

Quality of Groundwater in Eastern Osaka, Japan, and its Hydrogeological Interpretation

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Quality of Groundwater in Eastern Osaka, Japan, and its Hydrogeological Interpretation*

Michiji TSURUMAKI

(With 4 Tables and 10 Figures)

Introduction

The area studied is located in the east of geographic center of Osaka Prefecture, and is bounded on the west by Uemachi Upland, having the Yodo River as a northern border, and limited on the east by the Ikoma Range, with an edge of the Yamato River on the south (Fig. 1). In the area, called the Eastern Osaka, land subsidence has been remarkably increasing during a few recent years. Owing to the parallelism between the progress of the subsidence and the rapid increase of wells for industrial and municipal water supply, the use of groundwater being gradually regulated and substituted for the use of surface water. Under the circumstances, it is pushed on to throw light upon the hydrogeological conditions of the Eastern Osaka.

This report is based on the work supported by a research grant from Osaka Prefectural Government, and started in 1965 and extended to 1969. The work has been concentrated principally on water quality problems, with secondary attention given on hydrogeology of the area. The primary purpose of this report is to elucidate the distribution pattern of the main constituents in water, and then to illustrate the chemical character of water in connection with the occurrence of it. However, a general sketch on the geology of the area is included in order to give the background necessary for some interpretation of the water quality.

The measurement of natural tritium (^3H) content in well water can give a valuable information about groundwater hydrology (BEGEMAN & LIBBY, 1957). In recent years, the term "tritium hydrology" is mentioned in literature (GRANT-TAYLER & TAYLER, 1967). The use of tritium dating was carried out for the study of groundwater hydrology in Kanto Plain in this country (OCHIAI, 1968). Since 1967, in this research, the measurement of natural tritium in well water has been put into practice by courtesy of Dr. T. OCHIAI at the laboratory of Agricultural Engineering Research Station. A fuller discussion of the chemical data gathered during the study will be presented later to this Journal after obtaining the data of tritium dating.

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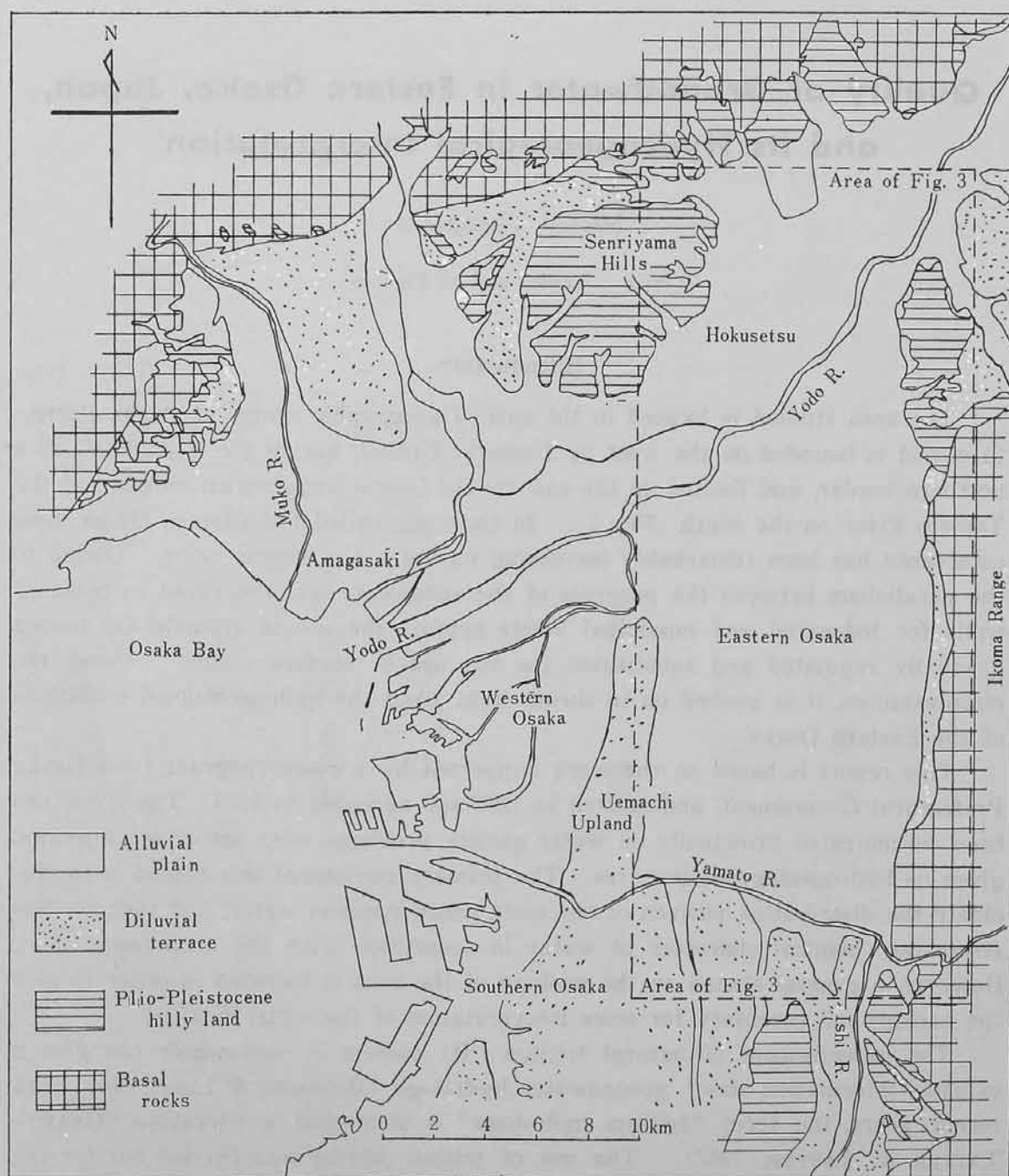


Fig. 1. Geological map (after IKEBE and TAKENAKA, 1969; modified).

General Sketch of Geology

Geology of the Osaka basin and the surroundings is summarized as the following table (IKEBE & TAKENAKA, 1969), and the simplified geologic map is shown in Fig. 1.

- Recent Alluvium (Umeda formation)
- Upper Pleistocene formation { Younger (Lower) terrace
Middle terrace
Older (Higher) terrace
- Osaka group (Late Pliocene–Early Pleistocene) { Upper part
Lower part
Lowest part
- Infra-Osaka group (Pliocene)
- Kobe group (Miocene)
- Pre-Neogene basement

The concept of a groundwater basin has become important for leading basin wide planning and development of groundwater (TODD, 1959). It is evident from the geological map that the Osaka basin consists a groundwater basin, but Uemachi Upland which lies in the middle part of the basin divides it into two areas,

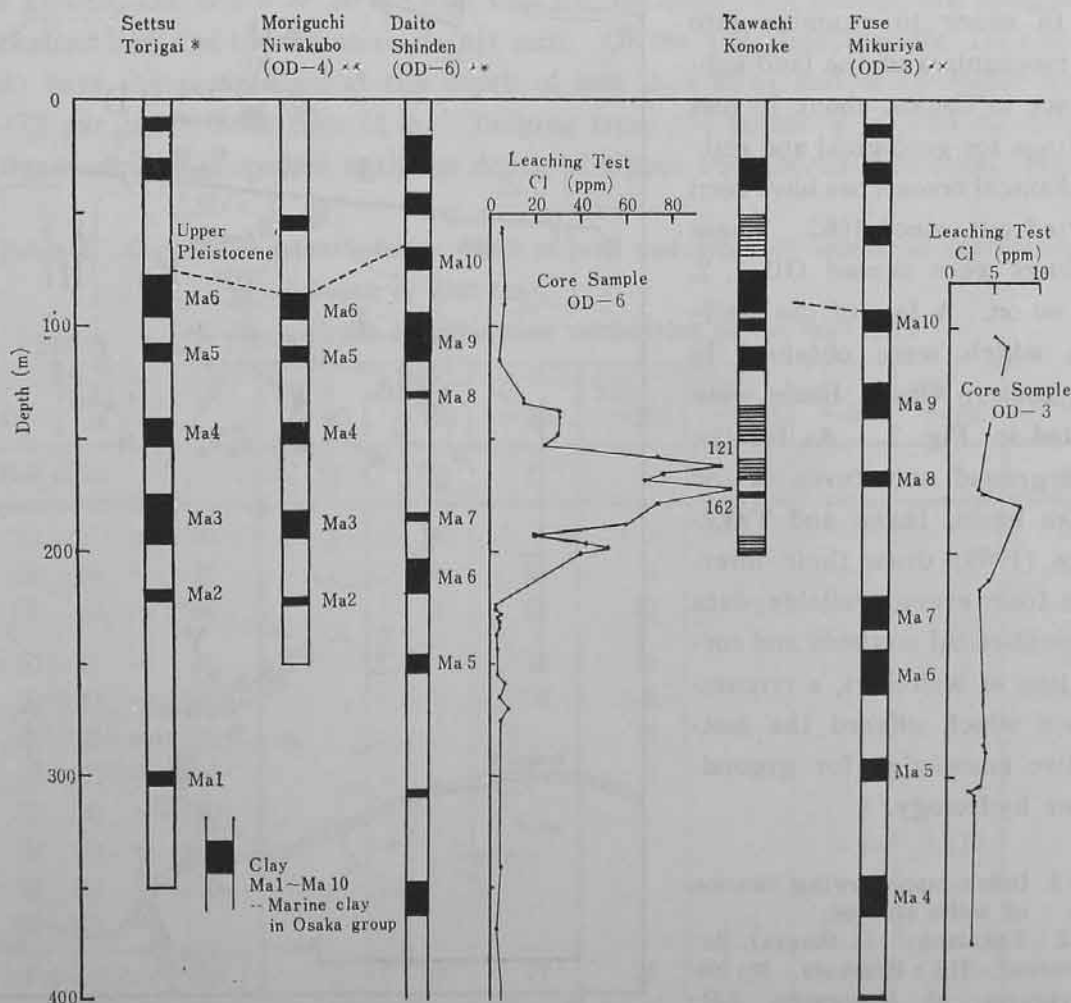


Fig. 2. Well logs and results of leaching test.

