NORTH CAROLINA DEPARTMENT OF WATER AND AIR RESOURCES

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REPORT OF INVESTIGATIONS NO. 8

GROUND WATER RESOURCES OF THE BELHAVEN AREA NORTH CAROLINA

Ву

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PREPARED COOPERATIVELY BY THE GEOLOGICAL SURVEY
UNITED STATES DEPARTMENT OF THE INTERIOR
AND THE NORTH CAROLINA
DEPARTMENT OF WATER AND AIR RESOURCES

RALEIGH

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The Honorable Dan K. Moore Governor of North Carolina Raleigh, North Carolina

Dear Governor Moore:

I am pleased to submit Report of Investigations Number 8, "Ground-Water Resources of the Belhaven Area, North Carolina" by Orville B. Lloyd and Edwin O. Floyd, U. S. Geological Survey.

This report contains the results of a study of ground-water conditions in the vicinity of Belhaven and evaluation of possible new sources of water supply for the Town of Belhaven. The study was conducted cooperatively by the U. S. Geological Survey and the North Carolina Department of Water and Air Resources.

George E. Richett

George E. Wickett

THE GROUND-WATER RESOURCES OF BELHAVEN AND VICINITY, NORTH CAROLINA

By
Orville B. Lloyd, Jr.,
and
Edwin O. Floyd

ABSTRACT

The area of study is located in northeastern Beaufort County in the Coastal Plain province of North Carolina and is about 150 square miles. Topography is relatively flat and elevations range from near sea level at the streams to about 20 feet above sea level in the northwestern part of the area. Average annual rainfall is about 52.5 inches, and average annual temperature is approximately 61° F.

The sediments that underlie the area are composed of about 2,500 feet of sand, silt, clay, shell, and limestone beds, that range in age from Cretaceous to Recent. Three aquifers of sand, shell, and limestone (herein designated as aquifers A, B, and C) containing fresh water, occur in the sedimentary section between land surface and a depth of about 300 feet. The sediments below 300 feet are saturated with salty water. Ground water occurs under non-artesian conditions in aquifer A (the water-table aquifer), and it occurs under artesian conditions in aquifers B and C. Theoretically, yields from properly constructed wells screening the full thickness of aquifers A, B, and C are less than 1, 10, and about 80 gpm (gallons per minute) per foot of water-level drawdown, respectively. The total of dissolved solids in water from these squifers is generally below 500 ppm (parts per million), except in the lower part of aquifer C where high concentrations of chloride are encountered. Water containing excessive concentrations of dissolved iron, hardness-causing constituents, and/or hydrogensulfide gas is common in each of the fresh water-bearing aquifers in the area.

GROUND-WATER RESOURCES OF BELHAVEN AND VICINITY

Large ground-water withdrawals associated with the open-pit mining of phosphate deposits southwest of the area have caused an extensive cone of depression in aquifer C, the major artesian aquifer in the area. At the time of this investigation, water levels had declined about 12 feet in aquifer C in the vicinity of Belhaven; this decline has increased the probability of salt-water contamination where heavy pumpage from this aquifer is anticipated in the Belhaven area. Aquifers A and B, to date, are relatively unaffected by substantial water-level declines and the threat of salt-water contamination. They should be considered as an alternative source of ground-water supply if further withdrawals from aquifer C create water-management problems.

INTRODUCTION

PURPOSE AND SCOPE OF INVESTIGATION

The purpose of this investigation was to determine the lithic character, areal extent, depth, thickness, and water-yielding characteristics of the water-bearing formations and the chemical quality of their contained water in the vicinity of Belhaven, North Carolina.

Pronounced lowering of water levels in the principal aquifer, increased municipal water needs, and the possibility of salt-water encroachment have caused extensive problems in this area. Consequently, in 1966 the town of Belhaven requested the U. S. Geological Survey to make a detailed investigation of ground-water resources in the area. The work was done between June 1966 and June 1967 by cooperative agreement between the town of Belhaven, the North Carolina Department of Water Resources and the U. S. Geological Survey.

DESCRIPTION OF THE AREA

The area of investigation is in the northeastern part of Beaufort County, North Carolina, and is about 150 square miles (fig. 1). About half of the land is cleared and used mainly for agriculture, and the remainder is forested. State Highway 99 enters the area from the north and U. S. Highway 264 crosses the southern part of the area in an east-west direction.

The town of Belhaven, the largest urban center, has a population of 2,365 according to the 1960 census. Estimated total population in the area is 5,000.

PHYSIOGRAPHY

The project area is in the Atlantic Coastal Plain province of North Carolina. The Coastal Plain is generally characterized by a relatively flat surface that slopes gently to the southeast. This surface is divided into a number of marine terraces that were formed by wave and current action during periods when portions of the Coastal Plain were submerged beneath the sea. The terrace surfaces have been dissected by streams and rivers since the sea retreated.

The Belhaven area is located on the lowermost terrace. In the northern and northwestern part of the area drainage is poorly developed, and a large part of the land is occupied by swamps. Elevations in this vicinity range from 8 to 18 feet above msl (mean sea level). In the southern and southeastern part of the area the terrace is better drained by natural streams and ditches, and swamps are less extensive. Land surface elevations in this vicinity range from near sea level at the streams to about 5 to 13 feet above sea level in the interstream areas. Generally all streams flow to the south and southeast and are tributary to the Pungo River (fig. 1).

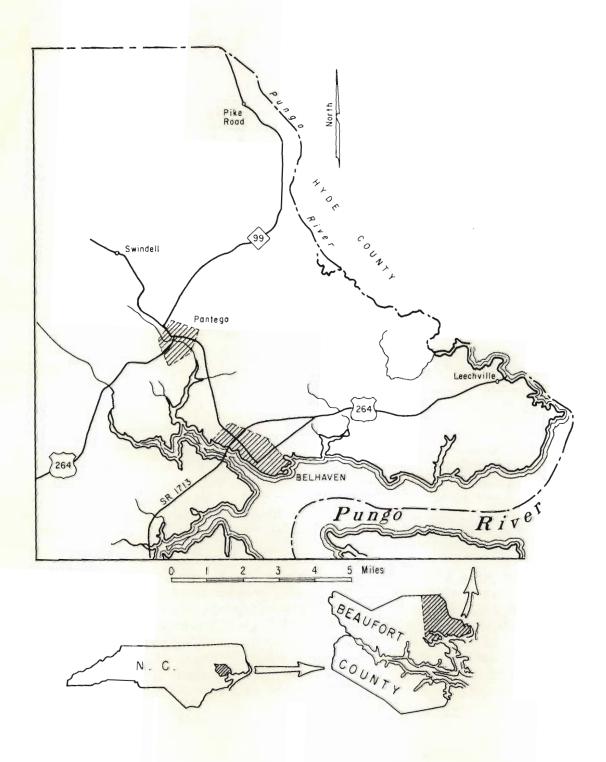


Figure 1. -- Map showing the location of the study area.

CLIMATE

The climate of the area is humid subtropical, characterized by warm summers, relatively mild winters, and precipitation that is well distributed throughout the year. U. S. Weather Bureau records indicate that the average annual temperature is approximately 61°F., and the average annual rainfall is about 52.5 inches. Figure 2 shows a graphic climatic summary for Belhaven.

PREVIOUS INVESTIGATIONS

Clark and others (1912) briefly described the geology and water resources of Beaufort County in a report on the Coastal Plain of North Carolina. This report includes 13 inventoried wells and 3 chemical analyses of water from wells in the vicinity of Belhaven. Mundorff (1945) reported the Yorktown Formation as the major water-bearing formation in the vicinity of Belhaven. He included 3 inventoried wells and 2 chemical analyses of water from wells in this area in his report. Brown (1958) described the relation of the middle Miocene phosphorite deposits to ground water in Beaufort County, and (Brown, 1959) described the geology and ground-water resources of Beaufort County in an eight-county reconnaissance report. This latter report includes 25 inventoried wells, and 9 chemical analyses of water from wells in the vicinity of Belhaven. Nelson and Peek (1964) described the geology and water-bearing units in Beaufort County with emphasis on the potential effects of phosphate mining near Aurora. Kimrey (1964) proposed the name Pungo River Formation for the middle Miocene phosphorites in Beaufort County, and (Kimrey, 1965) briefly described the ground-water hydrology of the county in a report emphasizing the detailed descriptions of the Pungo River Formation in Beaufort County.

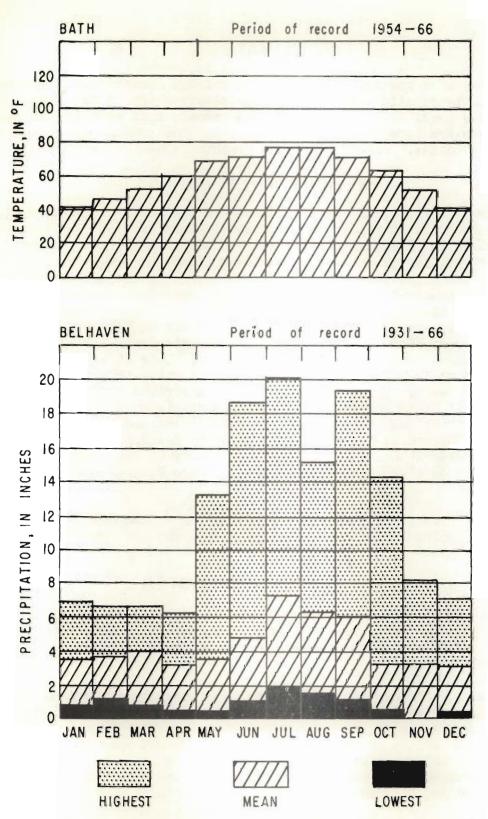
These previous investigations provided accurate and pertinent data that were extremely helpful during the field work and the preparation of this report.

WELL-LOCATION SYSTEM

The well-location system used in North Carolina conforms to the system adopted by the U. S. Geological Survey for the data card processing of well information. This system is intended to locate the position of a given well on the earth's surface.

Positions on the earth's surface may be located by a system of coordinates known as parallels of latitude and meridians of longitude. The parallels of latitude circle the earth parallel to the equator and are numbered from the equator to the poles in degrees, minutes, and seconds, depending upon the angular distance between them and the equator. The meridians of longitude traverse the earth north and south and are numbered east or west from the Greenwich, England, prime meridian in degrees, minutes, and seconds.

The well-location system derived from latitude and longitude coordinates, is based on a grid of 1-second parallels of latitude and meridians of longitude. The wells in a 1-second quadrangle are numbered consecutively in the order inventoried.



Note: Record of temperature taken at Back, N. C., for the period from 1954-66. Record of precipitation taken at Belhaven, N.C., for the period from 1931-66.

Figure 2. -- Climatic summary for Belhaven and vicinity.

