they handled the NDVI indices and soil vegetation adjustment index (SAVI), and estimated the electrical conductivity of the soil through the indices. Subsequently, Zewdu et al. [15] estimated and mapped salinity in southern Ethiopia with four classes of soil affectation according to FAO.

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Soca et al. [16] worked on determining salinity degradation on farmland using images from the Spot, Landsat 5, and Landsat 7 satellites for the northern coast of Peru. Poenaru et al. [17] worked to determine the sensitivity of remote sensing of the Landsat 8 satellite in the identification of saline zones for the care of crop lands. Their results show that saline soils have a higher reflectance in all bands of the images as well as that the NDVI index presents better values in estimating the quality of the crops and in the classification of those whose soils are affected by salinity. Some authors have indicated in Table 1 (indices for the determination of salt areas) the use of mathematical algorithms for the satellite images spectral bands in order to emphasize and facilitate the detection of the zones affected by salt. Among them stands out the normalized differential vegetation index (NDVI), the normalized differential salinity index (NDSI), and the brilliance index (BI), as well as several variants of the index of salinity.

Among the publications made with the use of these indices, several authors agree that the application of both NDVI and NDSI obtain similar results in the detection of salt areas [1, 3, 18-21]. However, most of them avoid using NDSI, despite having this algorithm closer to the process of salinization in soils and more related to the electrical conductivity of the same [1, 6, 18-21].

On the other hand, the passage of electric current through a saline solution is found in function of the concentration of salts and temperature; salinity of an extract can be characterized by measuring electrical conductivity (EC) with the use of a conductivity meter in laboratory [5]. Thus, the total dissolved salts are located in the space, due to the high values of electrical conductivity presented by the soil [22].

Mora et al. [23] analyzed the values of salinity by means of electrical conductivity, present ions (Ca<sup>2</sup> +, Mg<sup>2</sup> +, Na +), and sodicity through the ratio of sodium uptake and alkalinity by pH in a semi-arid irrigation district in Spain. The results of the measurements showed soils with a gradual increase of electrical conductivity that translates to higher concentrations of sodium throughout the temporary study.

This study aims to create a map identifying the presence of salts present in the soil beginning from the salinity and vegetation indices (NDSI and NDVI respectively) of the images of Landsat 8, in addition to measurements in situ of EC, humidity, pH, and laboratory analysis.

## Study Area

This work was accomplished in an agricultural land lot from the municipality of Ahome, Sinaloa, Mexico. In the study area, very dry climate is predominant, very warm and warm with summer rains and an average annual temperature of 25.3°C for the series 1986-2015 [24], located between the meridians 108°59'1'' and 109°06'19'' west and the parallels 25°38'25'' and 25°46'53'' north (Fig. 1). The total annual mean precipitation of 357.7 mm is for the series 1981-2010 [25]. The surface in the study area comes from the Cenozoic era of the quaternary period and the chromic Vertisol soil predominates, which is present in the area of study having medium and high fertility; however, it can present problems of drainage and salinity. These are cases where they are dry and found in place of plastics and sticky. The study of the soils in the region undoubtedly represents an alternative to mitigating problems related to agricultural management



Fig. 1. Location of the study area.

and risks in livestock farming and construction [25]. Additionally, the majority of the study area presents pH between 6.5 and 7.5, although in salt-affected soils this varies depending on the saline concentration of this one, specifically of sodium. Nevertheless, the region is characterized by its agricultural activity, and among its main crops are potato, sorghum, mango, corn, and beans [26].

#### **Materials and Methods**

Fig. 2 shows the methodology used in the present work, where it can be observed that the soil salinity can be estimated by satellite image processing, directly through measurements in situ or through the extraction of soil samples, and determine it in the laboratory, which in turn are related to some anions and cations previously filtered with the Kalman filter. The soil features of the study area are presented in a table of basic statistics.

The image of the Landsat 8 OLI satellite was consulted on the United States geological website (USGS) [27] dated June 2016, and location Phat 33, Row 42, in the area corresponding to the study area. Through the software Idrisi Selva 17.0, the bands were adjusted to the bands of the image, using the command Histo and Strech, bands belonging to the visible and near infrared spectrum were respectively measured and cut, that is bands 2, 3, 4, and 5 also perfored them a lineal stretching to their histograms with a 2.5% saturation. With the window function, the image was trimmed to the geographic size of the study area and the BI, NDSI, NDVI, and WATER indices were created with Image Calculator, whose equations are shown in Table 1.