* after INAI and ISHIDA (1966)

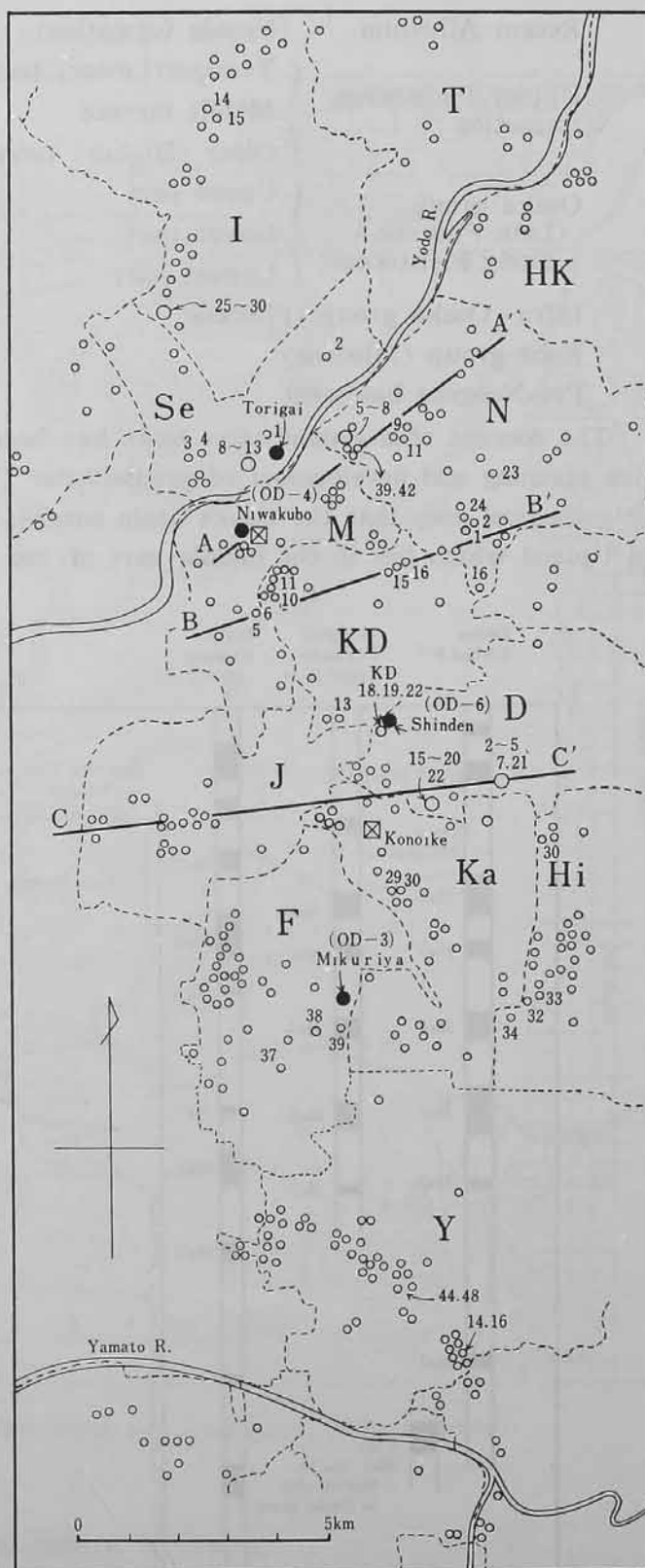
** after MATSUSHITA (1960, 1967)

that is to say, Western Osaka and Eastern Osaka. Owing to the concept of the groundwater basin, this research comprises the Hokusetsu area extending to north of the Yodo River, and the combined area of the Hokusetsu area and the Eastern Osaka is designated as the Eastern Osaka Basin. The chemical quality of groundwater in the Western Osaka was discussed by the author with special reference to the intrusion of sea water (TSURUMAKI, 1962).

In order to examine into the mechanism of the land subsidence in Osaka, about 10 test drillings for geological and soil-mechanical researches have been carried out, since 1963. These drillings were named OD-1, 2, and so on. A few of the well-logs which were obtained in the Eastern Osaka basin were quoted in Fig. 2. As for the underground structures of the Osaka basin, IKEBE and TAKENAKA (1969) draw their inference from every available data of geophysical methods and correlation of well-logs, a circumstance which offered the instructive knowledge for groundwater hydrology.

Fig. 3. Index map showing location of wells studied.

T: Takatsuki, I: Ibaraki, Se: Settsu, HK: Hirakata, N: Neyagawa, M: Moriguchi, KD: Kadoma, J: Joto, D: Daito, Hi: Hiraoka, Ka: Kawachi, F: Fuse, Y: Yao,



- 15 ○ Well sampled and number
- ⊗ Test well
- Deep drilling for geological investigation
- A A' Line of chloride profile (Fig. 9)

Conditions of Well

Fig. 3 shows the location of the well studied. To each of the wells a number is assigned at every district. In this text, each well is called by the number with the combination of an initial of the name of district; for example, N-10 and KD-20. The water-samples were taken from the wells which are deep, high-capacity and constructed by drilling. Most of the deep wells in this area have well screens or other opening in the casing at several different depths, accordingly, determination of the depth zone at which water can enter the well is a troublesome work. As an example, the depth of well and the perforated portion of the well casing are shown in Fig. 9.

On the whole of wells studied in the Eastern Osaka, the relation between the depth of well and the depth of the uppermost perforated portion was examined and the result is shown in table 1. Out of total of 188 wells, 84 wells lack for their casing records. For the convenience of examination, the wells are divided into two groups, the depth of 20 to 150m and 151 to 350m; the former are designated as *shallow well* and the latter as *deeper well*. Of the 132 *shallow wells*, 117 (89 per cent) have the perforation at the depth of less than 60 m, and of 72 *deeper wells*, 53 (73 per cent) more than 61 m. Judging from the tendency of well completion of this area, and compared with the depth of Upper Pleistocene formation (Fig. 2),

Table 1. Correlation table between depth of well and depth of uppermost perforation

A.....Depth of well (m)

B.....Depth of uppermost perforation of the well (m)

B \ A	20 ~ 50	51 ~ 100	101 ~ 150	151 ~ 200	201 ~ 250	251 ~ 300	301 ~ 350	Total
Not clear	13	40	21	7	3			84
① ~ 20	3	16	6	1				26
② 21 ~ 40	1	28	28	4				61
③ 41 ~ 60		13	22	12	2			49
④ 61 ~ 80			12	13	3		1	29
⑤ 81 ~ 100			3	9	4			16
⑥ 101 ~ 120				5	1			6
⑦ 121 ~ 140				1	6			7
⑧ 141 ~ 160						2		2
⑨ 161 ~ 180						2	1	3
⑩ 181 ~ 200					2			1
⑪ 201 ~					1	1	1	3
Total ① ~ ⑪	4	57	71	45	19	5	3	288
Total ① ~ ③	117			19				
Total ④ ~ ⑪	15			53				

it may be said that the *shallow wells* are supplied from the aquifers of the Upper Pleistocene, and *deeper wells* from those of Upper and/or Lower Part of Osaka group.

Methods for Collection and Analysis of Water-sample

A. Collection of water-sample.

The major parts of the water-samples were collected from the wells whose records of construction had been preserved. In the region not having proper distribution of wells, the samples were taken from the wells even lacking the records. The sample was taken from such a well as was able to be pumped up directly and not through pressure tank or any other distribution systems, because the water was in artesian condition. After the measurements in field, water-sample was collected in a polyethylen bottle. All bottles were washed with hot hydrochloric acid and rinsed with tap water in the laboratory, and they were rinsed again with the water to be sampled.

B. Field determinations.

Temperature, pH and electrical conductivity were determined simultaneously with sampling, and the addition of the reagents necessary for colorimetric determination of ferrous ion was performed.

(1) Temperature: Temperature measurement was made with mercury thermometer graduated per 0.2 °C.

(2) pH: A pH determination was made colorimetrically through addition of bromthymol blue or phenol red solution as indicator in comparison with standard color series.

(3) Electrical conductivity: A preliminary step in each analysis was the determination of electrical conductivity in micromhos ($\mu\Omega$) with a portable conductivity meter, model CM-3M fabricated in TOA Electronics Ltd.. Accuracy and sensitivity of the meter are $\pm 10\%$ and 2.5% respectively. The electrical conductivity is related to the concentration of ionized substance in water; therefore, a conductivity reading gives the analyst an idea as to the concentration of ions, and this speeds up the analytical procedures. Also the conductivity in micromhos at 18 °C was used as a check on the chemical determinations (LOGAN, 1961. TSURUMAKI, 1969).

C. Laboratory tests.

Alkalinity, nitrite, phosphate and ferrous ion were determined on the same day of collection, and then others in succession within a week. Analytical procedures were based mainly on "Testing Methods for Industrial Water (JIS K 0101)" published by Japanese Standards Association and partly on "Standard Methods (1965)" published by American Public Health Assoc. and American Water Works Assoc..

(1) Alkalinity: Titrated with 0.02N sulfuric acid in use of mixed bromcresol green-methyl red indicator. The value is considered to represent the amount of bicarbonate because of the absence of P. alkalinity in all of the water-samples.

(2) Chloride: Volumetric methods using mercuric nitrate and silver nitrate were adopted to meet the concentration of chloride.

(3) Sulfate: Turbidimetric method and colorimetric barium chromate method were adopted.

(4) Nitrate: Photometrically determined in use of phenoldisulfonic acid, excepting the specimen with higher content of chloride. For the specimen of high chloride, brucine method (AWWA, p 198, 1965) or 2, 4- xyleneol method (NAGASAWA, 1966) were used tentatively.

(5) Nitrite: Photometrically determined in use of G. R. reagent.

(6) Phosphate: Colorimetrically determined through stannous chloride method.

(7) Ammonia: After distillation colorimetrically determined by nesslerization.

(8) Calcium and Magnesium: EDTA titrimetric methods were adopted.

(9) Sodium and potassium: The measurements were performed with atomic absorption and flame emission spectrophotometer, model AA-1E fabricated in Nippon Jarrell-Ash Co., Ltd..

(10) Silica (soluble): Colorimetrically determined by molybdosilicate method.

(11) Ferrous iron: Ferrous ion was determined colorimetrically with the specimen treated in the field using o-phenanthroline method.

(12) Total iron: Colorimetrically determined through thiocyanate method after oxidizing the specimen.

(13) Manganese: Colorimetrically determined through persulfate method.

D. Expression of result.

The result of chemical determination was expressed in parts per million (ppm) and in terms of equivalents per million (epm). Epm can be obtained from ppm by dividing the ppm by the equivalent weight of the ion. As regard silica, the expression by epm was not adopted, for the silica in most waters is present as the particles of subcolloidal size (Hem, 1959, p. 56).

When a complete analysis was made, the totals of anions and cations expressed in epm should be equal. Difference between them can be used to check for the possible error in analytical procedures. Table 2 shows the results of analysis for the representative water-samples, as well as the totals of anions and cations, and the ratios of some important constituents.

In order to estimate the range of variation in the chemical quality, the water-samples of two groups of wells were taken and analyzed repeatedly from 1966 to 1969 once or twice in a year. One group locates at the western Neyagawa and consists of four *shallow wells* (N-5, 6, 7, 8) and one *deeper well* (N-39), and the other group locates at the southwest Daito and consists of five *deeper wells* (D-16, 17, 18, 19, 20).

Regional Distribution of Water Quality

Fig. 4 and 5 are the maps showing the regional distribution of the quality of water-samples taken from the *shallow wells*. There are two reasons why we adopted the *shallow wells* as the examination of regional distribution. (1) The depth zones from which water can enter the *shallow wells* are in the limited range of depth as seen in Table 1. (2) The range of seasonal or annual variation of the quality of the *shallow well* is smaller than that of *deeper well* as seen from the comparison of the analyses of N-5, 6, 7, 8 and D-16, 17, 18, 19 (Table 2-3, 2-6 and 2-7).