The well-location number is composed of fifteen numbers and letters (table 4): the first six numbers and one letter compose the digits of the degrees, minutes, seconds, and indicate northern (N) or southern (S) hemisphere that define the latitude of the 1-second quadrangle; the next seven numbers compose the digits of the degrees, minutes, and seconds that define the longitude on the east side of the 1-second quadrangle; the last number, following a decimal, indicates the order in which wells were inventoried within the 1-second quadrangle (fig. 3)

ACKNOWLEDGMENTS

Special gratitude is due the residents of northeastern Beaufort County for supplying pertinent information about their wells and for allowing tests to be made on their wells. Heber Wilkerson, Superintendent of Public Utilities of Belhaven, J. M. Hudson, and Joe Ratcliff, well drillers, were especially helpful in making well data available during the investigation.

The North Carolina Department of Water Resources drilled a test well about 2 miles north of Belhaven and supplied drill cuttings, logs, and pertinent data collected during and after the drilling.

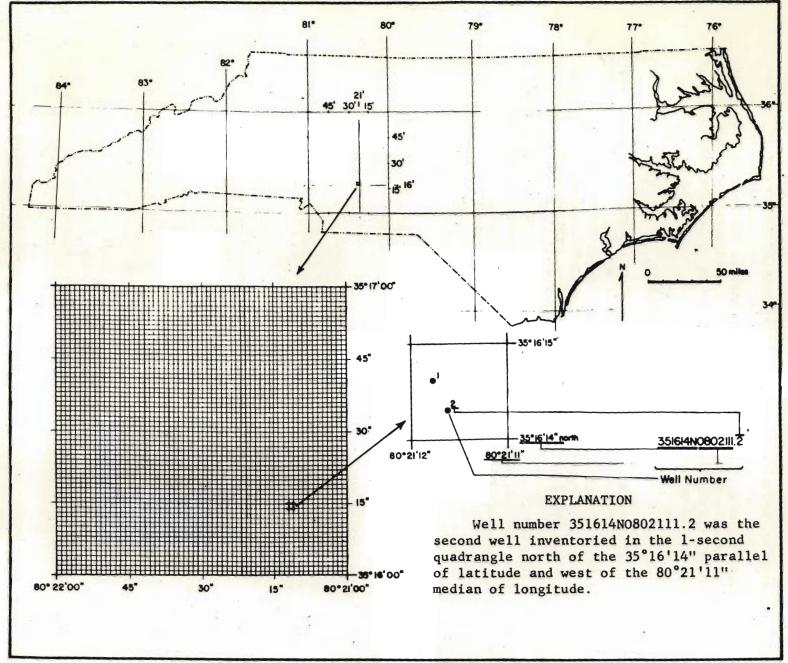


Figure 3.--Map of North Carolina showing the method of determining latitude-longitude well numbers.

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GEOLOGY

An understanding of the geology of the area is requisite to the evaluation of the occurrence, availability, and chemical quality of the ground-water resources. The type and distribution of the rock materials determine the capacity of the rocks to store and transmit water, and the chemical character of the rock materials has a great influence on the amount and kind of dissolved mineral matter found in the water.

GEOLOGIC FORMATIONS

The geologic formations that underlie the area comprise about 2,500 feet of sand, silt, clay, shell, and limestone beds that range in age from Cretaceous to Recent. Figure 4 shows the depths, thickness, and general lithology of the formations penetrated during the test drilling in the area of investigation. The sediments penetrated below the upper half of the Castle Hayne limestone contain brackish and saline water, and for this reason only the units of Eccene and younger age are discussed below.

TERTIARY SYSTEM

Eocene Series

Castle Hayne limestone. -- The Castle Hayne Limestone. (Clark and others, 1912), of middle Eocene age, is not present at the surface in any part of the area of investigation. The top of this formation is an irregular undulatory surface. The formation dips toward the east about 10 feet per mile. Depths to the top of the limestone range from about 175 feet below msl in the west to more than 325 feet below msl in the eastern part of the area (fig. 5). Average thickness is approximately 200 feet.

The upper half of this formation is composed of indurated, light-tan to dark-gray, fossiliferous limestone. The fossils, mainly pelecypods, are in the form of casts and molds; none of the original shell material remains. In the lower half of the formation, hard beds of the indurated limestone described above are interlayered with beds of medium- to coarse-grained, calcareous quartz sand. The sand beds predominate near the base of the formation. Thin layers of calcareous clay, crystalline limestone, and dolomitic limestone are found throughout the lower half of the formation. Glauconite and phosphate (collophane) are the common accessory minerals.

Miocene Series

Pungo River Formation. -- The Pungo River Formation (Kimrey, 1964), of middle Miocene age, occurs entirely in the subsurface and unconformably overlies the Castle Hayne limestone in the area of study. Generally the top of this formation strikes in a north-south direction and dips about 9 feet per mile toward the east. The depth to the top of this unit ranges from 150 feet below msl in the western part, to about 275 feet below msl in the eastern part of the area (fig. 6). The thickness of these sediments ranges from 25 feet in the west to 50 feet in the east, and averages about 35 feet.

The Pungo River Formation consists of greenish-brown to greenish-gray interbedded layers of phosphatic sand, silt, clay, and limestone. Concentrations of up to 50 percent fine- to medium-grained spherules of brown to black phosphate (collophane) are not uncommon in the phosphatic sands.

Yorktown Formation. -- The Yorktown Formation (Clark and others, 1912), of late Miocene age, unconformably overlies the Pungo River Formation. Like the Castle Hayne Limestone and Pungo River Formation, it is confined to the subsurface throughout the area. The top of this formation is found between 20 and 30 feet below land surface, and the thickness of the deposits ranges from about 120 feet in the west to more than 240 feet at the eastern limits of the area. The sediments are composed chiefly of gray, fine- to coarse-grained quartz sands and white to gray shell beds, interlayered with blue-gray silts and clays. Pelecypod, gastropod, coral, and bryozoan remains comprise the major portion of the shell beds. Fine-grained glauconite and phosphate (collophane) are prominent accessory minerals.

QUATERNARY SYSTEM

Post-Miocene Series

Surficial deposits. -- These sediments overlie the Yorktown Formation throughout the study area. Their top is represented by land surface, their thickness ranges from a thin veneer near the streams and rivers to more than 30 feet in some of the higher interstream areas. These deposits are composed of interbedded and lenticular quartz sand, silt, and clay. Iron-oxide stain is prominent on the sedimentary particles.

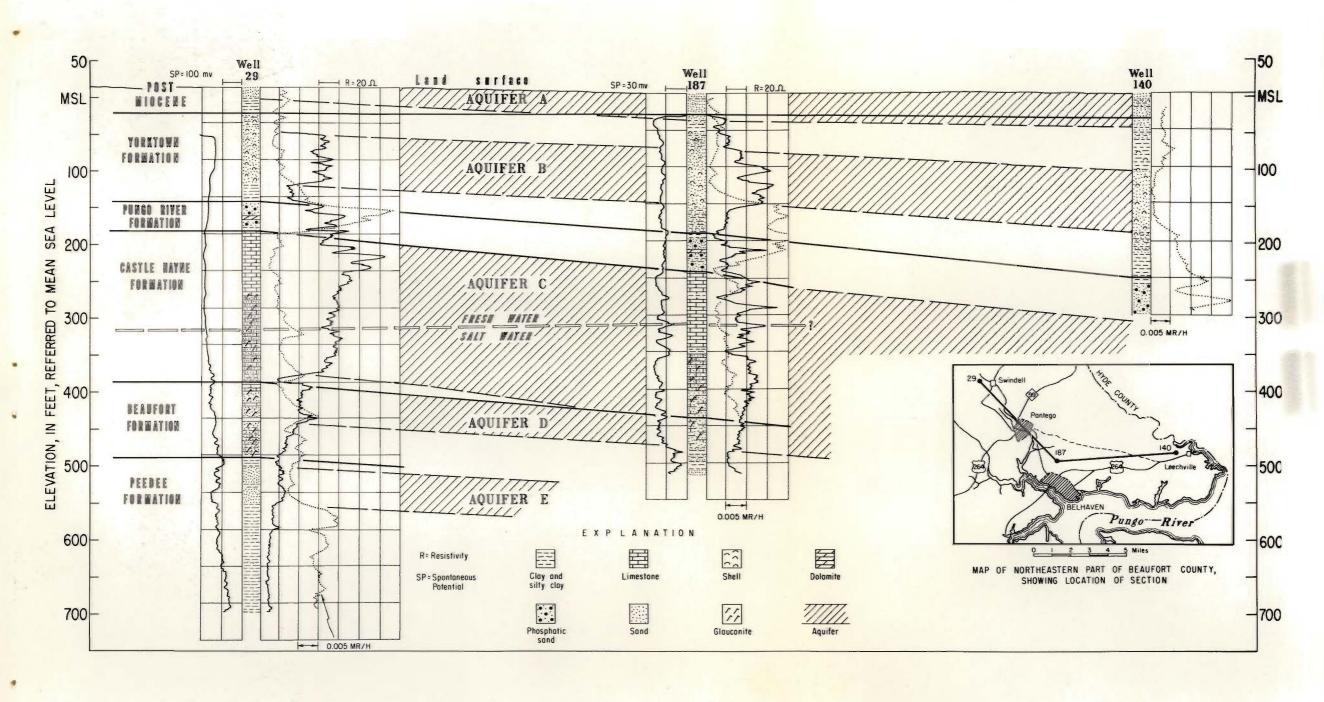
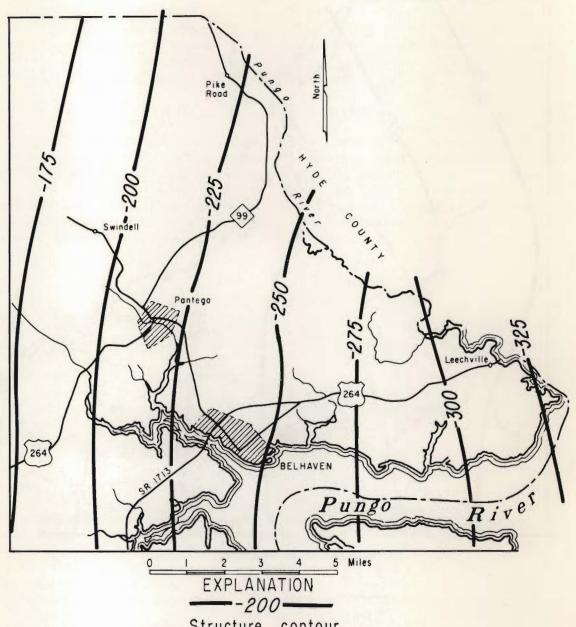


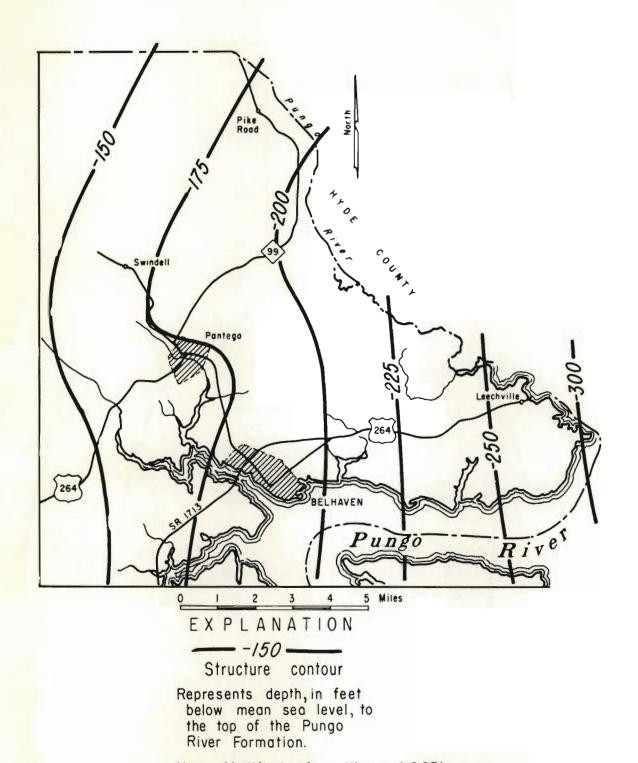
Figure 4.--Section showing the correlation of geologic formations and aquifers by using electric, gamma-ray, and lithic logs in the vicinity of Belhaven, N. C.