Twenty-six control points were determined with which physical soil parameters were requested. In each of them, wells were made at 30 cm depth, from which 3 kg of soil sample were extracted, and then with a portable conductivity meter HI 98331 Hanna Instruments brand, electrical conductivity readings were taken in situ (Fig. 1). Saline concentration of the soil depends on the type of soil and the moisture that it contains, so the equipment measures the ion concentration of the soil dissolution in function of its chemical composition and its property for retaining water and the chemical elements that it has [34]. The points were geo referenced with a GPS navigator.

Soil samples were subjected to a drying, milling, and sieving process. According to the official Mexican standard NOM-21-RECNAT-2000 [35], the detection of soluble ions and electrical conductivity was performed in laboratory. For determining the soil texture we used the Bouyoucus technique [36]. In the Statistica 7 software, statistical relationships of the variables (previously filtered as indicated in the following section) were realized, obtaining a correlation matrix between



Fig. 2. Methodology applied for determining salinity, and some anions and cations from satellite image processing.

INDEX NAME	FORMULA	REFERENCE
Brightness Index (BI)	$\sqrt{R^2 + NIR^2}$	[1, 21]
Normalized Differential Salinity Index (NDSI)	$\frac{R - NIR}{R + NIR}$	[18, 21, 28-33]
Normalized Differential Vegetation Index (NDVI)	$\frac{NIR - R}{NIR + R}$	[1, 2, 10, 15]
WATER	COMP124 NIR(strech 2.5%)	[19]
COMP124	B + (6 * G) + (36 * NIR)	

Table 1. Indices for determining saline zone

Where R: red band, B: blue band, G: green band, NIR: near infrared band

Table 2. EC test results in situ and salinity indices (NDSI, NDVI, BI, water).

Deint V	v	V	EC in situ	EC in laboratory	NDCI	NDVI	DI	WATED	
Point		Y	mS/cm	mS/cm	NDSI	NDVI	Ы	WAIEK	
1	701831	2848909	0.93	1	-0.1469	0.1469	27816.7	40.0	
2	699951	2848698	0.67	8.4	-0.1633	0.1633	27371.6	39.5	
3	698053	2849001	2.05	5	-0.1383	0.1383	26424.6	39.9	
4	697807	2851161	2.75	10	-0.1423	0.1423	27714.7	40.5	
5	695263	2852699	0.51	1.5	-0.1271	0.1271	27837.4	40.6	
6	694546	2852113	1.3	3.3	-0.1718	0.1718	27503.5	39.7	
7	696793	2849366	3.95	10.5	-0.1146	0.1146	28013.9	39.8	
8	699369	2848103	0.82	3.9	-0.1637	0.1637	28245.3	39.7	
9	701782	2846455	0.98	1.5	-0.1512	0.1512	26893.7	41.7	
10	697731	2844644	2.04	4.5	-0.1681	0.1681	27213.0	40.6	
11	691151	2847308	0.14	6.9	-0.1718	0.1718	27374.9	40.3	
12	694037	2848268	2.99	9.4	-0.1366	0.1366	26668.6	40.4	
13	691079	2847513	2.83	7.6	-0.1377	0.1377	25794.5	39.8	
14	697176	2845538	4.44	8	-0.1201	0.1201	27548.4	39.6	
15	697625	2845561	6.19	10.6	-0.1337	0.1337	26988.7	40.4	
16	697055	2842063	3.77	10.4	-0.1409	0.1409	25304.7	40.5	
17	690137	2844353	11.43	39	-0.0895	0.0895	24303.9	40.9	
18	690095	2845716	12.12	43	-0.0707	0.0707	24061.7	41.2	
19	689946	2846885	12.17	46	-0.0749	0.0749	26051.6	41.4	
20	691192	2843015	7.14	34	-0.133	0.133	27993.3	40.6	
21	692010	2842474	8.46	35	-0.0984	0.0984	27655.0	40.3	
22	692836	2842475	9.06	23	-0.0918	0.0918	26661.3	40.7	
23	696117	2838589	7.95	25	-0.1024	0.1024	25271.6	40.7	
24	695747	2837102	12.68	49	-0.0699	0.0699	24537.8	41.0	
25	692009	2842860	11.15	41	-0.0979	0.0979	24607.6	40.9	
26	690711	2842432	2.41	7	-0.1228	0.1228	27394.7	39.8	



Fig. 3. Correlation between the indices NDSI, NDVI, BI, WI, and electrical conductivity in situ.

the indices obtained with the satellite image and the electrical conductivity measured in the soil, in order to characterize the study area.