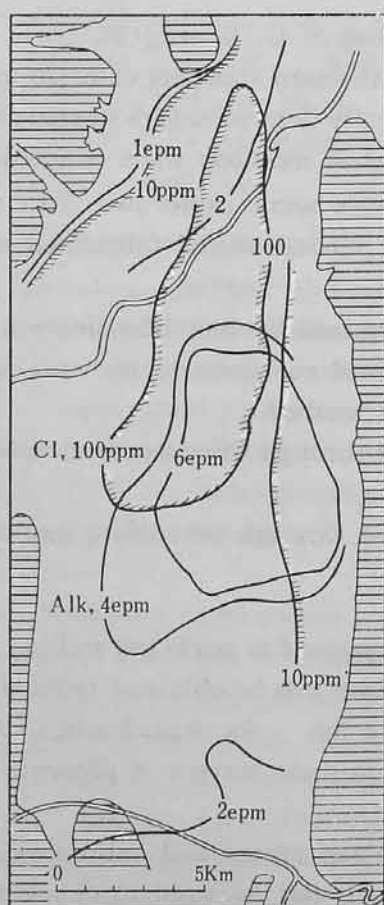


Fig. 4. Generalized alkalinity and chloride map of the *shallow well*.

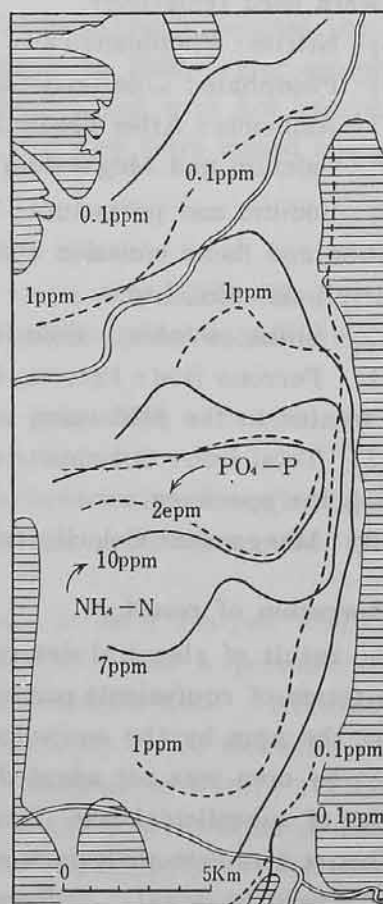


Fig. 5. Generalized ammonia and phosphate map of the *shallow well*.

Although not enough data are available for detailed maps, there are enough so that the pattern can be drawn with some confidence. The main feature of the pattern is a gradual increase in each of the constituents from low values in the surrounding hilly lands to high values in the area of central low land.

(1) Alkalinity (Fig. 4): In the southern area, low alkalinity line of 2 epm stretches from the junction of the Yamato and Ishi rivers to the northwest. The

pattern suggests that the river-bed water of the rivers is supplying the groundwater to the Upper Pleistocene formations of this area. The Yamato River, the downstream of which was excavated in recent times, had been flowing to the north in former times. As for the northern area, such type of replenishment of groundwater as is inferred in the southern area has not been defined. A few words may be said about the circumstance: (1) The alkalinity map in the northern area and the low content of natural tritium (ref. Table 2-10, N-50 m and N-100 m) make it possible to deny the replenishment. (2) Low silica content, for example, of 25-30 ppm (ref. Table 2-3, Group-B) suggests the likelihood of the replenishment, on the contrary. The analyses of surface waters of the Yodo, Yamato, and Ishi rivers are shown in Table 2-11.

There are pronounced high values of alkalinity (4~6 epm) in the central regions of this area. A marked tendency to increase alkalinity with groundwater flow is well accepted, and the alkalinity which shows the concentration of bicarbonate ion is produced by the decomposition of organic matter in ground (FOSTER, 1950; SUGISAKI, 1962). Accordingly, the groundwater flow in those regions must be very slow or stagnant. It seems that the pumping up of such groundwater plays one reason of a severe ground subsidence in this area.

(2) Chloride (Fig. 4): The zone of high chloride water shown by the isochlor pattern of 100 ppm lies along north to south trend in the northern part of central Eastern Osaka. The higher values of chloride appeared within the isochlor are ranged from 500 to 800 ppm. Chloride in this area is not supposed to be depend on the sea water intrusion because of the following reasons. (1) High chloride zone does not connected with the western area. (2) Chloride does not increase year after year. (3) Groundwater with high chloride sometimes occurs in a deeper stratum. (4) Alteration of water quality due to the contamination of sea water recognized with the groundwater in Western Osaka (TSURUMAKI, 1962) is not seen in this area. (5) Contents of natural tritium in the high chloride groundwater in this area are usually low (1.9, 2.7, 4.0 in T.U. unit). As a result, it can be said that the source of chloride in this area is connate sea water. Chloride may be present in resistates as the result of inclusion of connate water, and is to be expected in any incompletely leached deposit laid down under the sea or in a closed basin where chloride was present (HEM, 1959, p.104). Accordingly, the distribution of chloride in groundwater may offer some interesting explanation for the underground structure of the basin. More detailed consideration of the chloride distribution will be dealt presently in the later chapter.

(3) Ammonium and phosphate (Fig. 5): The general pattern of the two ions is that of a progressive increase from the marginal regions to the central region. Nitrogen and phosphorus are regarded as to be derived from decomposition of organic matter contained in the marine clayey stratum. Therefore, these maps represent the distribution of the younger marine formations in this area.

Distribution of Chloride with Depth

Of the various ions analyzed, chloride has proved the most useful for three reasons. (1) From geochemical point of view, chloride ion is not complicated by exchange reaction or by precipitation from solution. (2) From geological point of view, chloride has interesting meaning as described above. (3) Water containing much chloride is poor for all sides of utilization.

A. Introductory research.

(1) Relations among depth, temperature, and chloride.

In order to investigate chloride distribution with depth, it is necessary to determine the depth zone in which water enters the well. The relationships between the depth zone which was identified from the casing record and water temperature, the temperature and chloride content of the water-sample taken from the well were examined for five regions in this area, and then the general features of chloride profiles were drawn in 1967 (TSURUMAKI).

(2) Electric log.

Electric log is a useful method for locating salt-water leaks in water wells. Resistivity logs were performed at two localities with the different electrode spacings, one of which is nearly equal to the diameter of the bore hole and the other is three times of it. The results are shown in Fig. 6. The log of western Neyagawa indicates that the salty water locates at the depth of 60 m, and southern Kawachi at 150~200 m.

(3) Leaching test of core-sample.

The boring core samples taken by the double core tube and carefully stored must preserve the pore water at the depth of sample. Extraction and analysis of chloride ion in the sandy samples obtained from the test holes named OD-3 and OD-6 were performed with the following procedure. Sample in wet state was treated with 0.5N potassium nitrate solution of the same weight for 2 hours at a room temperature. After standing with several times of stirring, the suspension was filtered off, and the filtrate was submitted for analysis. Colorimetric method using mercuric nitrate and VOLHARD'S titrimetric method were adopted in accordance with concentration. The results were expressed in ppm of chloride of the filtrate, and shown in Fig. 2. OD-6 had the samples contained high chloride at the depth of

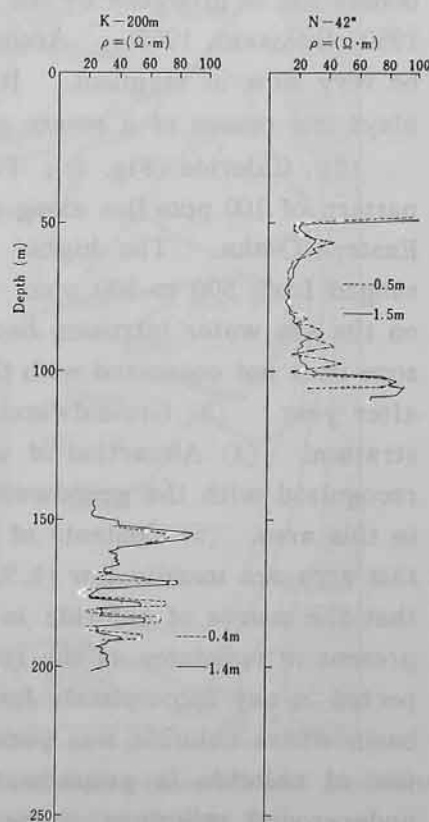


Fig. 6. Resistivity logs.
* measured by Nippon
Sakusen Tanko Co.,

150 to 180 m, whereas none of the samples of high chloride appeared in OD-3.

(4) Variation of chloride with time.

Seasonal or annual variations of the water quality observed at the well group of southwest Daito suggested the chloride variation with depth. Fig. 7 shows the variation of chloride and water temperature. The variation are often observed after the repair of well by swabbing. In the case of the wells of D-16, 17, 18, and 19, increment of chloride is accompanied by rising temperature. In contrast, well of D-20 shows decrement of chloride in close accordance with rising temperature. These wells have screens at several different depths ranging from 60 to 290 m as shown in line-C of Fig. 9, and so, the well water is the mixture of groundwaters of different depths, and the ratios of mixing are variable in every well and at an interval of time. The study of the relations makes it possible to allocate the waters of high and low chloride to the three zones as following; the shallow zone is low, the middle zone is high, and the deeper zone is low in chloride content.

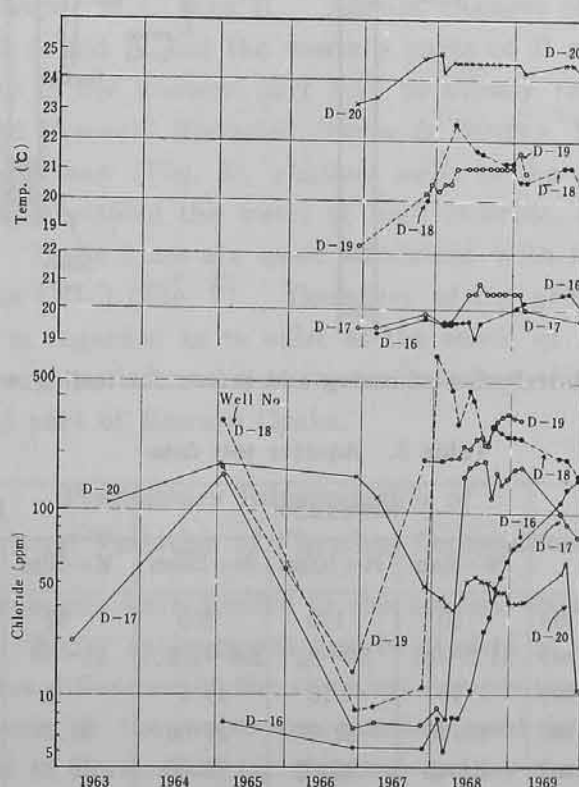


Fig. 7. Variation of chloride and temperature with time, at Daito well group.

(5) Test well.