Structure contour
Represents depth, in feet below
mean sea level, to the top of
the Castle Hayne limestone.

Note: After Kimrey (1965)

Figure 5.--Map showing the configuration of the top of the Castle Hayne limestone.



Note: Modified after Kimrey (1965)

Figure 6.--Map showing the configuration of the top of the Pungo River Formation.

GROUND WATER

SOURCE AND OCCURENCE

Of the 52.5 inches of precipitation that falls annually on the area, some runs off overland into the streams and rivers, some is returned to the atmosphere by evaporation or through transpiration by plants, and an estimated 15 to 25 percent of it percolates downward to the zone of saturation and becomes ground water.

The zone of saturation is the zone in which all the pore spaces in rock or soil are filled with water. This zone includes all the sedimentary material in the area between basement rock, about 2,000 feet below land surface, and the water table, about 5 feet below land surface.

Water in the zone of saturation occurs under water-table (unconfined) and artesian (confined) conditions. Water occurs under unconfined or non-artesian conditions in the permeable material between land surface and the first impermeable zone found below land surface. Water at the water table, the upper surface of the unconfined ground water, is at atmospheric pressure. The water table is free to rise and fall in response to changes in climatic factors. Under non-artesian conditions the water level in a well represents the water table.

Ground water in permeable material overlain by impermeable units is confined beneath the impermeable layers and occurs under artesian conditions. The upper surface of artesian water is not free to rise and fall in response to changes in climatic factors because it is fixed by the overlying confining layer. However, the hydraulic or artesian pressure exerted against the confining layer responds to changes in climatic factors. Under artesian conditions the water level in a well will rise above the top of the water-bearing zone to a height nearly equal to the hydraulic pressure in that zone. At any given time, the heights of water levels in wells tapping the same artesian zone define the piezometric surface of the water-bearing formation.

AQUIFERS

Any formation, part of a formation, or group of formations in the zone of saturation that will transmit usable quantities of water to wells is called an aquifer.

There are many aquifers within the various geologic formations that underlie the area. Those that are known to contain fresh ground water are found between land surface and about 300 feet below land surface, and are of primary importance here. These water-bearing zones are referred to by letter (A, B, and C) rather than by formation because they occur in only part of one formation or include more than one formation (fig. 4). The deeper aquifers that contain brackish or saline water exclusively will be mentioned only briefly below.

AQUIFER A

Aquifer A is composed of all the saturated permeable sands from land surface to the first impermeable material found below land surface. This generally includes most of the post-Miocene sediments throughout, and both the post-Miocene and the upper part of the Yorktown sediments in the central and eastern part of the area (fig. 4). The average thickness of aquifer A is about 35 feet. Water in this aquifer is unconfined and occurs under non-artesian or water-table conditions. Sustained yields from 1-1/4- to 36-inch diameter wells tapping aquifer A are low, usually less than 10 gpm. The water from this aquifer is generally corrosive and contains excessive concentrations of dissolved iron. Locally it is very hard.

AQUIFER B

Aquifer B includes the shell and sand beds found near the middle part of the Yorktown Formation, and is separated from aquifer A by lenticular silt and clay layers (fig. 4). The impermeable layers overlying aquifer B confine the water in this aquifer under artesian conditions where the silts and clays are thick, and under semi-artesian or leaky-artesian conditions where they are thin. The top of this aquifer strikes in a general north-south direction and dips about 5 feet per mile toward the east. Depths to its top range from 40 feet below msl in the western part, to about 110 feet below msl in the eastern part of the area (fig. 7). Average thickness is about 70 feet (fig. 4). Yields from existing 1-1/4- to 4-inch diameter wells range from 4 to 75 gpm. A properly constructed 10-inch diameter well screened in this aquifer should yield about 10 gpm per foot of drawdown. The water is generally hard and locally it contains excessive concentrations of dissolved iron.

AQUIFER C

Aquifer C is composed of the Castle Hayne limestone (fig. 4). It is separated from aquifer B by relatively thick, and continuous layers of gray silt and clay of the lower part of the Yorktown Formation and by layers of dolomitic limestone and clays in the Pungo River Formation. These confining beds range from about 10 to 90 feet thick from the western to the eastern limits of the area, respectively (fig. 4). Water in aquifer C is confined beneath these impermeable beds and occurs under artesian conditions.

The strike, dip, and depths to the top of this aquifer coincide with the top of the Castle Hayne limestone (fig. 6). Average thickness is about 190 feet (fig. 4). Aquifer C is by far the most permeable water-bearing zone that underlies the area. Yields from existing 1-1/4- to 8-inch diameter wells tapping this aquifer range from 5 to 300 gpm. Theoretically, a properly constructed well, 10 inches in diameter, drawing water from the full thickness of this aquifer should yield about 80 gpm per foot of drawdown. Water in the upper part of aquifer C contains excessive concentrations of hardness causing constituents, and locally excessive amounts of dissolved iron are found. Brackish or saline water is generally encountered in the lower part of the aquifer at about 300 feet below msl (fig. 4).

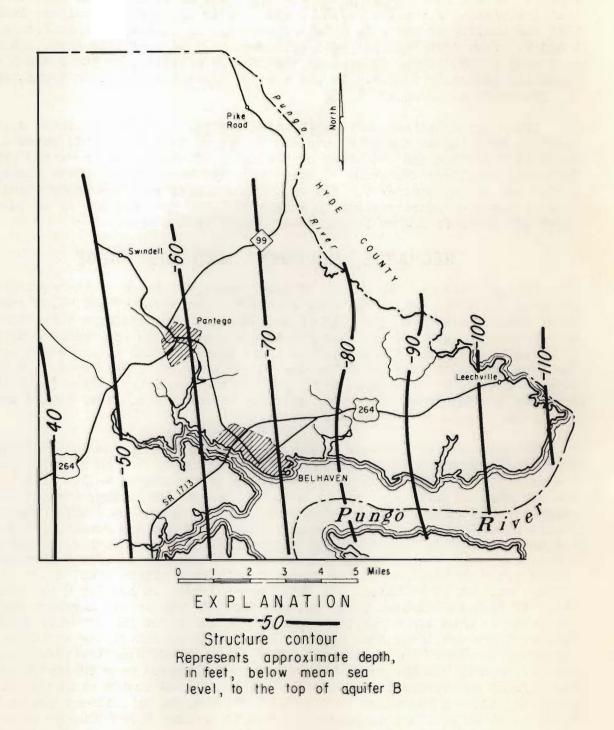


Figure 7. -- Map showing the approximate altitude of the top of aquifer B.

OTHER AQUIFERS

During the investigation, three wells were found that produced water from the Pungo River Formation. Of these wells one was used for domestic supply, one for stock supply, and the other was unused. It was determined that the yields of the two wells in use were low. Chemical analyses indicated that the quality of water from this formation was similar to that from aquifers B and C. From test-drilling data collected as a part of this investigation and by Kimrey (1965) it was determined that the permeability of the Pungo River Formation generally is low. In the area of investigation, this formation is not important as an aquifer.

Two other artesian aquifers (D and E of fig. 4) were penetrated during the test drilling in the area, one consisting of the glauconitic sands of the Beaufort Formation and the other of the sands of the Peedee Formation (fig. 4). Both these aquifers are about 50 feet thick and have approximately the same strike and dip as aquifer C. However, these deeper aquifers contain saline water (chloride concentrations in excess of 6,700 ppm) and are of no value regarding domestic and municipal water supply in the area.

RECHARGE, MOVEMENT, AND DISCHARGE

Precipitation is the source of ground-water recharge to the aquifers in the area of study. Major recharge occurs between November and March when water loss to evapotranspiration is negligible. Most recharge takes place in the higher areas. As the water enters the ground in the recharge areas, it moves in response to gravity from these areas of high water levels toward the areas along streams and estuaries where low-water levels occur. Thus it moves downward, by slow percolation through the confining layers (aquicludes), from one aquifer to another, and laterally within the aquifers toward areas of discharge such as streams and rivers.

Ground water withdrawals from an aquifer can modify the natural rechargedischarge conditions described above. Pumping water from an aquifer causes the water levels to decline in the vicinity of the pumped well. As a result, the piezometric surface or water-table of the aquifer is depressed so that it resembles the shape of an inverted cone (cone of depression) which has its lowest point at the center of pumping. All other factors remaining the same, the cone of depression will spread very rapidly in an aquifer with high permeability. Such a cone of depression has developed in aquifer C from the ground-water withdrawals at the open-pit phosphate mining area, about 15 miles southwest of Belhaven. To date, water levels in aquifer C have declined about 12 feet at Belhaven (fig. 8), and about 3 feet in the northern part of the Belhaven area since pumping began in the summer of 1965. Water levels in aquifer C are now lower than those in overlying and underlying aquifers throughout at least the southern half of the area of investigation. Consequently, nearly all the water that would be discharged from aquifer C into the streams and rivers in the area is being diverted to the mining area. In addition, some of the water that would be discharged by effluent seepage from the other aguifers is being diverted to aguifer C and through it to the mining area. Figure 9 shows the principal areas of recharge and discharge for aquifers A and B. Aquifer C receives recharge from the overlying aquifers throughout the southern half of the area of study and from the underlying aquifers throughout the area.

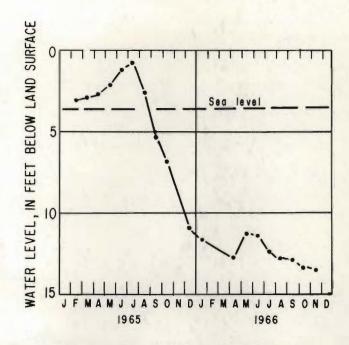


Figure 8.--Water level of aquifer C in observation well at Belhaven, 1965-66.