## The Kalman Filter

The Kalman filter (KF) is a recursive algorithm that optimally estimates signals by filtering white gaussian noise [37]. It can be seen as a low-pass filter [38]. The KF has been a tool successfully used in aerospace [39], data fusion [38], signals filtering, navigation [40], and control theory, among others.

The discrete linear system considered for the KF is as is shown:

$$x_k = A_{k-1} x_{k-1} + W_{k-1} \tag{1}$$

$$y_k = C_k x_k + v_k \tag{2}$$

...where  $x_k$  describes the unknown signal. The measurement with white gaussian noise is described by  $y_k$ . The matrices  $A_{k-1}$  and  $C_k$  are factors and  $W_{k-1}$ , and  $v_k$  represents uncertainties of white gaussian noise associated with the unknown signal and the measurement, respectively. Both are considered of zero mean, uncorrelated, and with matrices of variance  $Q_{k-1}$  and  $R_{k}$ , respectively.

The KF for the linear system described above is given by

$$\hat{x}_{k}^{-} = A_{k-1} \, \hat{x}_{k-1}^{+} \tag{3}$$

$$P_k^- = A_{k-1} P_{k-1}^+ A_{k-1}^T + Q_{k-1}$$
(4)

	EC in situ	Ec in laboratory	NDSI	NDVI	BI	WATER
EC in situ	1.00					
EC in laboratory	0.96	1.00				
NDSI	0.91	0.85	1.0			
NDVI	-0.91	-0.85	-1.0	1.0		
BI	-0.71	-0.65	-0.6	0.6	1.00	
WATER	0.59	0.58	0.5	-0.5	-0.55	1.00

Table 3. Correlation matrix of salinity indices and EC in situ and laboratory

Bold correlations are significant at p< .050



Fig. 4. Correlations between EC and concentrations of ions.

$$K_{k} = P_{k}^{-} C_{k}^{T} (C_{k} P_{k}^{-} C_{k}^{T} + R_{k})^{-1}$$
(5)

$$\hat{x}_{k}^{+} = \hat{x}_{k}^{-} + K_{k}(y_{k} - C_{k}\hat{x}_{k}^{-}) \tag{6}$$

$$P_k^+ = P_k^- - K_k C_k P_k^- \tag{7}$$

...where  $\hat{x}_{k}^{-}$  is the estimate signal of  $x_{k}$  before processing the measurement  $y_{k}$  in the instant k.  $P_{k}^{-}$  is the variance of the estimation error  $\hat{x}_{k}^{-}$ .  $K_{k}$  is the optimization factor, usually named Kalman gain.  $\hat{x}_{k}^{+}$  describes the estimate signal of  $x_{k}$  after processing the measurement  $y_{k}$  in the instant k and  $P_{k}^{+}$  is the variance of the estimation error  $\hat{x}_{k}^{+}$ .

For the particular case of this work that correspond to signals filtering, the system (1) and (2) as well as the KF are specified with the matrices  $A_{k-1} = I_{8*8}$ ,  $C_k = I_{8*8}$ ,  $Q_{k-1} = 0.1/I_{8*8}$ ,  $R_k = 0.1/I_{8*8}$  and initial conditions  $\hat{x}_k^+ = [1 \quad 0.93 \quad 0.1 \quad 0.1469 \quad 5 \quad 3.5 \quad 0.5 \quad 1.6 \quad 2]^T$  and  $P_0 = I_{8*8}$ .