The results mentioned above have formed a fundamental guiding data for solving the chloride distribution with depth, but do not clarify the actual chloride concentration of groundwater which lies in ground. They reveal only the comparative values of each of the zones. In order to ensure the chemical quality and the

hydraulic characteristic of the aquifer individually, two groups of test wells which were constructed so that water could enter from a specified aquifer were established at Niwakubo and Konoike, and they were named N-50m, -100m, -250m, and K-50m, -100m, -200m. Driller's well logs and casing records are illustrated in Fig. 2 and 8, and the results of analysis and pumping test are shown in Table 2-10, and 3, respectively. Niwakubo test wells located in northern Moriguchi may be looked upon as representative of the northern part of this area, and Konoike test wells

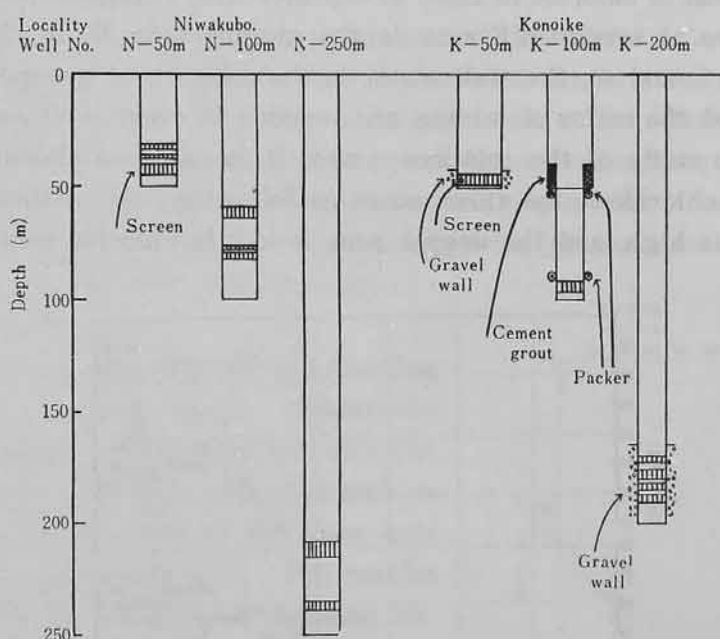


Fig. 8. Installation of casing and screen for testing wells.

Table 3. Aquifer test data

Locality	Niwakubo			Konoike		
Well No.	N-50m	N-100m	N-250m	K-50m	K-100m	K-200m
Total Depth (m)	50	100	250	51	100	200
Depth of Perforation (m)	31.5~45	59~82	208~238.5	45~50	92~97	175~193
Length of Perforation (m)	10	10	11.5	5	5	13
Static Level (m)	30.46	32.07	27.06	33.37	41.72	42.70
Pumping Level (m)	34.40	35.02	30.05	40.53	48.31	45.05
Specific Capacity (m ³ /day/m)	150	200	170	100	30	300
Coefficient of Transmissibility						
{ Drawdown (m ² /sec)	3.1×10^{-3}	5.4×10^{-3}	5.9×10^{-3}	8×10^{-4}	3.8×10^{-4}	1.3×10^{-2}
{ Recovery	2.9×10^{-3}	8.3×10^{-3}	5.1×10^{-3}	5.6×10^{-4}	3.4×10^{-4}	8.8×10^{-3}
Coefficient of Permeability (m/sec)	3.0×10^{-4}	6.8×10^{-4}	4.8×10^{-4}	1.1×10^{-4}	7.2×10^{-5}	8.5×10^{-4}
Date of Test	Dec. 1967			Mar. 1968		

as that of the central part of it. The N-50m and -100m wells at Niwakubo and the K-200 m well at Konoike yielded the water of high chloride, concentration of which were ranged from 160 to 210 ppm.

B. Profile of chloride.

Fig. 9 shows three profiles of chloride content in groundwater.

(1) Line-A is a profile along the south of the Yodo River. The groundwater of high chloride prevails at the depth zone of 40 to 80m. The zone coincides with the Uper Pleistocene formations distributed in this region as recognized from Fig. 2. At the northeastern part of this line, high chloride water is not seen in the shallow zone, but the deeper well (N-31) whose record of casing is not clear, yields the high chloride.

(2) Line-B shows the profile along Keihan railway line.

(3) Line-C is the profile along Katamachi railway line.

There are some similarities between the profiles of B and C, but the depth of high chloride zone is deeper in C than B. Abrupt changes of chloride are seen at the eastern parts of A and B, and the western parts of B and C. The discontinuity of water quality in the western part may be closely related with the geological structure called Uemachi Elevation (IKEBE & HUZITA, 1966).

As shown in chloride map (Fig. 6), *shallow wells* in the southern regions of Eastern Osaka have never yielded the water of high chloride, and this holds true for the *deeper wells*. Those facts are quite consistent with the leaching test of core sample taken from OD-3 (Fig. 2). Boundary of the groundwater bodies of low and high chloride is regarded as to exist at the south of line-C. Owing to the discontinuity, it would be inferred that the tectonic line of northeast trend lies water of in the central part of Eastern Osaka.

Preliminary Interpretation of Regional Variation of Chemical Composition

The foregoing discussions were limited to the spacial distribution of selected constituents in water. Water analyses can also be studied in various ways to demonstrate similarity and difference of the chemical composition. In this chapter the dominant ions present in the water are examined and interpreted using the most prevailing method of illustration for chemical-quality data.

Trilinear plotting system proposed by PIPER (1944) is one of the efficient methods which enable to illustrate a large number of chemical analyses. The trilinear diagram illustrates the various percentages of anions and cations in the two triangular fields and a combined position of all major ions in the diamond-shaped field. Percentages of anions and cations are based on total equivalents per million of the major ions. In the PIPER's original diagram, the diamond-shaped field indicates the character of water as represented by the relationships among $\text{Na} + \text{K}$, $\text{Ca} + \text{Mg}$, $\text{CO}_3 + \text{HCO}_3$, and $\text{Cl} + \text{SO}_4$ ions. The waters in this area, however, contain

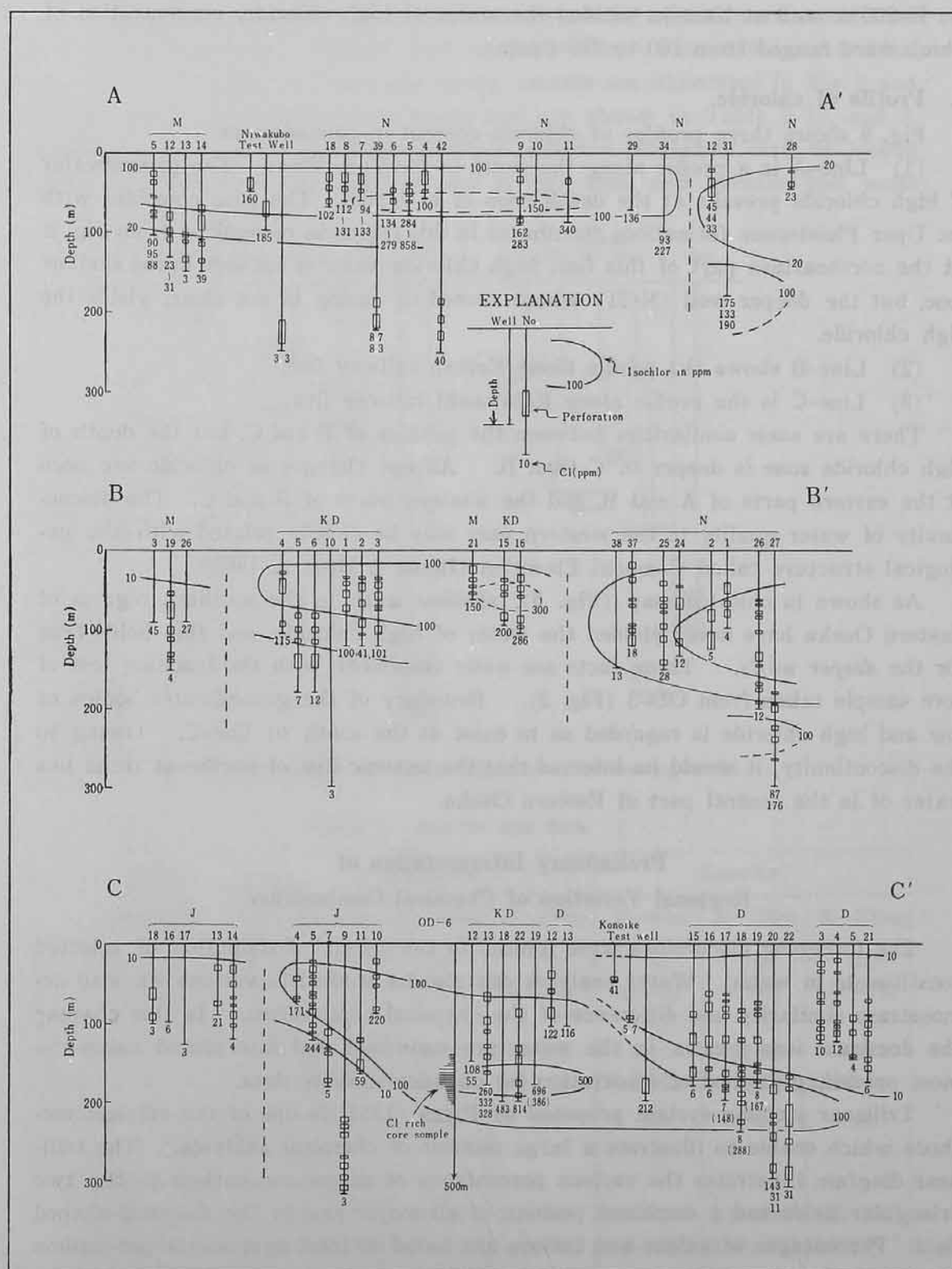
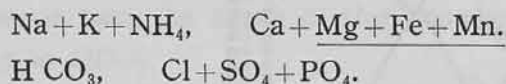


Fig. 9. Profiles of chloride concentration. Location of profiles is shown on Fig. 3.

considerable amounts of phosphate, ammonia, iron and manganese, and these ions should not be overlooked for calculating the total concentration. In this report the relationships are represented by the combination of;



Carbonate is not determined in most waters in this area. The combinations of ions underlined are used for the cation and anion triangular fields. The arrangement of constituents mentioned above is a matter of convenience, and has no severe geochemical meaning. The arrangements of Na, Ca and Mg (SUGISAKI, 1962), and $\text{Fe}^{2+} + \text{NH}_4^+$, $\text{Ca}^{2+} + \text{Mg}^{2+}$ and $\text{Na}^+ + \text{K}^+$ (KONISHI, 1966) are found in literatures. In the diamond-shaped field concentrations of dissolved matter are usually represented by the circles, the radius of which are proportional to the concentrations. This method of illustration requires considerable space for plotting a single one water-sample, and is inadequate for plotting a number of the analyses of similar composition. Each of the diagrams shown in Fig. 10 contain from 15 to 20 water analyses. The concentrations are indicated by epm unit near by the diagram.

Fig. 10 illustrates the representative water analyses which are obtained from the northern area of Eastern Osaka basin. The water-analysis data are classified into three regions and five groups.

(1) Analyses of waters taken from Hokusetsu area are shown in Diagram-A. Along the foot zone of northern hilly land, both of *shallow* and *deeper wells* yield the waters which are low in total mineral content. The waters are high in the ratio of bivalent cations ($\text{Ca} + \text{Mg} + \dots$) to total cations, and high in sulfate content except for *deeper wells*. This type of water quality is close agreement with that the hills and the foot zone may be physiographically taken as the intake area of groundwater for this area and, perhaps, partly in Eastern Osaka. In the southern part of this area, Settsu, the concentration of total soluble matters varies widely than the above region. Of them, the waters of high chloride may be accounted for the transitional type between the above type and the type of next group. The waters of low chloride in this region are high in the ratio of monovalent cations ($\text{Na} + \text{K} + \text{NH}_4$) to total cations, which fact means that the waters are soft. Sulfate is scarce in both of the high and low chloride waters and this fact is attributed to the reduction of sulfate accompanied by the inactive flow or stagnation of groundwater (FOSTER, 1950; CEDERSTROM, 1946).