AQUIFER TESTS

The ability of an aquifer to transmit water and its capacity to store water can be determined by pumping a well screened in the aquifer in question and measuring the discharge in the pumped well and the amount and rate of water-level decline in observation wells that are screened in that same aquifer. The hydraulic characteristics (coefficients of storage and transmissibility) of the aquifer can then be determined from the collected data by mathematical calculations.

The coefficient of transmissibility is defined as the amount of water, in gallons per day, that will flow through a vertical strip of an aquifer 1 foot wide under a hydraulic gradient of 1 foot per foot, at the prevailing water temperature. The coefficient of storage is defined as the volume of water released from or taken into storage in each column of the aquifer having a base of one square foot and a height equal to the saturated thickness of the aquifer, when the water level is lowered or raised one foot. The field coefficient of permeability is the rate of flow of water, in gallons per day, through a 1 square foot cross-section of the aquifer under a hydraulic gradient of 1 foot per foot at the prevailing temperature.

The average hydraulic characteristics for aquifers B and C, the major sources of municipal, irrigation, and domestic water supplies in the area, are listed in table 1. The values for aquifer C have been extrapolated

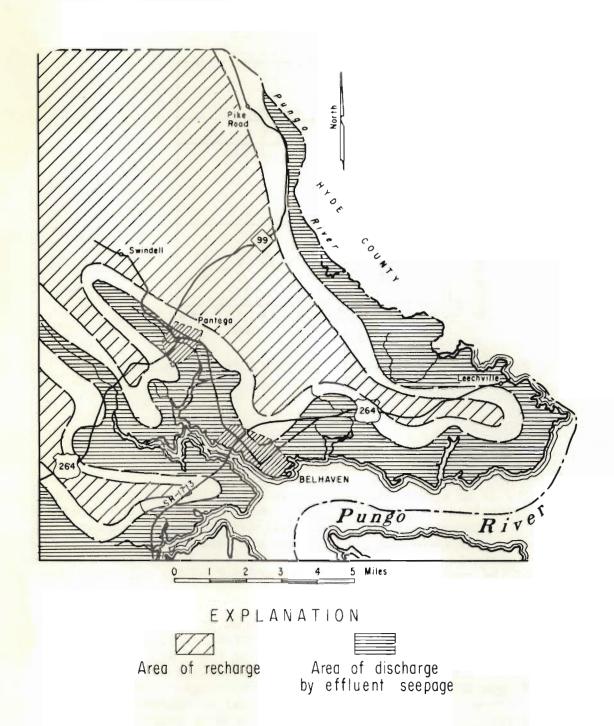


Figure 9. -- Principal areas of recharge and discharge for aquifers A and B.

Table 1. -- Average quantitative values for aquifers B and C

	Aquifer B	Aquifer C
Thickness in feet	67	190
Coefficient of transmissibility, in gallons per day per foot	20,000	180,000 - 230,000
Coefficient of storage, dimensionless	.0004	.0004
Field permeability, in gallons per day per square foot	300	950 - 1220
Calculated specific capacity, in gallons per minute per foot of drawdown		
for 2" wells for 10" wells for 36" wells	9 10 12	60 - 80 82 - 104 95 - 119

from pumping and recovery tests made by J. W. Harshbarger and others on wells finished in this aquifer southwest of the area of study. The results of these tests are assumed to be valid approximations for the area of study because there is little or no appreciable change in the lithic character and thickness of the aquifer between Belhaven and the area of the tests. The values for aquifer B were determined from pumping and recovery tests made on wells in the area of investigation.

The coefficients of transmissibility and storage can be used to determine the water-yielding capacity of wells finished in the aquifer, the effects of pumping on water levels in the aquifer, and the volume and velocity of water moving through the aquifer.

As mentioned above, pumping water from a well causes the water level to decline in the vicinity of the pumped well. The relation between the pumping rate and the amount of drawdown in the well is referred to as specific capacity. It is usually reported in gallons per minute per foot of drawdown after some specified period of continuous pumping, generally one day.

Specific capacity is controlled by the coefficients of storage and transmissibility of the aquifer and the efficiency of the well tapping the aquifer. Calculated specific capacities for aquifers B and C, assuming 100 percent well efficiency, are listed in table 1. The measured specific capacity of a well is generally less than the calculated specific capacity and the difference can usually be related to improper construction and/or poor well development. If a well were pumped for 24 hours at 500 gpm, and the drawdown was 100 feet at the end of this time, the measured specific capacity would be 5 gpm per foot of drawdown. If the calculated specific capacity was 10 gpm the efficiency of the well would be 50 percent.

The size, shape, and growth rate of a cone of depression developed by pumping water from an aquifer depend on the rate and duration of pumping, the coefficients of storage and transmissibility of the aquifer, the amount of ground-water recharge to the aquifer, and the location of impermeable boundaries of the aquifer. Other factors remaining the same, relatively flat cones that spread rapidly develop in aquifers with high transmissibility, and steep cones that spread slowly develop in aquifers with low transmissibility. This is true because water can move more freely in aquifers with high transmissibility; thus smaller head differentials are required to move any given amount of water.

Figures 10 and 11 represent half-sections through calculated cones of depression in aquifers B and C, respectively. These graphs show the theoretical drawdown in water level at selected times and various distances from a well being pumped at a constant rate. In the derivation of the formula used for constructing these distance-drawdown plots, it is assumed that the coefficients of storage and transmissibility are constant and the same in all directions, the aquifer is infinite in areal extent, and confined between impermeable beds so that all the water is drawn from storage in the aquifer, the well penetrates and draws water from the full thickness of the aquifer, and that ground water is released from storage instantaneously with a decline in artesian head. Variations from these calculated plots can be expected because the above conditions are seldom, if ever, completely met, and the distance-drawdown graphs are constructed from average values of the hydraulic characteristics of the aquifers. Therefore, when a well is constructed it is important to test-pump it and determine the hydraulic characteristics of the aquifer at the well. Actual drawdown will generally be less than the calculated drawdown after extended periods of pumping except where impermeable boundaries of the aquifer are intercepted by the cone of depression.

During the first few minutes of pumping, the water withdrawn from the well is largely derived from storage in the aquifer. This period is marked by a relatively rapid decline in water level and/or artesian pressure in the vicinity of the pumped well. In artesian aquifers, such as aquifer C, as the artesian pressure declines, the gradient between the pumped aquifer and the overlying aquifers is reduced. This reduction in gradient results in a reduction of the rate of natural discharge through the confining beds. As this natural discharge is diverted to the pumping well there is a corresponding decline in the rate at which water is derived from storage. If a cone of depression in the Belhaven area extends into areas where the aquifers are being recharged, such as the nearby Pantego Swamp, there will be an increase in the rate of recharge. When the reduction in natural discharge and the increase in recharge equal the rate of pumping, the removal of water from storage and the decline in artesian pressure cease, and the cone of depression becomes stabilized.

Figures 10 and 11 may be used for estimating drawdowns at short distances from a pumped well over periods of time up to 100 days. Similar plots can be constructed for any desired yield, because drawdown is approximately proportional to the pumping rate for wells screened in artesian aquifers. To calculate the drawdown for some desired yield other than that shown on the figures, divide the drawdown shown on the graph by the indicated pumping rate, then multiply by the desired pumping rate.

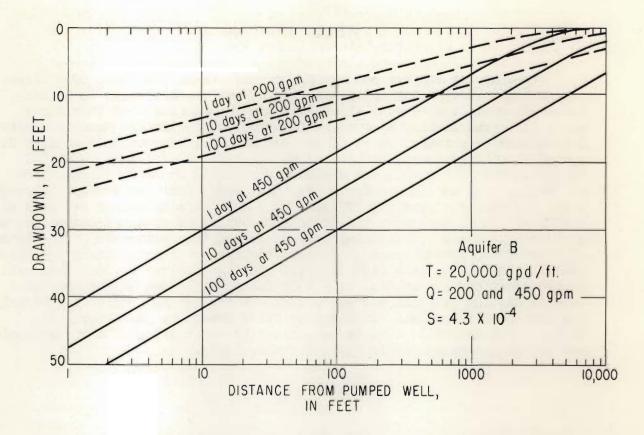


Figure 10.--Calculated distance-drawdown curves determined for pumping from wells finished in aquifer B.

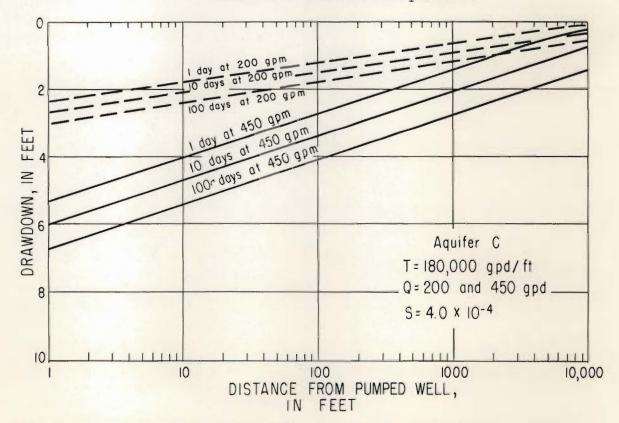


Figure 11. -- Calculated distance-drawdown curves determined for pumping from wells finished in aquifer C.

WELL SPACING

Accurate information about the hydraulic characteristics of aguifers is essential for estimating the spacing of two or more wells to be constructed in the same area. When wells are spaced close together their cones of depression may overlap. Where overlap occurs drawdown is additive. The calculated drawdown half-way between two wells screened in aquifer C, spaced 2,000 feet apart, and pumped at 450 gpm each for 1 day would be about 2.8 feet, or double the drawdown that would occur 1,000 feet from one well pumped at 450 gpm for 1 day (fig. 11). Additive drawdown effects lower the specific capacity of wells, and may cause excessive water-level declines. It is desirable to reduce the overlap of cones of depression as much as possible to insure maximum well and aquifer efficiency. The methods of determining proper spacing of wells tapping artesian aquifers are discussed in detail by Lang (1961). Well spacing involves economics as well as hydrology and hydraulics. Costs of installation and maintenance of interconnecting pipeline and electrical systems may be prohibitive if wells are spaced to completely eliminate additive drawdown. Therefore, the distance between wells must be determined by considering both the economic and hydrologic-hydraulic factors (Bentall, 1963).

QUALITY OF WATER

The slightly acidic rainwater that percolates to the zone of saturation in the area reacts with and dissolves portions of the soil and rock materials. The amount and kind of dissolved mineral matter in the ground water depend upon such factors as the amount and type of organic material in the soil, the kind of rock or soil through which or over which the water moves, and the length of time the water is in contact with the soil and rocks.

Once in solution, the chemical constituents in ground water generally exist as positively and negatively charged ionic particles called cations and anions, respectively. The amount of these constituents in water is determined by chemical analyses, the results of which are reported by weight concentration of each constituent in a million unit weights of water, or parts per million (ppm). Table 2 shows the source or cause and significance of the principal chemical constituents of water reported in chemical analyses.