## **Results and Discussion**

## Relationship between Salinity and Electrical Conductivity Indices *in situ*

Twenty-six points were located at which electrical conductivity was measured in situ and for each point we determined the BI, NDVI, NDSI, and water, which can be seen in Table 2. By means of the use of simple correlation, the salinity indices with the electrical conductivity of the soil were analyzed, finding significant differences for the indices NDSI (r = 0.91), NDVI (r = -0.91), BI (r = -0.71), and water (r = 0.59), which can be seen in Fig. 3. Table 3 shows that the NDSI salinity index used in this work was the one that obtained the highest correlation with the electrical conductivity of the soil, so it was used to characterize the salinity in the study area and is given by the expression:

$$NDSI = 0.006698EC \text{ in situ} - 0.15986 \quad (8)$$

				-									
		OBTAINED IN LABORATORY						FILTERED DATA					
No.	рН	FC lab	EC lab SOLUBLE IONE (Meq/			E (Meq/lt)		FC lab	SOLUBLE IONE (Meq/lt)				
			Ca <sup>2+</sup>	Mg <sup>2+</sup>	Na <sup>+</sup>	HCO <sub>3</sub> -	Cl-		Ca <sup>2+</sup>	Mg <sup>2+</sup>	Na <sup>+</sup>	HCO <sub>3</sub> -	Cl-
1	6.9	1	5	3.5	0.5	1.6	2	1.0	5.0	3.5	0.5	1.6	2.0
2	7.5	8.4	40	30	40	1.6	20	7.9	47.4	35.5	12.0	1.1	21.4
3	7.3	5	25	15	10	0.7	15	5.3	26.1	17.2	16.2	1.0	14.6
4	7.5	10	50	45	35	1.1	20	8.0	40.0	29.5	20.4	1.2	28.0
5	7.6	1.5	8	5	1	1.3	2.5	1.0	5.0	3.5	0.5	1.6	2.0
6	7.4	3.3	18	9.5	4.5	2	7	3.6	19.0	12.1	7.1	1.7	7.5
7	7.3	10.5	65	50	15	0.8	30	9.6	31.5	20.7	67.8	1.0	29.0
8	7.5	3.9	22.5	10	5.5	0.8	6	3.9	18.9	13.3	4.5	1.0	11.6
9	7.4	1.5	8	5	1	0.9	2.5	3.0	17.2	10.6	3.7	0.9	6.1
10	7.4	4.5	20	15	5	1	15	4.1	20.6	16.2	11.2	1.2	8.4
11	7.1	6.9	40	25	5	0.8	20	5.4	32.0	19.7	8.0	0.9	11.9
12	7.2	9.4	45	35	30	1.4	35	7.9	33.8	21.8	32.0	1.2	27.4
13	7.1	7.6	40	35	5	1.7	20	5.9	28.0	20.9	26.5	1.6	13.8
14	7.3	8	30	15	45	1	30	7.8	40.0	32.9	10.9	1.5	23.1
15	7.9	10.6	30	20	90	0.8	30	10.1	30.6	20.3	69.2	0.9	42.0
16	7.5	10.4	30	20	70	0.8	50	8.2	40.9	34.4	27.8	1.1	17.9
17	7.2	39	175	100	325	0.6	300	37.3	169.4	134.7	256.8	0.6	277.8
18	7.2	43	200	175	275	0.5	325	40.3	181.3	130.5	379.4	0.7	333.2
19	7	46	425	275	150	0.5	375	40.7	161.9	111.6	330.3	0.7	328.1
20	7.6	34	175	100	225	0.6	225	28.0	119.8	69.5	227.3	0.7	201.4
21	7.4	35	175	125	200	0.7	250	35.9	197.2	133.2	204.6	0.7	256.9
22	7.6	23	125	75	50	0.6	100	26.1	151.0	98.9	103.5	0.7	184.7
23	7.3	25	150	100	100	0.8	200	27.9	152.6	97.2	109.0	0.6	159.9
24	7	49	200	150	550	0.6	425	42.7	327.4	221.4	190.8	0.5	337.9
25	7.3	41	150	100	300	0.7	325	37.3	233.2	146.4	211.9	0.6	268.1
26	8.1	7	25	15	30	1.5	30	5.8	31.9	20.5	4.8	0.9	16.8
L								,					

Table 4. Behavior of electrical conductivity and anions and cations in the study area

Fig. 4 also shows that the electrical conductivity of the soil in laboratory (EClab) can be correlated with that obtained in situ, finding a lineal relationship of equation 9, with an adjustment of r = 0.957.

$$EC_{lab} = 3.48069 \cdot EC_{in\,situ} \tag{9}$$

## Determining Relationships between Electrical Conductivity in Laboratory, pH, and Some Ions

From the values obtained in the laboratory of Ce, pH, and concentrations of the ions of Cl<sup>-</sup>, Na<sup>+</sup>, Ca<sup>++</sup>, and Mg<sup>++</sup> found in Table 4, we observed the strong correlation between Ce lab and Cl<sup>-</sup>, Na<sup>+</sup>, Ca<sup>++</sup>, and

Mg<sup>++</sup>. Once you know the electrical conductivity of the soil, it is possible to estimate each of the elements analyzed for the study area, with the exception of pH, for which no significant differences were found with an adjustment of r = -0.29 (see Table 5).