(2) The *shallow wells* in northern Moriguchi and western Neyagawa yield the water of high chloride with a range of 100 to 300 ppm. Water analyses of this region are shown in Diagram-B. Dominant constituents of the water are chloride and sodium.

(3) The *deeper wells* in the same region as above and the *shallow well* in the southern Neyagawa yield the water of low chloride and low alkalinity. The difference in the water quality between them is the cationic composition as seen in Diagram-C. The waters of *deeper well* which are conceived to be drawn from

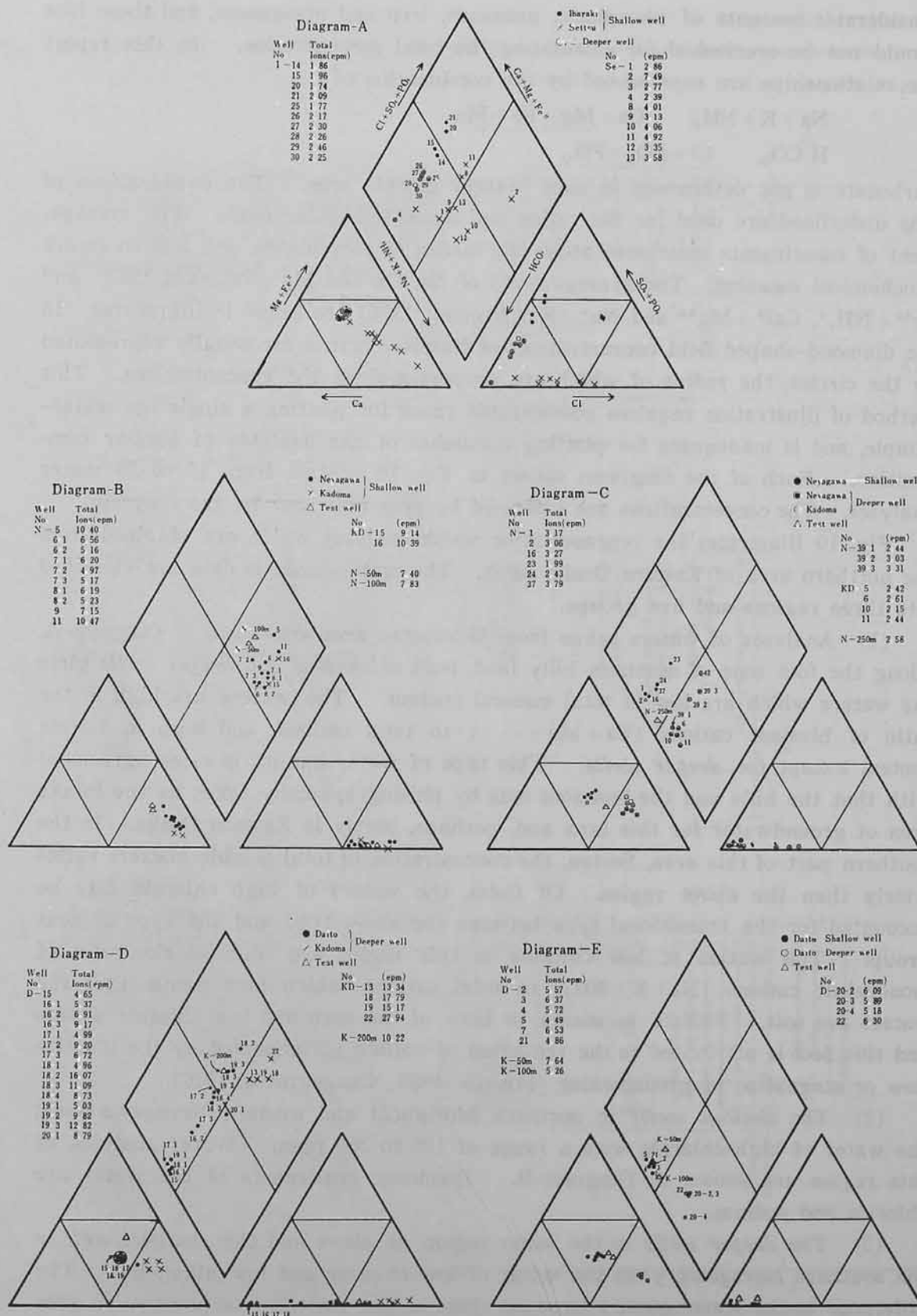


Fig. 10. Analyses represented by three points plotted in trilinear diagram.

the aquifers correlated to the strata of the Lower Part of Osaka group are soft by comparison with the shallow waters. As for the correlation, it is mentioned that the depth of the uppermost perforation of the wells of N-250m, N-39 and KD-10 is deeper than the depth of the boundary between the Upper and Lower Osaka group at OD-4.

(4) Diagram-D illustrates the chemical analyses of the water samples obtained from the *deeper wells* in the region of southern Kadoma and eastern Daito. The chloride concentration in the water samples of D-16~20 varies widely with time as already shown (Fig. 9), and in the diamond-shaped upper field the analyses plot on nearly a straight line. When analyses represent two original waters and their mixtures, the analyses will plot on a straight line. On the basis of such informations, it appears well established that the waters are the mixtures of two kinds of groundwaters; namely shallow water of high alkalinity and deeper water of high chloride. In the cationic triangular diagram the analyses plot on a limited field, therefore, those waters are assumed to be similar in cationic composition. The waters taken from the wells of D-16~20 have higher content of divalent cations than those from the wells of KD-18, 19 and 20, which fact means that the former is harder than the latter. As for the waters of high chloride in this region, it may be said that the deeper the aquifer lies from north to south, the harder the water becomes.

(5) The analyses of six *shallow wells* and two *deeper wells* which have the depth of 300m are illustrated in Diagram-E. All the waters are high in alkalinity

Table 4. Summary of water quality

Occurrence			Water Quality		
Region	Group*	Sub-group	Total Ions (epm)	Predominant Ions	
				Anions	Cations
Hokusetsu	A	Ibaraki	1.8~ 2.5	HCO ₃ +SO ₄	Ca (+Mg+Fe)
		Settsu	2.4~ 7.5	HCO ₃ or Cl	Na
North of Eastern Osaka	B	shallow	5.0~10.5	Cl	Na
	C	shallow	2.0~ 3.8	HCO ₃	Ca+Mg
		deeper†	2.2~ 3.6	HCO ₃	Na
	D	deeper	5.0~18	Cl or Cl \rightleftharpoons HCO ₃	Ca+Mg
Middle of Eastern Osaka	E	shallow	4.5~ 7.6	HCO ₃	Ca+Mg
		deeper††	5.2~ 6.3	HCO ₃	Na

* Group name refers to Table 2 (Analytical table) and diagram in Fig. 10.

† The deeper wells are in the same region of group-B.

†† The depth of wells are deeper than the wells of group-D.

and low in chloride. The waters taken from *shallow wells* are harder than those from *deeper wells*, and the fact was seen in Diagram-C also. As for the waters of low chloride, there is a tendency that the hardness in water is lessing with the increase in the depth of aquifer.

Thus the variations in chemical quality derived from the differences in the location and the depth of wells were summarized in Table 4.

Summary and Conclusion

Groundwater samples taken from the wells in the Hokusetsu and the Eastern Osaka area have been analyzed since 1966.

Water analyses of the shallow wells the depths of which are less than 150 m are illustrated by the maps showing the regional distribution of alkalinity, chloride, ammonia and phosphate. These maps indicate the career of groundwater as follows; (1) Alkalinity map suggests that the river-bed water of the Yamato and Ishi rivers is supplying the artesian groundwater to the Upper Pleistocene formations in the southern region of Eastern Osaka. (2) Chloride map indicates that the groundwater of the central part of the area is contaminated with connate saline water. (3) Ammonia and phosphate, and alkalinity maps show that the groundwater in the central region is confined by the younger marine formation.

On the basis of the several preliminary reseaches, the profiles of chloride concentration in the groundwater were compiled from the water analyses of the shallow- and deeper wells in the northern regions of the area. The profiles and the chloride map indicate markedly the discontinuities of water quality, and this fact suggests the existence of two tectonic lines. One which lies in the western region and runs parallel to Uemachi upland is connected with the tectonic movement which produced the upland, and the other lies in the central region, running across the basin from northeast to southwest. The latter is merely a matter of surmise, and is not yet connected with any of the tectonic movement. Such an underground structure as this must be restraining the flow of groundwater.

The dominant ions in the water were examined using the trilinear diagram which is wellknown as PIPER's key diagram and which is extensively used for the comparison of water analyses. As a result, it has led to the clarification of characters of groundwater due to the difference of occurrence, such as region and depth, as follows; (1) Along the foot zone of the Senriyama hills the shallow wells and deeper wells yield the waters characterized by low alkalinity and high sulfate. (2) The waters of high chloride increase their hardness ($\text{Ca} + \text{Mg}$) with depth of aquifer. (3) The waters of low chloride increase their sodium and potassium concentrations with depth.

There remains the problem of how to interpret these results geochemically and hydrologically. This article is a preliminary report and a fuller account of the data will be discussed together with the result of tritium measurement.

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In this research, field and laboratory works were carried out through the co-operation by the author and Mr. M. NAGASAWA of Osaka Prefectural University, and each one had his own allotted regions or wells for the research. The analyses accomplished by NAGASAWA are quoted in Table 2 with the notation of "M.N.". His contribution to this research has been very great and the author wishes to express deepest appreciation to him. The author is also very grateful to Dr. T. OCHIAI who is executing the tritium dating.