Nearly 200 water samples were collected from wells that are finished in the major aquifers of the area, and were analyzed by the Quality of Water Branch, U. S. Geological Survey. Results of the analyses are given in table 3 and table 4.

QUALITY OF WATER IN THE AQUIFERS

AQUIFER A

Concentrations of dissolved solids in water from aquifer A are usually below 500 ppm (the maximum amount recommended by the U. S. Public Health Service for public water supplies). The only dissolved mineral matter found in excessive amounts in water from this aquifer were iron and hardnesscausing constituents. Dissolved iron concentrations ranged from 0.12 to 19 ppm and were above 0.3 ppm in 95 percent of the observed cases. Hardness values ranged from 15 to 345 ppm and were in the moderately-hard to very-hard range in about 50 percent of the water samples collected from this aquifer. The hard waters were found where wells are screened in lenticular shell beds that occur in the lower part of this aquifer in the central and eastern part of the area. The water is subject to possible contamination from human and animal wastes, fertilizers, detergents, etc., and unusually high concentrations of nitrate, chloride, or phosphate should be regarded as indicators of such contamination. Determinations of pH were made in the laboratory and average about 7.5. A more acidic and corrosive water is expected in such shallow aquifers, and the near neutral values cited above probably resulted from the escape of carbon dioxide during collection and transportation to the laboratory. A typical chemical analysis of water from aquifer A is shown in table 3.

Water from this aquifer will generally require treatment for the removal of excessive concentrations of iron and/or hardness-causing constituents to make it suitable for most uses.

GROUND-WATER RESOURCES OF BELHAVEN AND VICINITY

Table 2.--Source and significance of dissolved mineral constituents and properties of ground water

	Constituent	Source or cause	Significance
	Silica (SiO ₂)	Dissolved from practically all rocks and soils, usually in small amounts from 1-30 ppm. High concentrations, as much as 100 ppm, generally occur in highly alkaline waters.	Forms hard scale in pipes and boilers. Inhibits the action of zeolite-type water softeners.
	Iron (Fe)	Dissolved from practically all rocks and soils. May also be derived from iron pipes, pumps, and other equipment.	On exposure to air, iron in ground water is oxidized to reddish-brown sediment. More than about 0.3 ppm stains laundry and utensils reddish-brown and is objectionable for food processing, beverages, dyeing, bleaching, ice manufacture, brewing, and other processes. U. S. Public Health Service (1962) drinkingwater standards state that iron concentration should not exceed 0.3 ppm. Larger quantities cause unpleasant taste and favor growth of iron bacteria.
Cations	Manganese (Mn)	Dissolved from some rocks and soils. Not so common as iron. Large quantities often associated with high iron content and with acid waters.	Same objectionable features as iron. Causes dark brown or black stain. U.S.P.H.S. drinking-water standards provide that maganese should not exceed 0.05 ppm.
	Calcium (Ca) and Magnesium (Mg)	Dissolved from practically all soils and rocks, but especially from limestone, dolomite, and gypsum. Calcium and magnesium are found in large quantities in some brines. Magnesium is present in large quantities in sea water.	Cause most of the hardness and scale-forming properties of water; soap consuming. Water low in calcium and magnesium is desired in electroplanting, tanning, dyeing, and in textile manufacturing. These hardness-causing constituents are reported together as equivalent amounts of calcium carbonate (CaCO ₃). Hardness scale used by U. S. Geological Survey is as follows: 0-60 ppm = soft, 61-120 ppm = moderately hard, 121-180 ppm = hard, above 180 ppm = very hard.
	Sodium (Na) and Potassium (K)	Dissolved from practically all rocks and soils. Found also in ancient brines, sea water, some industrial brines, and sewage.	Large amounts, in combination with chloride, give a salty taste. Moderate quantities have little effect on the usefulness of water for most purposes. Sodium salts may cause foaming in steam boilers and a high sodium ratio may limit the use of water for irrigation.
Suc	Bicarbonate (HCO ₃) and Carbonate (CO ₃)	Action of carbon dioxide in water on carbonate rocks such as limestone and dolomite.	Bicarbonate and carbonate cause alkalinity. Bicarbonates of calcium and magnesium decompose in steam boilers and hot water facilities to form scale and release corrosive carbon-dioxide gas. In combination with calcium and magnesium cause carbonate hardness.
Anions	Sulfate (SO ₄)	Dissolved from rocks and soils containing gypsum, iron sulfides, and other sulfur compounds.	Sulfate in water containing calcium forms hard calcium sulfate scale in steam boilers. In large amounts, sulfate in combination with other ions gives bitter taste to water. Some calcium sulfate is considered beneficial in the brewing process. Drinking-water standards recommend that the sulfate content should not exceed 250 ppm.

Table 2.--Source and significance of dissolved mineral constituents and properties of ground water--Continued

Constituent	Source or cause	Significance
Chloride (Cl)	Dissolved from rocks and soils. Present in sewage and found in large amounts in ancient brines, sea water, and industrial brines.	In large amounts in combination with sodium gives salty taste to water. In large quantities increases the corrosiveness of water U.S.P.H.S. drinking-water standards recommend that the chloride content should not exceed 250 ppm.
Fluoride (F)	Dissolved in minute quantities from most rocks and soils.	Fluoride in drinking water reduces the inci- of tooth decay when the water is consumed di ing the period of enamel calcification. Ho- it may cause mottling of the teeth depending on the concentration of fluoride, the age of the child, amount of water consumed, and susceptibility of the individual (Maier, F. 1950). U.S.P.H.S. drinking-water standards recommend that fluoride not exceed 1.7 ppm where 5 year average of daily maximum air temperature is 53.0° to 53.7°F.
Nitrate (NO ₃)	Decaying organic matter, sewage, and nitrates in soil.	Concentrations much greater than the local average may suggest pollution. There is evidence that more than 45 ppm of nitrate (NO ₃) may cause a type of methemoglobinemia in infants, sometimes fatal. Water of high nitrate content should not be used in baby feeding. Nitrate is helpful in reducing intercrystalline cracking of boiler steel, but it encourages growth of algae and other organisms that produce undesirable tastes and odors.
Phosphate (PO ₄)	Dissolved in very small quantities from rocks and soils.	Concentrations much greater than local averages may indicate contamination from phosphate detergents and/or fertilizers.
Dissolved solids	Maybe determined by evaporation	constituents dissolved in a particular water. on-weight or computation methods. U.S.P.H.S. namend that public supplies contain no more
Specific conductance	that is conducted by one cubic index to the total amount of a	°C, of the amount of electrical current c centimeter of the liquid. Can be used as an dissolved solids in a particular water. Gen- onductance by about 0.6 will give a close of dissolved solids.
рН	of water. A pH of 7 is neutra	concentration or an indication of the acidity al, above 7 is alkaline and below 7 is acid. ily much more corrosive than those whth a

Table 3. -- Typical chemical analyses of water from aquifers A, B, C, D, and E

			Aquifers	1	
	A	В	С	D	E
Well Number*	48	190	186	29 - D	29 - E
Silica (SiO ₂)	40	40	48	17	10
Iron (Fe)	3.5	.19	.10	.24	.38
Manganese (Mn)		.02			
Calcium (Ca)	44	53	41	25	145
Magnesium (Mg)	11	16	37	. 26	163
Sodium (Na)	20	83	206	615	4,180
Potassium (K)	2	14	30	40	108
Carbonate (CO3)	0	0	0	0	0
Bicarbonate (HCO3)	231	444	472	488	460
Sulfate (SO ₄)	5	3.6	22	62	598
Chloride (Cl)	15	8.0	221	765	6,720
Fluoride (F)	.2	. 4	.9	1.6	1.3
Nitrate (NO3)	.0	.2	.1	.3	. 4
Phosphate (PO ₄)	.0		.0	.0	.0
Dissolved solids	256	436	838	1,760	12,500
Specific conductance	427	700	1,500	3,140	20,300
Hardness as CaCO3	154	200	257	170	1,030
рН	7.2	7.5	7.9	8.1	7.7

^{*}See figure 13 for location of wells and table 4 for well data.

Note: All constituents, except specific conductance and pH reported in parts per million.

TABLE 4. RECORDS OF WELLS IN THE VICINITY OF BELHAVEN, N. C.

OWNERSHIP: F-Federal Government; M-Municipal; P-Private; S-State agency; N-Corporation.

USE: H-Domestic; P-Public supply; S-Stock supply; I-Industrial; U-Unused.

TYPE OF QW ANALYSIS AVAILABLE: P-Partial; C-Complete.

LOG DATA AVAILABLE: J-Gamma-ray log; Y-Geologist, electric, and gamma-ray logs.

WELL FINISH: O-Open-end; S-Screen; T-Sand point.

WATER-BEARING FORMATION: pM-Post Miocene; My-Yorktown; Mpr-Pungo River; Ech-Castle Hayne limestone; Pb-Beaufort; Kpd-Peedee. AQUIFER: Letter shown is the same as described in this report.

REMARKS: Quality of water reported in parts per million except for specific conductance and pH. Reported well depth noted by *. Diameter of well footnoted by A denotes actual diameter of 1-1/2 inches; B denotes actual diameter of 1-1/4 inches.

Well lo	ocation	Sequence No.	l No.	Owner	ership	analys	data	~	pth cased feet)	Diameter (in Finish	Altitude of LSD (feet)	er level ow LSD feet)	Yield (gallons per minute)	Drawdown (feet)	Specific		Chloride	luoride	a	Hardness	Water- bearing formation	Aquifer	Remarks
Lat	Long	Seq	Wel		Own	M.O.	Log	Depth	Dep (f	Diamet	Alt LSD	Water below (fee	Yie (ga	Dra (Spe	Hď	Ch1	Flu	Iron	Har	Wat bea for	Aqu	
354115N 354111N 354104N 354101N 354030N 354009N	0763847 0763839 0763737 0763722	1 1 1 1 1 1 1	5	E ELLIS	PH	I P I P I P I P	**	*300 *375 *150 *300 260 386	 240	2 0 2 0 2 0 2 0 2 0	14 14 14 8 7	 4 3	10 5 5 15		627 627 700	8.0	20 20 19 30 24 29	0.6 .6 .8 .6 .7	0,10 .17 1.8 .14 .07	190 226 170 212 212 188	Ech Ech My Ech Ech	000 000	
353950N 353934N 353922N 353902N 353846N 353838N	0763640 0763639 0763628 0763640	1 1 1	8 9 10 11	TRINITY BAPT CH J S WINDLEY S C DAVIS UNKNOWN R C COOPER MRS EARL ROSE	PIPI	I P I P I P	**	*280 *280 *292 *300 *285 *308		2 0 A1 0 2 0 A1 0 A1 0 2 0	7 7 5 4 5 4	 c 2	10 10 10		666 676 767	7.5 8.0	26 35 22 21 28 42	.5 .5 .6 .3 .7	.40 .18 3.2 3.9 .13 4.2	208 230 232 392 228 244	Ech Ech Ech Ech Ech	000000	
353818N 353750N 353704N 353611N 353600N 353553N	0763650 0763858 0763932 0763940	1 1 1	14 15 16 17	L H ALLEN W J ALLEN R BISHOP J L BOOMER E R JONES THOMAS DAW	P I P I P I P I	H P H P H P	*	*250 *300 247 15 *275 241	=	2 0 A1 0 2 0 B1 T 2 0 B1 0	3 5 8 9 10 10	8 4 13	=======================================		676 724 87	8.3 8.0 8.2 5.8 7.8 7.5	30 28 37 10 30 30	.5 .6 .6 .3 .7	.20 .12 .30 14 .26 1.0	234 241 266 18 231 234	Ech Ech pM Ech Ech	CCCACC	E
353602N 353604N 353644N 353654N 353848N 353813N	0764005 0764022 0763959 0764006	1 1 1	20 21 22		PH	I I I I P	K	*250 *260 *50 *50 221 218	240 42 42 	B1 0 2 0 2 0 2 0 2 0 2 0 2 0	10 8 10 10 14 13	 10 10	75 50 50 3 5		656	7.7 8.4 8.1	29 11 13	.7	.55 .12 .90	224 275 264	Ech Ech My My Ech Ech	CCBBCC	COMPLEX CO.