Likewise, when comparing the NDSI index it provides greater accuracy in the study area, so that from their values obtained, they are correlated with the electrical conductivity of the soil and with the concentrations of Cl<sup>-</sup>, Na<sup>+</sup>, and Ca<sup>++</sup>, as well as Mg<sup>++</sup> ions determined in laboratory. In Fig. 4 these concentrations are observed, obtaining a higher correlation between electrical conductivity and the chloride ion (r = 0.99).

		pH	EC lab	Ca <sup>2+</sup>	Mg <sup>2+</sup>	Na <sup>+</sup>	Cl-
	pН	1					
	EC lab	-0.29	1				
	Ca <sup>2+</sup>	-0.35	0.9	1			
OBTAINED IN LABORATORY	Mg <sup>2+</sup>	-0.38	0.91	0.99	1		
	Na <sup>+</sup>	-0.26	0.89	0.65	0.68	1	
	Cl-	-0.35	0.99	0.89	0.9	0.91	1
	EC lab		1				
	Ca <sup>2+</sup>		0.95	1			
FILTERED DATA	Mg <sup>2+</sup>		0.95	0.99	1		
	Na <sup>+</sup>		0.93	0.80	0.81	1	
	Cl-		0.99	0.94	0.95	0.95	1

Table 5. Correlation matrix pH, CElab, and some ions

Bold correlations are significant at p<.050

In this way, the relationships between electrical conductivity and the concentrations of anions and cations previously determined in the laboratory are obtained and make faster calculations, and the linear correlation was determined and in the case of chloride, the exponential was also performed.

In equations 10, 11, 12, 13, and 14 the equations are observed, obtaining greater adjustment with the chloride, sodium, calcium, and magnesium ions with an r greater than 0.91.

$$Cl^{-} = 8.3404 EC lab - 31.724$$
 (10)

$$Cl^- = 1.12386 EC lab^{1.52648}$$
 (11)

 $Na^+ = 6.31112EC lab$  (12)

 $Ca^{++} = 5.49824EC lab - 4.49$  (13)

Mg<sup>++</sup> = 3.72839EC lab - 2.646 (14)

## Determining Ions as from Salinity Index

Practically, from the values of the NDSI it is possible to estimate the electrical conductivity in situ and from this, the value of the electrical conductivity in laboratory and estimate the concentration of the ions Cl<sup>-</sup>, Na<sup>+</sup>, Ca<sup>++</sup>, and Mg<sup>++</sup>.

In Figs 5, 6, 7, and 8, the concentrations of saline ions were estimated within the study area in addition to being able to map and compare them with the prediction of NDSI to each of the ions:  $Cl^-$ ,  $Na^+$ ,  $Ca^{++}$ , and  $Mg^{++}$ .

Chlorine is absorbed by the plant in the form of chloride ion, in suitable concentrations, from 0.2 to 0.4 mg/g in most plants, it contributes as an essential factor for photosynthesis, in addition to developing a

regulatory role in stomatal opening, acting as a cation neutralizer and reducing the effect of diseases caused by fungi. While in excessive amounts, it causes thinning of leaves and even dries plants, being a factor that limits the growth of plants in arid and semi-arid soils [41]. Fig. 3 shows the correlation of the chlorine ion with electrical conductivity. It presents an average of 110 Meq/ 1, presenting a minimum of 2 Meq/1 and a maximum of 225 Meq/1 in the sodic saline areas, in the map of Fig. 5 which is shown on the left, it is observed the map of the estimates as from the values of the NDSI index.

On the other hand, sodium can be beneficial during periods of drought, since it slows plant wilt while maintaining the cellular osmotic potential, as well as being beneficial in the soil structure when dispersing clays [41]. Sodium has a critical role in soil health, and excessive amounts of sodium affect the physical structure of soil because it aggregates fragmenting by causing a decline in water and air permeability of soil, adversely affecting root development [42, 43]. The same behavior is observed in both comparative maps of Fig. 6, of the analyzed samples, an average of 98.8 Meq/l was obtained showing in agriculture regions minimum levels of 0.5 Meq/l, while in arid regions, values of 550 Meq/l.