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Table 2-1. Chemical analyses of groundwaters from Hokusetsu area (1)

Group	A		A		A		A		A		A		A		A		A		A	
Well No.	1-14		1-15		1-20		1-21		1-25		1-26		1-27		1-28		1-29		1-30	
Depth of Perforation(m)	47~116		26~137.5		40~109.5		30~111		50~144		110~172		114~186.5		185~234		201~240		190~276	
Total Depth (m)	130		140		120		120		154.5		182		188.5		235.5		250		293	
Date of Collection	Mar. 12, '68		Mar. 12, '68		Mar. 15, '68		Mar. 15, '68		Mar. 13, '69		Mar. 13, '69		Mar. 13, '69		Mar. 13, '69		Mar. 13, '69		Mar. 13, '69	
Temp. (°C)	16.8		16.8		16.1		17.4		16.1		16.1		16.2		17.2		17.2		17.3	
EC. ($\mu\text{U} \cdot 18^\circ\text{C}$)	152		167		148		173		137		164		159		162		175		163	
pH	6.3		6.2		6.1		6.2		6.3		6.4		6.7		6.6		6.6		6.6	
HCO ₃	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
Cl	63.6	1.042	64.8	1.062	47.3	0.776	53.1	0.870	70.5	1.156	88.6	1.452	96.4	1.580	103.2	1.692	105.5	1.730	104.2	1.708
SO ₄	3.69	0.104	3.19	0.090	4.41	0.124	4.96	0.140	4.58	0.129	3.69	0.104	6.77	0.141	4.20	0.119	4.34	0.122	4.20	0.118
NO ₃ -N	33.0	0.687	39.3	0.818	40.4	0.842	52.5	1.092	22.5	0.468	29.0	0.695	28.0	0.583	21.5	0.448	29.5	0.615	20.5	0.427
PO ₄ -P	0.45	0.032	0.051	0.004	0.034	0.002	0.02	0.001	0.045	0.003	0.04	0.003	0.045	0.003	0.045	0.003	0.050	0.004	0	0
Total Anions(A)	0.134	0.013	0.056	0.005	0.075	0.007	0.015	0.001	0.098	0.009	0.082	0.008	0.125	0.012	0.060	0.006	0.095	0.009	0.10	0.010
		1.878		1.979		1.751		2.104		1.765		2.172		2.319		2.268		2.480		2.263
NH ₄ -N	0.078	0.005	0.012	0.001	0.023	0.002	0.039	0.003	0.14	0.010	0.13	0.009	0.15	0.011	0	0	0.093	0.007	0.135	0.010
Na	9.7	0.422	9.4	0.408	8.6	0.374	9.3	0.403	9.7	0.421	10.8	0.470	11.8	0.513	11.7	0.508	12.8	0.557	11.7	0.508
K	3.42	0.08	3.55	0.091	2.58	0.066	2.83	0.073	2.95	0.075	3.13	0.080	2.90	0.074	2.76	0.071	2.59	0.066	2.78	0.071
Ca	12.08	0.604	13.55	0.677	11.48	0.574	14.50	0.725	9.67	0.483	12.90	0.643	13.13	0.656	14.53	0.723	15.30	0.763	14.08	0.703
Mg	6.72	0.552	7.82	0.645	6.14	0.506	6.93	0.571	6.58	0.542	7.85	0.646	7.85	0.646	7.32	0.602	8.03	0.660	7.13	0.587
Fe II	4.08	0.146	3.00	0.107	5.18	0.185	8.10	0.290	5.90	0.221	7.88	0.282	9.34	0.335	8.42	0.301	9.46	0.336	8.60	0.308
Mn	0.71	0.026	0.62	0.023	0.35	0.012	0.45	0.016	0.65	0.023	1.08	0.039	1.28	0.047	1.26	0.045	1.28	0.047	1.13	0.041
Total Cations(C)		.843		1.952		1.719		2.081		1.775		2.169		2.282		2.252		2.436		2.228
SiO ₂	66.8		61.5		52.2		47.8		58.2		60.0		53.6		59.4		56.2		61.0	
(A)/(C)		1.02		1.015		1.02		1.01		0.995		1.00		1.02		1.01		1.02		1.01
Ca/Na		1.43		1.66		1.54		1.80		1.15		1.37		1.28		1.425		1.37		1.38
K/Na		0.21		0.22		0.18		0.18		0.18		0.17		0.14		0.14		0.12		0.14
Mg/Ca		0.91		0.95		0.88		0.79		1.12		1.005		0.985		0.83		0.865		0.835
Analyst	M. Tsurumaki		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.	

Table 2-2. Chemical analyses of groundwaters from Hokusetsu area (2)

Group	A		A		A		A		A		A		A		A		A		A	
Well No.	Se-1		Se-2		Se-3		Se-4		Se-8		Se-9		Se-10		Se-11		Se-12		Se-13	
Depth of Perforation(m)	57~143		44~134		56.5~118.5		78~208.5		21~130		22.5~130.5		34~101		49.5~142		38.5~123.5		53~167	
Total Depth(m)	150		150		120		230		140		135		106		150		130		180	
Date of Collection	Mar. 1, '68		Mar. 1, '68		Mar. 1, '68		Mar. 1, '68		Apr. 3, '68		Apr. 3, '68		Apr. 3, '68		Apr. 3, '68		Apr. 3, '68		Apr. 3, '68	
Temp. (°C)	16.9		16.4		16.9		18.0		18.3		18.5		18.3		17.3		17.0		17.6	
EC. (μS , 18°C)	335		690		222		187		365		275		355		460		290		315	
pH	6.8		7.1		6.7		6.7		7.0		7.0		7.2		6.8		7.2		7.2	
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
HCO ₃	125.1	2.050	141.3	2.317	137.3	2.251	136.8	2.243	123.0	2.016	124.9	2.047	152.6	2.501	128.2	2.102	144.3	2.366	133.1	2.182
Cl	27.42	0.772	174	4.91	14.37	0.406	3.30	0.093	67.95	1.915	35.59	1.004	51.61	1.456	97.73	2.756	32.26	0.908	45.59	1.286
SO ₄	2.46	0.051	10.5	0.218	5.5	0.115	2.14	0.045	3.96	0.083	3.04	0.063	0.72	0.015	1.38	0.029	2.18	0.045	2.28	0.048
NO ₃ -N	0	0	0.01	0.001	0	0	0	0	0.076	0.005	0.032	0.002	0.097	0.007	0.059	0.004	0.033	0.002	0.092	0.006
PO ₄ -P	0.265	0.026	0.258	0.025	0.082	0.008	0.013	0.001	0.21	0.020	0.24	0.023	0.78	0.075	0.26	0.025	0.67	0.065	0.31	0.030
Total Anions(A)		2.899		7.471		2.780		2.382		4.039		3.139		4.054		4.916		3.386		3.552
NH ₄ -N	0.95	0.069	0.84	0.060	0.45	0.032	0.31	0.022	2.13	0.152	1.65	0.118	1.09	0.078	2.22	0.159	0.93	0.067	1.87	0.133
Na	25.3	1.100	127	5.52	16.8	0.731	10.8	0.470	33.2	1.443	27.8	1.209	56.0	2.435	40.6	1.765	41.0	1.783	33.5	1.457
K	6.05	0.155	7.3	0.19	4.8	0.12	4.7	0.12	10.2	0.261	8.7	0.222	5.8	0.148	10.8	0.276	6.6	0.169	9.2	0.235
Ca	12.85	0.642	15.93	0.796	16.27	0.843	14.48	0.724	18.65	0.932	13.80	0.690	11.64	0.582	24.26	1.213	10.53	0.527	16.52	0.826
Mg	8.85	0.728	9.90	0.814	10.21	0.840	9.06	0.745	14.28	1.074	9.55	0.786	7.24	0.596	16.22	1.335	7.87	0.657	10.13	0.836
FeII	2.96	0.106	3.20	0.115	4.86	0.174	7.65	0.274	2.76	0.099	2.20	0.079	6.00	0.215	4.24	0.152	3.18	0.114	2.67	0.096
Mn	0.41	0.015	0.48	0.017	0.845	0.031	1.15	0.042	0.745	0.027	0.60	0.022	0.525	0.019	0.79	0.029	0.395	0.014	0.39	0.014
Total Cations(C)		2.814		7.512		2.771		2.397		3.988		3.126		4.073		4.929		3.321		3.597
SiO ₂	45.2		40		51.6		62.4		54		44.3		34.5		42.5		46.5		53	
(A)/(C)		1.03		0.995		1.00		0.99		1.01		1.00		0.995		1.00		1.02		0.99
Ca/Na		0.58		0.14		1.15		1.54		0.65		0.57		0.24		0.69		0.30		0.57
K/Na		0.14		0.034		0.17		0.26		0.18		0.18		0.06		0.16		0.095		0.16
Mg/Ca		1.13		1.02		0.995		1.03		1.15		1.14		1.02		1.10		1.23		1.01
Analyst	M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.	

Table 2-3. Chemical analyses of groundwaters from Eastern Osaka (1)

Group	B		B		B		B		B		B		B		B		B		B	
Well No.	N-5		N-6(1)		N-6(2)		N-7(1)		N-7(2)		N-7(3)		N-8(1)		N-8(2)		N-9		N-11	
Depth of Perforation(m)	37~62		45~65				25~49						25~49				21~86		33~67	
Total Depth(m)	78.8		75				60						60				91		91	
Date of Collection	Apr. 5, '67		Sep. 5, '67		Nov. 25, '68		Dec. 10, '65		Dec. 30, '67		Dec. 27, '69		Feb. 23, '66		Nov. 25, '68		Oct. 6, '67		Oct. 6, '67	
Temp. (°C)	16.6		17.1		17.1		16.6		16.7		17.5		16.6		17.2		16.8		16.9	
EC. (μ U.18°C)	963		600		464		598		445		485		540		479		710		1043	
pH	7.4		6.9		7.0		7.0		7.1		7.2		7.0		7.0		7.0		7.0	
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	ppm
HCO ₃	138.4	2.269	132.6	2.174	134.1	2.198	134.4	2.203	137.1	2.248	126.7	2.077	138.3	2.268	134.1	2.198	168.7	2.766	151.7	2.487
Cl	283.7	8.00	147.8	4.168	97.0	2.736	133.2	3.757	94.1	2.654	82.76	2.334	131.0	3.695	103	2.905	148.3	4.183	272.8	7.694
SO ₄	9.4	0.196	5.4	0.113	6.6	0.138	5.5	0.115	1.18	0.046	32.8	0.683	4.5	0.094	3	0.063	4	0.083	8.8	0.183
NO ₃ -N	0.38	0.027	0.24	0.017	0.077	0.005	0	0	0.24	0.017	0	0	0.09	0.006
PO ₄ -P	0.26	0.025	0.363	0.035	0.46	0.044	0.35	0.034	0.255	0.025	0.59	0.058	0.31	0.030	0.43	0.047	0.84	0.080	0.32	0.030
Total Anions(A)	10.49		6.490		5.143		6.126		4.978		5.152		6.104		5.213		7.112		10.40	
NH ₄ -N	1.20	0.086	0.90	0.064	1.01	0.072	0.85	0.061	0.95	0.068	0.955	0.068	0.86	0.061	0.72	0.051	1.26	0.090	1.25	0.089
Na	158	6.87	105	4.565	82.2	3.574	109	4.739	77.5	3.370	74.1	3.222	107	4.652	82.1	4.004	115	5.00	180	7.826
K	9.4	0.240	7.3	0.187	6.43	0.166	7.6	0.194	6.6	0.169	6.5	0.166	7.8	0.199	6.65	0.170	9.2	0.236	11.5	0.294
Ca	25.22	1.261	14.20	0.710	8.88	0.444	11.38	0.568	9.68	0.484	11.56	0.578	10.74	0.536	6.26	0.313	13.70	0.685	17.80	0.890
Mg	20.64	1.700	11.70	0.963	8.84	0.727	7.39	0.608	9.32	0.766	11.05	0.909	8.71	0.716	7.49	0.616	13.06	1.073	16.49	1.356
Fe II	3.48	0.127	2.97	0.106	4.56	0.163	2.40	0.087	2.9	0.106	5.74	0.205	2.50	0.091	1.76	0.063	2.35	0.084	1.36	0.019
Mn	0.745	0.027	0.725	0.026	0.72	0.026	0.55	0.020	1.225	0.045	0.50	0.018	0.59	0.021	0.655	0.023	0.73	0.026
Total Cations(C)	10.31		6.621		5.172		6.277		4.963		5.193		6.273		5.238		7.191		10.53	
SiO ₂	27.6		26.3		30.8		24.9		23		25.4		28.6		28.4		25.6		30.2	
(A)/(C)		1.02		0.98		0.99		0.975		1.00		0.99		0.97		0.99		0.99		0.99
Ca/Na		0.18		0.155		0.12		0.22		0.14		0.18		0.115		0.078		0.14		0.11
K/Na		0.035		0.041		0.046		0.041		0.050	25.4	0.051		0.043		0.042		0.047		0.038
Mg/Ca		1.35		1.36		1.64		1.07		1.58		1.57		1.33		1.97		1.57		1.52
Na/Cl		0.86		1.095		1.31		1.26		1.27		1.38		1.26		1.38		1.19		1.02
Mg/Cl		0.21		0.23		0.27		0.16		0.29		0.39		0.19		0.21		0.26		0.18
Analyst	M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.	