TABLE 4. RECORDS OF WELLS IN THE VICINITY OF BELHAVEN, N. C .-- Continued

Well	location	Sequence No.	Well No.	Owner	wnership	OW analysis	Log data	Depth (feet)	Depth cased (feet)	Diameter (in)	1 1	Altitude of LSD (feet)	Water level below LSD (feet)		Drawdown (feet)	Specific	Нq	Chloride	Fluoride	ron	Hardness	Water- bearing formation	Aquifer	Remarks
353822N 353804N 353752N 353737N 353730N		1 1 1 1 1 1	25 26 27 28 29	LEONARD BESS RUTH CREOLE T SLUNVERSON H BENSON USGS USGS	P P P P	H P	Y	22 3 220 *250 82 128	-	B1 36 A1 2 4	0 0	14 10 12 14 12 12 12	10 2 10 7 7	3 4 5 23 21	G 5 8	124 550 579 801	7.9 7.7 8.3	8 9 10 25 24	0.1 .3 .5 .4 .4	8.9 .14 .38 8.5 3.2	33 258 197 316 419	pM pM Ech Ech My My	A A C C B B	BE-T1-66
353730N 353730N 353706N 353700N	0764200 0764200 0764200 0764119 0764112 0764121	1 1 1 1	30 31	USGS USGS USGS G A DUNSHEE WALTER CANNADY UNKNOWN	FPP	U C C C P P I P	Y	382 412 555 168 *80 205	372 402 545 	4 4 2 4	888000	12 12 12 11 12 11	7 10 11 8 3 10	19 20 20 5 	45 26 46 7	1980 3140 20300 868 560	8.1 7.7 8.0	386 765 6720 18 14	1.4 1.6 1.3 .5	.80 .24 .38 14.	173 170 1030 303 210	Ech Pb Kpd My My Ech	C D E B B C	Test well Depth 711 feet
353633N 353613N 353546N 353605N	0764200 0764207 0764308 0764315 0764146 0764135	1 1 1	34 35 36 37	MATTIE RIDDICK	P P P		J	172 181 *84 *260 *280 215	240 	B1 2 2 2	000000	6 8 13 12 12 10	10 12 14	3 3 75 10		888 637 608 729 608	7.8	16 15 19 14 23	.5 .6 .5 .3	2.4 8.7 .36 1.2 .40	172 204 243 250 238	Mpr Mpr My Ech Ech Ech	BCCC	
353611N 353622N 353610N 353557N		1 1 1	40 41 42 43	MRS P WILKINSON MACEDONIA CHURC WALTER CANNADY VERNON DAW COASTAL LUM CO E H BISHOP JR	P P P			*80 214 293 *300 105 *250	60	2 2 2 2	000000	10 10 8 8 8 8	5 12 10 11 4 8	75 10 3 35 4		603 695 656 676	8.2	23 14 21 26 12 81	.3 .4 .4 .7 .6	19. .80 1.1 .58 3.4	398 233 193 223 340 162	My Ech Ech My Ech	BCCCBC	
353550N 353551N 353521N 353509N	0764044 0764033 0764045 0764045 0764032 0764018	1 1 1 1	46 47 48 49	T ALLEN	PIPI	H P P P P P	J	16 20 234 28 *220 23	17 25 	B 1	TOTO	7 7 8 8 8 8	6 6 14 7 5	5 5 1		241	6.5 8.0 7.2 7.7 7.8	12 23 134 54 51	.7	16 .16 19. 6.9	58 212 154 34 78	pM pM Ech pM Ech pM	A A C A C A	
353519N 353458N 353505N 353443N	0764013 0763952 0764018 0764026 0764020 0764003	1 1 1	52 53 54 55	P BROWN	PIPI	P P H P H P		*200 *290 194 *270 *300 *275	250	2 2 2 2	0 0 0	8 7 10 10 9 4	12 	10 50 10		637 500	7.9 8.2 8.2 8.1 8.3	24 34 28 31 31	.7 .88	.28 .27 .09 .10 .53	218 158 197 195 210	Ech Ech Ech Ech Ech Ech	000000	

GUALITY OF

TABLE 4. RECORDS OF WELLS IN THE VICINITY OF BELHAVEN, N. C .-- Continued Ownership Use of water QW analysis per (feet) Water level below LSD (feet) No of (Specific Depth cased (feet) Altitude o LSD (feet) Yield (gallons minute) Water-bearing formation Drawdown (feet) data Diameter Sequence Chloride Fluoride Hardness No. Finish Aquifer Well location Owner Remarks Depth Well Log hd Lat Long PHP 353437N 0764026 1 57 H SCHAVENDER *265 2 0 598 8.2 C 10 30 0.7 162 Ech 0.18 353441N 0764035 1 58 T W SAWYER 114 7.7 PHP *37 --2 8 .76 A 12 22 My .1 PHP PHP 353441N 0764052 1 59 H RUSS *250 --B1 0 9 647 8.1 C 31 .7 .24 184 Ech 231 | 2 0 0 353447N 0764057 1 60 C RADCLIFF *255 9 15 ------C Ech 61 G DANIELS C .20 260 353508N 0764206 1 *200 12 5 651 8.2 11 .5 Ech PHP 20 353505N 0764158 1 62 DAVID LEE B 98 12 5 5 627 8.2 .2 8.0 272 --12 My P H P P P P P H P -- B1 O .50 353458N 0764152 1 63 J T SLADE *250 11 618 8.4 14 . 5 222 Ech C 353448N 0764148 1 598 7.8 64 CHURCH 80 -120 7 5 5 10 .2 4.8 300 My B 353425N 0764113 1 65 C J CARAWAN 29 26 B1 T 8 5 989 7.7 64 .2 89 A 11 Mg PHP 353408N 0764159 66 G BELLMAN *240 20 22 C 1 10 676 8.5 . 5 .42 234 Ech 1 353353N 0764157 67 MATTHEW LILLY PHP 206 20 9 C 16 695 8.2 27 .7 224 ---5 .44 Ech 353347N 0764158 1 68 W CRADDACK PHP 20 C *250 10 695 8.2 24 .7 .54 213 Ech ------2 0 2 0 PHP 40 353326N 0764208 1 69 ROBERT GASKINS *250 10 704 8.0 30 .7 .40 237 C Ech 353240N 0764222 1 PHP 704 8.5 70 STEVE HARRIS *200 C 10 32 .6 .42 242 Ech P H P P H P 353237N 0764212 1 71 S H DAVENPORT *300 A1 0 10 10 724 8.2 30 .6 .24 255 Ech C 353222N 0764219 1 72 J CUTHRELL C *185 --20 10 320 7.4 43 .3 .14 84 Ech PHP 20 353245N 0764427 1 73 JESSE KEECH 733 8.4 C *195 ---11 ----48 . 5 1.7 231 Ech 353228N 0764416 1 74 J GADSKINS PHP C *260 -- B1 O 10 __ 811 8.0 53 .6 .28 245 Ech -P H P P S P H P P H P 353226N 0764404 1 75 C CLARK 20 801 8.0 .6 .09 C *185 10 50 246 Ech 353213N 0764348 1 76 RALEIGH KEECH 7 C *180 A1 0 10 820 8.0 52 .06 242 Ech , 6 353203N 0764342 1 77 IDA KEECH 22 B1 T 135 8.1 A 6 25 5 14 .1 7.4 38 pM 7 C 353158N 0764357 1 78 C ATWOOD JR *297 20 10 801 8.1 250 --52 .6 .13 Ech --353151N 0764352 1 79 S W JAVESON PHP *20 17 B1 T 5 156 7.5 9 .90 39 A .6 Mq ~-4 C 353148N 0764324 1 80 H LATHAM PHP *210 -- B1 0 8 811 8.1 51 .6 .06 246 Ech PHP 353149N 0764311 1 C 81 W GAYLORD *220 -- A1 O 10 811 8.1 54 .18 249 Ech .6 353114N 0764257 1 82 GEORGE WINDLEY PHP 25 22 B1 T 133 7.9 10 9.0 A 4 13 .1 55 Mg PHPPUPU 353107N 0764243 1 -- A1 O 10 C 83 J HESSE *250 8 830 8.4 58 .21 Ech --.7 236 16 B1 T 353100N 0764237 1 84 UNKNOWN 8 2 502 7.6 5.7 19 3 110 . 1 18 pM AC 353051N 0764228 1 85 ROBERT WINFIELD *195 -- | 2 0 9 5 1130 8.5 164 .7 .22 233 Ech 353006N 0764225 1 86 R WINFIELD 181 -- A1 O 6 18 --Mpr --353018N 0764200 1 87 R WINFIELD PSPJ *200 2 0 7 1180 7.9 150 .56 220 C 4 1.1 Ech 2 0 2 0 353027N 0764156 1 88 R WINFIELD PHP *300 8 1720 8.2 312 1.0 .12 271 C --Ech -----PHP PHP 353040N 0764157 1 89 LISA BLAND *180 955 8.5 C 8 8 8 101 1.0 .05 274 Ech 353040N 0764150 1 90 R WINFIELD *150 B B -- A1 O 422 8.1 8 .2 1.7 216 My 353038N 0764135 1 91 DENNIS GIBBS 93 -- | 2 0 2 5 492 8.0 8 .1 2.2 236 My A1 0 353031N 0764134 1 92 G ROOSEVELT PHP 5 B *65 508 8.1 2.3 13 260 My