Meanwhile, calcium is generated by several types of rocks, of mother rock, mainly its abundance is found in function of the presence of organic matter present in soil, to a lower content of organic matter, the greater the presence of this mineral [28]. It is the cation of greater abundance in soil, it improves in plants the development of leaves and roots, activates enzyme-forming systems, in addition to reducing nitrate in plants [32], Fig. 7 shows the comparative maps for the cation  $Ca^{++}$ , which presents a similar behavior than the prediction of the NDSI to the



Fig. 5. Comparative map of concentrations of Cl<sup>-</sup>: a) concentration measured in laboratory and b) estimated concentration as from NDSI.

concentrations obtained in laboratory presenting a maximum value of 425 Meq/l in areas marked as saline.

The accumulation of magnesium in its ionic form  $Mg^{++}$  can occur by the same nature of soil as with calcium, as well as by using water with high concentrations for irrigation use helping raise the sodium level in clays. However, at toxic levels it acts as a flocculant, in addition to reducing plant growth and promoting the dispersion and expansion of clay due to the loss of water-conducting properties [33], comparative maps of the concentrations of  $Mg^{++}$  are observed in Fig. 8, which present a similar significance to the



Fig. 7. Comparative map of Ca<sup>++</sup> concentrations: a) concentration measured in laboratory and b) estimated concentration as from NDSI.

estimate from the NDSI index to that obtained in laboratory, which presents low levels in an average of 59.7 Meq/l.

According to the classification of salinity proposed by Richards [44] and the points taken in the study area shows that most of the area presents problems of salinity which varies from slightly saline to strongly saline. Small portions of non-saline soils were found to the north and northeast of the study area. The application of classification of soils considering the classification of Richards has been used by diverse authors [4, 45-47] in diverse geological environments and is useful for classifying the risk of salinity of the soils. The Kalman



Fig. 6. Comparative map of Na<sup>+</sup> concentrations: a) concentration measured in laboratory and b) estimated concentration from NDSI.



Fig. 8. Comparative map of concentrations of Mg<sup>++</sup> : a) concentration measured in laboratory and b) estimated concentration starting from NDSI.



Fig. 9. Behavior of soil salinity in the study area.

filter is useful for eliminating dispersity in a variable's relationships [48].

## Practical Application

Taking the NDSI values and using the previous models (Equations 8 to 14), as can be seen in Table 6, it is possible to determine the concentrations of chlorine, sodium, calcium, and magnesium by means of interpolations of values between the correlations from the NDSI and electrical conductivity within the study area, which makes it favorable in the practical agricultural application, that is, in the control of soil nutrients required by the crop.

#### Conclusions

In regions where agricultural activities predominate, the NDSI index presents better results to detect the

Table 6. Practical examples of the application of the study.

NDSI	-0.1	
EC in situ = (NDSI+0.15986)/0.006698	8.937	mS/cm
EC lab = $3.48069 \text{ EC}_{\text{in situ}}$	31.106	mS/cm
Na <sup>+</sup> = 6.31112 EC lab	196.31	Meq/l
Cl <sup>-</sup> = 8.3404 EC lab-31.724	227.71	Meq/l
$Ca^{2+} = 5.49824 \text{ EC} \text{ lab-}4.49$	166.53	Meq/l
Mg <sup>2+</sup> = 3.72839 EC lab-2.646	113.33	Meq/l

degree of salinity than the BI and water index. On the whole, with the NDSI index, electrical conductivity, interchangeable ion concentrations (previously filtered with the Kalman filter) and statistical functions, it is possible to determine the degree of salinity and concentration of interchangeable ions in the soil, and any point in the study area. On the other hand, the water index proves to be a good indicator in the presence of wet areas by flood or low phreatic level, which together with the NDSI index allows for the determination of saline areas caused by irrigation with salt water or by leaks of water of the same characteristics.

Of the anions and cations examined, the chloride ion had a higher correlation with electrical conductivity, which represents a greater ease of detection in the NDSI index. However, it is important to note that this is not only found alone, and therefore also functions as an indicator of the presence of the other analyzed ions.

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#### **Conflict of Interest**

The authors declare no conflict of interest.

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