Table 2-4. Chemical analyses of groundwaters from Eastern Osaka (2)

Group	B		B		C		C		C		C		C		C		C		C	
Well No.	KD-15		KD-16		N-1		N-2		N-16		N-23		N-24		N-37		KD-5		KD-6	
Depth of Perforation(m)	37~90		31~99		42~85		75~124		42~57		50~71		45~120		35~120		80~150		81~150	
Total Depth(m)	96		106		100		130		60		76		135		121		180		180	
Date of Collection	Nov. 21, '67		Nov. 21, '67		Dec. 10, '65		Dec. 10, '22		Dec. 22, '65		Nov. 8, '66		Nov. 8, '66		Mar. 27, '67		Feb. 8, '66		Feb. 8, '66	
Temp. (°C)	16.5		16.5		16.9		17.8		16.5		16.5		16.5		16.3		19.3		19.7	
EC. (μ U. 18°C)	800		935		256		250		270		185		240		307		212		222	
pH	7.0		6.8		6.7		6.7		6.8		6.6		6.6		6.7		7.0		7.0	
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
HCO ₃	217.9	3.572	164.3	2.694	178.5	2.926	176.1	2.887	181.6	2.977	85.6	1.404	123.5	2.025	194.5	3.188	134.1	2.198	138.4	2.269
Cl	187.7	5.295	267.1	7.535	3.77	0.106	4.52	0.127	4.09	0.115	18.0	0.507	12.5	0.352	17.9	0.505	6.09	0.172	7.61	0.215
SO ₄	6.9	0.144	0.94	0.020	1.9	0.040	0.7	0.014	3.0	0.062	3.0	0.062	0	0	2.4	0.050	0.9	0.019	1.7	0.035
NO ₃ -N	0.045	0.003	0.054	0.004	0.31	0.022	0.39	0.028	0.16	0.011	0.01	0.001	0.01	0.001	0.05	0.004	0.35	0.025	0.50	0.036
PO ₄ -P	1.00	0.097	0.78	0.075	0.97	0.095	0.41	0.040	1.24	0.120	0.173	0.017	0.52	0.050	0.405	0.040	0.36	0.035	0.38	0.037
Total Anions(A)		9.111		10.23		3.189		3.096		3.285		1.991		2.428		3.787		2.449		2.592
NH ₄ -N	3.48	0.249	3.18	0.227	2.88	0.206	2.30	0.164	3.27	0.234	0.47	0.033	2.14	0.153	4.09	0.292	3.20	0.229	3.12	0.223
Na	160	6.957	176	7.652	15.0	0.652	16.0	0.695	17.0	0.739	14.5	0.630	14.2	0.618	19.0	0.826	21.0	0.913	23.0	1.000
K	9.8	0.251	9.7	0.248	6.4	0.164	7.1	0.182	5.9	0.151	1.8	0.046	4.5	0.115	10.0	0.256	10.5	0.269	10.8	0.276
Ca	15.50	0.775	20.64	1.032	22.68	1.134	20.36	1.018	22.24	1.112	14.81	0.740	14.30	0.715	25.39	1.269	8.78	0.438	9.72	0.485
Mg	10.41	0.857	14.26	1.174	8.97	0.738	8.61	0.708	8.99	0.740	3.92	0.322	6.33	0.522	9.73	0.802	6.02	0.495	6.97	0.573
Fe II	2.12	0.076	4.54	0.163	6.55	0.234	7.00	0.251	7.10	0.254	5.90	0.211	8.35	0.298	9.05	0.324	1.00	0.036	1.38	0.050
Mn	0.355	0.013	1.22	0.044	0.48	0.017	0.37	0.013	0.57	0.021	0.235	0.009	0.46	0.017	0.625	0.023	0.44	0.016	0.55	0.023
Total Cations(C)		9.178		10.54		3.145		3.031		3.251		1.991		2.438		3.792		2.396		2.627
SiO ₂	43.8		39.2		43.4		53.0		48.6		51.8		52.4		57.5		54.0		58.6	
(A)/(C)		0.99		0.98		1.01		1.02		1.01		1.00		1.00		1.00		1.02		0.99
Ca/Na		1.11		1.35		1.74		1.47		1.51		1.17		1.15		1.54		0.48		0.485
K/Na		0.036		0.032		0.25		0.26		0.20		0.073		0.19		0.31		0.29		0.28
Mg/Ca		1.11		1.14		0.65		0.69		0.67		0.435		0.73		0.63		1.13		1.18
Na/Cl		1.31		1.02																
Mg/Cl		0.16		0.16																
Analyst	M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.	

Table 2—5. Chemical analyses of groundwaters from Eastern Osaka

Group	C		C		C		C		C		C				D		D		D	
Well No.	KD—10		KD—11		N—39 (1)		N—39 (2)		N—39 (3)		N—42		M—22		KD—13		KD—18		KD—19	
Depth of Perforation(m)	230~286		90~159		182~215						181~235.5		67~133		62~166		?		?	
Total Depth(m)	300		180		223						250		137		180		200		180	
Date of Collection	Feb. 8, '66		Dec. 13, '65		Nov. 30, '67		Nov. 25, '68		Nov. 27, '69		Nov. 27, '69		Sep. 28, '66		Jan. 22, '68		Nov. 8, '66		Dec. 5, '67	
Temp. (°C)	21.7		18.5		20.5		20.1		20.2		19.8		17.5		19.1		20.0		19.5	
EC. (μ U. 18°C)	192		238		218		262		304		330		870		1020		1450		1164	
pH	7.4		7.2		7.2		7.1		7.2		7.0		7.2		7.1		7.0		6.9	
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
HCO ₃	122.2	2.003	152.9	2.506	131.9	2.163	132.2	2.173	132.1	2.166	134.3	2.202	261.4	4.285	247.2	4.053	237.7	3.896	245.7	4.028
Cl	3.59	0.101	4.42	0.125	8.34	0.234	28.3	0.798	38.48	1.085	49.38	0.393	167.6	4.727	328	9.255	483	13.625	386	10.89
SO ₄	0.7	0.015	3.1	0.065	0	0	0.4	0.008	2.0	0.042	1.6	0.033	3.8	0.079	2.3	0.049	3.4	0.071	6.4	0.134
NO ₃ —N	0.36	0.026	0.57	0.041	0.05	0.004	---	---	0.025	0.002	0.02	0.001	---	---	0.07	0.005	0.028	0.002	0.063	0.005
PO ₄ —P	0.44	0.043	0.51	0.049	0.20	0.020	0.21	0.020	0.23	0.022	0.23	0.022	0.58	0.057	0.85	0.082	1.56	0.150	1.50	0.145
Total Anions(A)		2.188		2.786		2.421		2.999		3.315		3.651		9.148		13.44		17.74		15.20
NH ₄ —N	2.67	0.191	4.37	0.312	2.14	0.156	2.72	0.194	2.63	0.188	3.375	2.241	7.83	0.559	13.1	0.936	16.7	1.195	18.7	1.336
Na	18.0	0.783	24.8	1.078	22.0	0.956	28.3	1.231	33.4	1.454	31.1	1.354	80.0	3.478	125	5.435	206	8.96	153	6.652
K	8.8	0.225	13.6	0.348	9.9	0.253	10.6	0.271	10.5	0.269	11.65	0.298	18.5	0.473	28.8	0.738	30.0	0.767	26.2	0.670
Ca	9.46	0.472	8.78	0.438	8.57	0.428	9.70	0.485	10.50	0.525	12.12	0.606	46.69	2.335	57.31	2.865	63.23	3.162	60.82	3.041
Mg	4.91	0.404	6.02	0.495	7.29	0.600	9.42	0.775	9.44	0.776	12.09	0.994	24.72	2.033	38.44	3.161	44.05	3.625	40.21	3.307
Fe II	0.79	0.029	0.83	0.030	1.78	0.064	2.10	0.075	2.17	0.078	2.00	0.072	1.12	0.041	2.00	0.072	2.40	0.086	2.52	0.090
Mn	0.45	2.016	0.32	0.011	---	---	0.56	0.020	0.50	0.018	0.58	0.021	0.41	0.015	0.845	0.031	0.93	0.033	0.82	0.030
Total Cations(C)		2.120		2.712		2.457		3.051		3.308		3.586		8.934		13.24		17.883		15.13
SiO ₂	58.4		52.2		55		54		55		55		49.2		57.8		65.3		54	
(A)/(C)		1.03		1.03		0.99		0.99		1.00		1.02		1.02		1.015		0.995		1.005
Ca/Na		0.60		0.41		0.45		0.39		0.36		0.45		0.67		0.53		0.35		0.46
K/Na		0.29		0.32		0.26		0.22		0.19		0.22		0.14		0.14		0.086		0.11
Mg/Ca		0.86		1.13		1.40		1.60		1.48		1.64		0.87		1.10		1.15		1.09
Na/Cl														0.74		0.59		0.66		0.61
Mg/Cl														0.43		0.34		0.27		0.30
Analyst	M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.	