TABLE 4	1.	RECORDS	OF	WELLS	IN	THE	VICINITY	OF	BELHAVEN,	N.	C Continued
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Well I	Long	Sequence No.	Well No.	Owner	Ownership	QW analysis	Log data	Depth (feet)	Depth cased (feet)	Diameter(1n)	Finish	Altitude of LSD (feet)	Water level below LSD (feet)	Yield (gallons per minute)	Drawdown (feet)	Specific	Hď	Chloride	Fluoride	Iron	Hardness	Water- bearing formation	Aquifer	Remarks
353035N 353036N 353020N 353040N	0764058	1 1 1 1	93 94 95 96 97	RANDOLPH MOORE L M EBOR JOE EBOR P SPENCER L EBOR C MARSLENDER	P H P H P H P H P H	P P P		*130 40 *100 *237 33 *260	37 30 	2 B1	O T O O	8 5 5 3 5 5	 4 4	5 4 3 10		507 262	8.0 8.2 8.3	16 14 22 108 5 114	0.2 .1 .1 .7 .1	4.3 3.3 11 .08 5.2 .13	243 110 166 258 160 244	My My My My pM Ech	B B G A C	
353047N 353100N 353113N 353154N	0764007 0763952 0764053	1 1 1	100 101 102 103		P H P H P H P H P H	P P P		*255 *200 22 *260 192 *225	20 	2 B1 2 2	0 0 T 0 0	6 7 4 5 6	2 14	3 10 5 10		1010 965 106 1270 1010 647	7.9 7.3 8.2	119 96 10 198 139 25	.8 .7 .1 .7 .9	.10 .10 22 .01 .38 5.5	260 274 26 259 208 287	Ech Ech pM Ech Ech	CCACCC	
53208N 53213N 53143N 53136N	0764114 0764120 0764017 0764008	1 1 1	106 107 108 109	C D ALLIGOOD C D ALLIGOOD J H GADSTON ALICE BESS M MIDGETT HUBS RECK	PHPHPHPHPPPP	P P P		*267 *105 *190 *300 *380 70	 	2 2 2	0 0	5 6 6 6 6	15 4	10 5 10 5 		695 811 1150 907		91 29 74 176 75 17	.7 .3 .7 .7 1.0	.26 5.6 .10 .12 .08 2.8	260 307 216 198 279 194	Ech My Ech Ech Ech My	СВСССВ	
53138N 53146N 53151N 53308N	0763931 0763915 0763905 0763823	1 1 1	112 113 114 115	C SMITH W EDWARDS	PHPHPHPH	P P P		*75 *260 *30 *30 *87		B1 B1	OTTO	5 6 5 5 5 5	 3	10 10 5 		1120 218 232 627	7.6 8.3 7.3 7.5 7.5 6.0	46 165 45 56 10 56	.1 .7 .1 .1 .4	4.6 .13 8.6 10 2.2 .78	102 243 56 36 282 56	My Ech pM pM My pM	B C A A B	
53323N 53330N 53338N 53353N	0763821 0763839 0763840 0763841	1 1 1 1	118 119 120 121		P H P H P H P H P H	PPPPPPPPPPPPPPPPPPPPPPPPPPPPPPPPPPPPPPP		*30 *380 6 73 17 18		A1 2 24 B1 B1 B1	000 T	5 6 3 5 5 3	2 4 3 2	5 		685 294 285	6.8 7.6 6.0 6.8 7.8	8 57 30 30 27	.1 .8 .2 .0 .3	.50 .14 .95 1.1 1.4	15 206 48 29 311	pM Ech pM My pM pM	A C A B A	
353438N 353455N 353500N 353506N	0763843 0763854 0763900 0763859	1 1 1	124 125 126 127	E S WHITLEY D TAYLOR A D JOHNSON PANTEGO PROVISI L WHITLEY D W ALLEN	P H P H P H P H P H	P		20 *275 256 230 23 *300	18 20	2 2 2 1	T 0 0 0 T 0	5 4 7 8 8	3 8 7 4	10 3 		110	7.9 7.9 6.1 8.1	37 38 10 43	.6 .7 .0 .7	.10 .10 5.0 .28	237 243 22 175	pM Ech Ech Ech pM Ech	A C C C A C	

Well :	location	Sequence No.	II No.	Owner	Ownership	Use of water	data	1 10	Depth cased (feet)	Diameter(in) Finish	Altitude of LSD (feet)	Water level below LSD (feet)	Yield (gallons per minute)	Drawdown (feet)	Specific conductance		Chloride	Fluoride	uc	Hardness	Water- bearing formation	Aquifer	Remarks
Lat	Long	Sec	Well		OWI	N O	Log	Del	Del	Dig	A1.	₩a. be.	Yie (gg	Dra	Spe	Hd	Ch	Flu	Iron	На	Wa beg fo	Aqu	
3440N 3446N 3418N 3423N	0763714 0763703 0763605 0763554	1 1 1 1	130 131 132 133	H T SPENCER G C SPENCER H T SPENCER RONALD LANIER RONALD LANIER DOUGLAS SPENCER	P P P	H II H II S II H II		*325 *275 98 21 25 26	19 23	2 0 A1 0 B1 0 B1 T B1 T	10 10 11 10 11 10	5 4 5 5			555 332 492	8.2 7.7 7.2 6.8 7.1 7.7	79 66 5 12 22 29	0.7 .8 .2 .2 .2 .3	0.12 .87 16 5.0 11 14	188 241 344 150 239 146	Ech Ech My pM pM pM	C C B A A	
3405N 3427N 3356N 3355N	0763522 0763504 0763202 0763144	1 1 1 1	136 137 138 139		PPP	H II H II H II H II S	J	*165 250 82 79 105 *350	=	A1 2 0 2 0 A1 T B1 0 2 0	11 5 11 9 9	9 8 8 5 6	10 10 5 5		579 695	8.5 9.0 7.7 8.1 8.0	37 68 9 12 14	.6 1.2 .1 .4 .3	.03 19 4.8 .09 .06	234 50 266 170 214	My Ech My My My Ech	B C B B C	
3418N 3420N 3413N 3405N	0763029 0763017 0763007 0763007	1 1 1 1	142 143 144 145	ETHEL BLUNT DOTTIE MALSCH S KEIFER H C HARRIS DOTTIE MALSCH R L MARTIN JR	PPP	H II H II H II H II		32 122 *100 *300 4 102	=	B1 0 2 0 2 0 36 0 A1 0	4 4 3 4 3	2 1 2 0	5 5 3 5		262 917 868 878 917	8.1 7.9 8.3	7 29 29 30 30	.4 .2 .2 .4 	1.1 1.2 1.4 4.2 3.9	121 410 284 298 345	pM My My Ech pM My	A B B C A B	
53411N 53317N 53313N 53325N	0763208 0763218 0763229	1 1 1 1	148 149 150 151	E GRIFFIN M JONES FRANK BILLUPS FRANK BILLUPS HARRY PETERSON MAGNEBAR MINING	P P P	H F H F H F		92 120 25 *100 *150 280	22	B1 0 B1 0 B1 T 2 0 2 0 10 0	3 4 5 5 6 0	1 1 1 6	4		936 897 253 618 618	8.3 7.7 8.0	32 29 10 13 13	.5 .3 .2 .3 .3	4.2 1.1 .36 .40 4.8	389 102 129 328	My My pM My My Ech	B B A B C	
3328N 3329N 3331N 3340N	0763258 0763311 0763323 0763413	1 1 1	154 155 156 157	F O DAVIS LULA CARY CARTER ADAMS M NIXON L M DILDAY CLARA DILDAY	PPPP	H F H F H F S F		*325 25 *150 *150 95 87		2 0 B1 T A1 0 2 0 2 0 2 0	8 8 8 7 4 4	0 2 2	 10 3		589 656	8.3 8.2 8.2	216 13 13 13 16 14	1.1 .1 .3 .3 .2 .2	.10 4.4 5.2 3.5 3.8 3.0	204 140 307 302 345 334	Ech pM My My My My	C A B B B	
53327N 53330N 53329N 53326N	0763447 0763504 0763522	1 1 1 1	160 161 162 163	M ARMSTRONG MRS PINNEFORD J PINNEFORD J DAVIS MARY EBOR MARION HARRIS	PPPP	H F H F H F		96 24 18 53 79 *100	15	2 0 B1 T B1 T B1 0 B1 0	3 4 4 4 4	2 1 2 4	5 5 5		378	8.1 8.2 7.4	16 20 2 11 8 7	.1 .1 .3 .4 .3	3.4 11.4 .12 3.3 3.0 1.9	133 164 28 345 328 308	My pM pM My My	B A A B B	

	ocation	Sequence No.	ell No.	Owner	Ownership	Use of water	Log data	Depth (feet)	Depth cased (feet)	Diameter (in)	Finish	Altitude of LSD (feet)	Water level below LSD (feet)	Yield (gallons per minute)	Drawdown (feet)	Specific	Hd	Chloride	Fluoride	Iron	Hardness	Water- bearing formation	Aquifer	Remarks
Lat	Long	S	*		0		3 17	Q .	Ω	A	124	4 H	≥ D	A 0	-	0 0	Ω,			-				
353240N	0763606 0763600 0763552 0763600	1 1 1 1 1 1	169	PURNELL BARBER IRVING WHITE LLOYD HUDNAL J E MCKINNEY REGINALD BISHOP D N ROUSE	P P P	H II H II H II S II H II		79 23 *100 *360 240 *105	21 	B1 B1 A1 2 2 2	O T O O O O	4 4 4 5 4	2 2 13 9	3 10 		618 160 647 1050 1110 618	8.1 8.2 8.3 8.2	12 28 10 117 101 28	0.3 .1 .3 .8 1.2 .4	11 5.1 3.5 .22 11 .50	316 36 324 276 158 322	My pM My Ech Ech My	B A B C C B	
353237N 353217N 353212N 353210N	0763657 0763655 0763653 0763645	1 1 1	172 173 174 175	WALTER JOHNSON AXOOM SMITH CTY OF BELHAVEN CTY OF BELHAVEN A B CUTHRELL CTY OF BELHAVEN	P M M	H II H II P II H II P II		230 *220 *275 265 *275 28	250 245 25	B1	0	4 4 5 4 4	9 11	10 150 50 5		965	8.0	1050 14 220 154 84 70	.8 .3 1.0 .1 .7	1.8 1.0 .18 4.1 	674 338 262 200 211 98	Ech Ech Ech Ech Ech	CCCCCA	
353210N 353214N 353215N 353231N	0763703 0763708 0763725 0763727	1 1 1	178 179 180 181	CTY OF BELHAVEN CTY OF BELHAVEN CTY OF BELHAVEN JACK LEE CTY OF BELHAVEN CTY OF BELHAVEN	M M P M	P I P I H I	P	131 223 221 133 265 255		B1 2 2 2 2 2 2	000000	4 4 4 4 4	2 11 5 11 11	5 5 4		1180 811 840	7.6 7.4 7.9 8.2 	116 260 136 26 117	.3 .0 1.0 .4 .5	37 26 18 .06 	102 378 105 205 54	My Ech Ech My Ech Ech	BCCBCC	
353253N 353253N 353253N 353343N	0763746 0763747 0763748 0763718	1 1 1 2	184 185 186 187	CTY OF BELHAVEN CTY OF BELHAVEN CTY OF BELHAVEN CTY OF BELHAVEN N C WATER RES N C WATER RES	M M M S	P C P C U I		*275 *275 *275 *275 *275 *340 *370	250 250 250 250 250 335 365	4 4 8 4	000000	5 5 5 5 4 4	=======================================	150 150 150 300 		1300 1500 1300 1500 2900 3000	7.6	186 225 185 221 820	=======================================	.03 .01 .04 .10	226 238 206 257	Ech Ech Ech Ech Ech	000000	Analysis 4/29/65 " " " " " " " " " " " " " " " " " " "
353343N 353343N	0763718 0763718	1 3	188 189	N C WATER RES N C WATER RES N C WATER RES N C WATER RES	S	U	PY	*467 24 124 150	462 114 100	2 A1	O T S O	4 4 4 4	4 4 3	20 75	16	7800 396 700	7.7	3540 20 33	=	.08 5.4 .17	615 171 171	Ech pM My My	C A B B	Casing set at 298

AQUIFER B

In nearly all cases, concentrations of dissolved solids in water from aquifer B were below 500 ppm and averaged about 380 ppm. However, hardness-causing constituents, dissolved iron, and hydrogen-sulfide gas were found in objectionable amounts in the majority of the water samples collected from this aquifer. Hardness ranged from 29 to 419 ppm and was above 180 ppm throughout most of the area. Concentrations of dissolved iron ranged from 0.03 to 37 ppm and were above 0.3 ppm in 85 percent of the samples analyzed for this constituent. The pH values of the water were above 7.0 in most cases. A typical chemical analysis of water from aquifer B is shown in table 3.

Generally, treatment for the removal of excessive hardness, iron, and hydrogen-sulfide gas will be required to make this water suitable for domestic use.

AQUIFER C

The concentration of dissolved solids in water from aguifer C ranged from 150 to 4,680 ppm and averaged about 575 ppm. Excessive amounts of iron, chloride, hardness-causing constituents, and hydrogen-sulfide gas were encountered in various places throughout the area. Concentrations of dissolved iron ranged from 0.01 to 19 ppm and were above 0.3 ppm in about 50 percent of the observed cases. Hardness ranged from 34 to 674 ppm and averaged 234 ppm. Ninety percent of the water samples collected from wells finished in this aquifer contained more than 180 ppm hardness. Chloride content ranged from 9 to 3,540 ppm. Concentrations of chloride are generally below 250 ppm in water in the upper part of aquifer C. However, concentrations of this anion are very near 250 ppm in the extreme southern part of the area, and have equaled or exceeded this limit at various times in water from the municipal wells at Belhaven. Figure 12 shows the approximate distribution of chloride concentrations in water contained by the upper part of this aquifer. The lower part (below about 280-300 feet below land surface) is saturated with brackish and saline water. Observed pH values of water samples collected from aquifer C averaged 8.1 and were above 7.0 in all cases. A typical analysis of water from the upper part of aquifer C is shown in table 3.

Where concentrations of chloride are within acceptable limits, the removal of hardness-causing constituents and excessive concentrations of iron and hydrogen sulfide will make water from this aquifer suitable for most uses.

OTHER AQUIFERS

The waters in the deeper artesian aquifers of the area contain large amounts of dissolved solids (up to 12,500 ppm) mainly due to excessively high concentrations of sodium and chloride. Observed chloride content ranged up to 6,720 ppm and was well above 250 ppm in all cases. In addition, excessive concentrations of iron, sulfate, and hardness-causing constituents are common, and there is no economical way to make this water suitable for domestic use. Table 3 shows a typical chemical analysis of water from aquifers D and E.

FACTORS AFFECTING FUTURE USE AND DEVELOPMENT

There are several factors that must be considered when planning the development of ground-water supplies from the various aquifers in the area of investigation. The most important factors are the lowering of water levels and the change in chemical quality of the waters as a result of pumping.

Large quantities of water are pumped from aquifer C, about 15 miles southwest of the area of investigation, to allow the open-pit mining of phosphate deposits that occur in the overlying Pungo River Formation. The influence of this concentrated pumpage has spread for some 30 miles from the center of pumping. Within the area of influence, the withdrawals have reduced the artesian pressure, caused water levels to decline, and induced the movement of water from all directions toward the center of pumping. To date this pumping has caused water levels in aquifer C to decline about 12 feet in the vicinity of Belhaven. Data collected during this investigation indicate that there is little immediate threat of lateral salt-water encroachment in the upper part of aquifer C in the Belhaven area under the present hydrologic conditions. In fact, it appears that fresher water in the upper part of aquifer C in the north and northeastern part of the area of study is moving toward Belhaven as a result of pumping in the mine area (fig. 12).

However, the lower part of aquifer C, in this area, contains water having chloride concentrations in excess of 250 ppm. In response to the lowered water levels, this salt water is probably moving slowly towards the upper part of the aquifer. Under the present hydrologic conditions it is possible that the entire aquifer in the area may eventually become contaminated from this source.

Aquifers A and B are separated from aquifer C by layers of dolomitic limestone and clay in the Pungo River Formation and by silt and clay layers found near the base of the Yorktown Formation. Aquifer D is separated from aquifer C, in the western part of the area, by a layer of tight silty clay about 10 to 15 feet in thickness. This clay layer thins towards the southeast and pinches out between Pantego and Belhaven where aquifers C and D become one hydrologic unit. Some leakage of ground water from aquifers A and B to aquifer C occurs due to higher water levels in the shallower aquifers. However, because of ample recharge, the water levels in the shallower aquifers are essentially the same as they were prior to the decline of water levels in aquifer C. Ground-water withdrawals from aquifers A and B are well distributed and relatively small throughout the area of study. Thus, these shallow aquifers, particularly the more productive aquifer B, should be considered as good alternative sources of ground water if salt-water contamination prohibits the use of aquifer C.

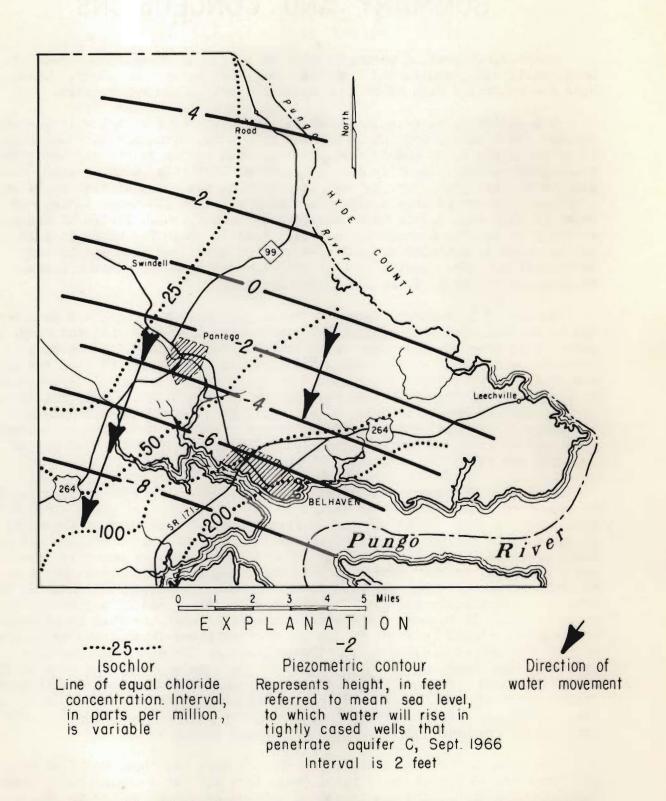


Figure 12.--Map showing the approximate distribution of chloride concentrations, water level, and water movement in the upper part of aquifer C, 1966.

SUMMARY AND CONCLUSIONS

About 2,500 feet of unconsolidated to partially consolidated beds of sand, silt, clay, shell, and limestone underlie the area of study. These beds are saturated with water from basement rock to the water table.

The sediments between land surface and about 300 feet below land surface contain fresh water, and can be divided into three permeable water-bearing zones (aquifers A, B, and C) that are separated by two relatively impermeable zones (aquicludes). Aquifer A is composed of lenticular beds of sand, silt, clay, and shell that occur between land surface and about 35 feet below land surface. The top of this aquifer is coincident with the water table, and water in this zone occurs under non-artesian conditions. Yields of wells screened in aquifer A generally are less than 10 gpm. The water in this aquifer usually contains excessive amounts of dissolved iron and is very hard locally. Other dissolved constituents are below the maximum amounts recommended by the U. S. Public Health Service.

Aquifer B is composed of about 70 feet of permeable shell and sand beds, and it is separated from aquifer A by lenticular layers of silt and clay. The water in aquifer B is confined under artesian conditions. The top of this zone occurs at about 40 feet below sea level in the western part of the area, and it slopes about 5 feet per mile toward the east. Theoretically, yields of properly constructed 10-inch diameter wells screening the full thickness of this aquifer should be about 10 gpm per foot of water-level drawdown in the wells at the end of one day's continuous pumping. Excessive concentrations of dissolved iron, hardness-causing constituents and hydrogen-sulfide gas commonly are found in the water in this aquifer.

Aquifer C is composed of shell limestone, and calcareous sand beds. It is separated from aquifer B by layers of silt and clay that range from 60 to more than 100 feet thick from the western to the eastern part of the area, respectively. Water in aquifer C is confined beneath this aquiclude and is under artesian conditions. The top of aquifer C occurs at about 160 feet below sea level in the western part of the area, and increases in depth at the rate of about 9 feet per mile toward the east. Average thickness of this zone is approximately 190 feet. Theoretically, the yield of a properly constructed well, 10-inches in diameter, screened through the full thickness of this aguifer should be about 80 gpm per foot of water-level drawdown in the well at the end of one day of continuous pumping. Water in aquifer C is generally very hard, and in many places it contains excessive amounts of dissolved iron and hydrogen-sulfide gas. Water in the lower part of this aquifer (below about 300 feet below msl), contains concentrations of chloride in excess of 250 ppm. Deeper artesian aquifers were penetrated during the test drilling in the area and were found to contain saline water.

The natural hydrologic conditions in the area have been modified by large and concentrated ground-water withdrawals from aquifer C during recent years. Water-level declines and the threat of salt-water contamination in aquifer C (the major artesian aquifer in the area), have concerned all those who use this aquifer for water supply. If ground-water withdrawals from aquifer C intensify these problems, aquifers A and B afford good alternative sources of ground water.

BASIC DATA

The wells listed in table 4 were inventoried during the investigation and are representative of the types, depth, yields, etc., of wells found in the vicinity of Belhaven. Figure 13 shows the location of the inventoried wells.

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Figure 13.--Map showing inventoried wells in the vicinity of Belhaven, N. C.