Table 2-6. Chemical analyses of groundwaters from Eastern Osaka (4)

Group	D		D		D		D		D		D		D		D		D		D	
Well No.	KD-22		D-15		D-16(1)		D-16(2)		D-16(3)		D-17(1)		D-17(2)		D-17(3)		D-18(1)		D-18(2)	
Depth of Perforation(m)	82~195		86~169		60~162						64~187						81~230			
Total Depth(m)	200		182		182						200						240			
Date of Collection	Dec. 5, '67		Nov. 29, '66		Nov. 29, '66		Feb. 7, '69		Dec. 8, '69		Nov. 29, '66		Feb. 7, '69		Dec. 8, '69		Nov. 29, '66		Feb. 23, '68	
Temp. (°C)	19.5		20.2		19.3		20.2		20.0		19.4		20.0		19.6		21.1		21.5	
EC. (μ U. 18°C)	1960		314		319				878		315				618		333		606	
pH	7.0		7.2		7.0		6.9		6.8		7.0		7.0		6.9		7.0		6.9	
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
HCO ₃	244.1	4.002	271.5	4.450	312.8	5.128	314.5	5.155	305.7	5.011	286.0	4.688	288.5	4.73	283.5	4.647	281.1	4.608	247.1	4.050
Cl	814	22.96	6.3	0.177	5.12	0.144	62.4	1.76	139.8	3.944	6.58	0.186	158.6	4.473	69.5	1.961	8.13	0.230	419	11.82
SO ₄	30.1	0.627	0.7	0.015	1.56	0.032	4.08	0.085	1.1	0.023	1.43	0.030	4.34	0.090	1.2	0.025	0.88	0.019	3.04	0.063
NO ₃ -N	0.034	0.002	0		0		0.024	0.002	0		0		0.070	0.005	0		0		0.09	0.006
PO ₄ -P	1.60	0.155	0.64	0.062	0.81	0.079	1.08	0.105	1.82	0.176	1.20	0.116	1.24	0.120	1.355	0.131	0.53	0.051	1.27	0.123
Total Anions(A)		27.75		4.704		5.383		7.107		9.154		5.020		9.418		6.739		4.908		16.06
NH ₄ -N	35.7	2.550	6.35	0.453	8.30	0.592	10.28	0.734	14.74	1.053	6.40	0.457	9.00	0.642	10.34	0.739	6.40	0.457	19.4	1.384
Na	298	12.96	22.4	0.974	26.3	1.144	37.0	1.610	54.5	2.372	25.0	1.087	60.0	2.610	39.1	1.700	25.3	1.198	108	4.696
K	30.8	0.787	18.9	0.483	18.9	0.483	16.0	0.409	19.9	0.509	14.2	0.363	17.0	0.435	15.8	0.404	15.9	0.407	22.3	0.571
Ca	104.50	5.225	28.74	1.434	32.44	1.618	43.14	2.154	59.56	2.878	33.10	1.651	57.55	2.870	40.88	2.044	31.57	1.575	90.60	4.520
Mg	77.60	6.382	15.07	1.239	17.40	1.431	20.06	1.652	26.75	2.200	16.30	1.340	27.62	2.270	20.84	1.714	15.86	1.304	56.60	4.655
Fe II	4.74	0.169	0.60	0.021	2.52	0.090	3.46	0.124	3.64	0.130	1.70	0.061	3.46	0.124	2.30	0.073	1.82	0.065	5.36	0.192
Mn	1.42	0.051	---	---	---	---	0.72	0.026	0.97	0.035	---	---	0.81	0.029	0.56	0.020	---	---	1.22	0.044
Total Cations(C)		28.12		4.604		5.358		6.709		9.177		4.959		8.980		6.694		5.006		16.07
SiO ₂	49.5		54.5		61.0		69		65		58.3		62.8		55.5		41.4		68.3	
(A)/(C)		0.99		1.02		1.00		1.06		1.00		1.01		1.05		1.01		0.97		1.00
Ca/Na		0.40		1.47		1.41		1.34		1.21		1.52		1.10		1.20		11.31		0.96
K/Na		0.061		0.50		0.42		0.25		0.22		0.33		0.17		0.24		0.34		0.12
Mg/Ca		1.22		0.86		0.88		0.77		0.76		0.81		0.79		0.84		0.83		1.03
Na/Cl		0.56								0.60				0.58						0.40
Mg/Cl		0.28								0.56				0.51						0.39
Analyst	M. T.		M. Nagasawa		M. N.		M. N.		M. T.		M. N.		M. N.		M. T.		M. N.		M. N.	

Table 2-7. Chemical analyses of groundwaters from Eastern Osaka

Group	D		D		D		D		D		D		E		E		E		E	
Well No.	D-18(3)		D-18(4)		D-19(1)		D-19(2)		D-19(3)		D-20(1)		D-20(2)		D-20(3)		D-20(4)		D-22	
Depth of Perforation(m)					74~178						178~290								167~296	
Total Depth(m)					185						300								306	
Date of Collection	Feb. 7, '69		Dec. 8, '69		Nov. 29, '66		Nov. 7, '67		Feb. 7, '69		Nov. 29, '66		Nov. 7, '67		Feb. 7, '69		Dec. 8, '69		Dec. 8, '69	
Temp. (°C)	20.5		20.2		20.2		20.5		20.8		23.2		24.7		24.2		24.2		24.0	
EC. (μS , 18°C)			830								590		470				442		520	
pH	6.9		7.0								7.1		7.3		7.2		7.2		7.1	
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
HCO ₃	267.2	4.38	276.6	4.534	282.8	4.636	289.8	4.750	281.5	4.615	280.1	4.592	302.9	4.965	304.4	4.99	290.9	4.769	325.9	5.343
Cl	236.0	6.656	142.2	4.011	12.83	0.362	177.8	5.016	287.0	8.094	147.0	4.146	37.6	1.061	30.8	0.860	10.59	0.299	30.68	0.865
SO ₄	5.53	0.115	1.3	0.027	1.15	0.024	0.10	0.002	6.26	0.131	1.42	0.029	0.08	0.0	0	0	0.9	0.019	0.7	0.015
NO ₃ -N	0.037	0.003	0	0	0	0	0	0	0.019	0.001	0		0		0.03	0.002	0		0	
PO ₄ -P	0.72	0.070	1.10	0.107	0.73	0.071	0.97	0.094	0.81	0.079	0.53	0.051	0.74	0.072	0.70	0.068	0.93	0.090	0.88	0.085
Total Anions(A)		11.22		8.679		5.093		9.862		12.92		8.818		6.098		5.920		5.177		6.308
NH ₄ -N	8.64	0.616	10.32	0.737	6.90	0.493	13.80	0.985	13.76	0.982	10.70	0.764	12.10	0.864	6.24	0.446	9.70	0.693	10.19	0.727
Na	78.8	3.428	58.1	2.526	22.6	0.983	57.8	2.513	90.0	3.915	73.2	3.084	57.7	2.505	69.0	3.002	58.8	2.557	65.95	2.867
K	16.6	0.424	16.1	0.412	17.0	0.435	22.2	0.581	20.8	0.532	28.0	0.716	22.2	0.667	16.6	0.424	16.0	0.409	18.1	0.463
Ca	65.80	3.284	50.36	2.518	31.76	1.584	58.25	2.907	77.10	3.850	41.86	2.086	17.86	0.891	19.17	0.957	14.38	0.719	22.00	1.100
Mg	37.00	3.042	30.00	2.467	17.50	1.439	32.28	2.655	39.21	3.224	24.86	2.044	13.52	1.112	12.14	0.998	9.61	0.790	13.39	1.101
FeII	3.64	0.130	2.62	0.094	1.10	0.039	3.58	0.128	4.88	0.175	2.06	0.074	0.68	0.024	0.54	0.019	0.51	0.018	0.93	0.033
Mn	0.76	0.028	0.68	0.025	0.83	0.030	0.90	0.033	0.58	0.021	0.23	0.008	0.16	0.006	0.23	0.008
Total Cations(C)		10.95		8.779		4.973		9.799		12.71		8.768		6.084		5.854		5.192		6.299
SiO ₂	59.7		55.5		55.0		60.7		64.2		69.1		67.5		67.2		62		66	
(A)/(C)		1.025		0.99		1.02		1.005		1.02		1.005		1.00		1.01		1.00		1.00
Ca/Na		0.96		0.99		1.61		1.16		0.985		0.675		0.355		0.32		0.28		0.38
K/Na		0.12		0.16		0.44		0.23		0.14		0.23		0.27		0.14		0.16		0.16
Mg/Ca		0.93		0.98		0.91		0.91		0.84		0.98		1.25		1.04		1.10		1.00
Na/Cl		0.52		0.63				0.50		0.48		0.74								
Mg/Cl		0.46		0.62				0.53		0.40		0.49								
Analyst	M.N.		M.T.		M.N.		M.N.		M.N.		M.N.		M.N.		M.N.		M.T.		M.T.	

Table 2—8. Chemical analyses of groundwaters from Eastern Osaka

Group	E		E		E		E		E		E							
Well No.	D—2		D—3		D—4		D—5		D—7		D—21		Hi—30	Hi—32	Hi—33	Hi—34		
Depth of Perforation(m)	57~102		23~122		21~118		92~147		52~116		69.5~157		83.5~179.5	66~121	60~193	56~205		
Total Depth(m)	125		127		125		147		120		175		183	150	200	210		
Date of Collection	Jan. 28, '66		Jan. 28, '66		Nov. 11, '66		Jan. 28, '66		Nov. 11, '66		Apr. 9, '69		Apr. 11, '69	Nov. 20, '69	Nov. 20, '69	Nov. 20, '69		
Temp. (°C)	18.2		18.0		17.5		19.5		18.9		19.2		17.7	18.8	20.0	19.1		
EC. (μ U. 18°C)	408		450		320		345		385		404		490	306	262	324		
pH	6.9		6.8		7.4		7.0		6.9		7.0		7.1	7.2	7.4	7.1		
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
HCO ₃	322.1	5.280	370.9	6.080	332.1	5.444	263.2	4.315	377.9	6.195	276.7	4.536	368.0	6.032	229.2	3.757	190.7	3.126
Cl	5.99	0.169	6.63	0.187	7.60	0.214	4.49	0.127	7.02	0.198	6.16	0.174	6.32	0.178	4.74	0.134	5.18	0.146
SO ₄	3.58	0.075	7.42	0.155	1.88	0.039	0.56	0.012	1.81	0.038	1.44	0.030	2.34	0.049	0.8	0.017	0.8	0.017
NO ₃ —N	0.20	0.014	0.46	0.032	0		0.14	0.010	0		0.03	0.002	0.06	0.004	0.025	0.002	0.024	0.002
PO ₄ —P	0.62	0.060	0.58	0.056	0.27	0.026	0.895	0.087	1.34	0.130	1.76	0.170	2.31	0.224	0.49	0.047	0.275	0.027
Total Anions(A)		5.598		6.510		5.723		4.551		6.561		4.912		6.487		3.957		3.318
NH ₄ —N	6.86	0.489	12.28	0.877	13.30	0.949	3.85	0.274	15.10	1.078	6.85	0.489	9.8	0.700	1.53	0.109	1.24	0.089
Na	28.4	1.234	31.3	1.361	18.4	0.800	23.3	1.013	26.8	1.165	25.3	1.100	27.5	0.196	19.1	0.829	23.9	1.038
K	16.0	0.409	9.6	0.246	16.1	0.412	10.5	0.269	18.9	0.483	10.2	0.261	4.7	0.120	3.36	0.086	4.02	0.103
Ca	36.78	1.835	38.32	1.912	35.46	1.769	32.76	1.635	38.15	1.903	32.96	1.648	56.40	2.820	42.72	2.136	28.08	1.404
Mg	16.78	1.380	18.22	1.498	18.13	1.491	14.72	1.211	19.92	1.638	15.21	1.252	16.05	1.320	8.35	0.687	7.31	0.601
FeII	5.32	0.191	9.64	0.345	8.12	0.291	0.76	0.027	6.74	0.241	0.90	0.033	4.28	0.153	0.35	0.012	0.30	0.011
Mn	0.48	0.018	0.95	0.035	0.24	0.009	0.16	0.006
Total Cations(C)		5.538		6.239		5.712		4.429		6.508		4.801		6.344		3.868		3.252
SiO ₂	67.6		66.2		41.3		58.8		66.1		53.0		52.5		40		39	
(A)/(C)		1.01		1.04		1.00		1.03		1.01		1.02		1.02		1.02		1.02
Ca/Na		1.49		1.41		2.21		1.61		1.63		1.50		2.36		2.58		1.35
K/Na		0.33		0.18		0.51		0.27		0.41		0.24		0.10		0.10		0.10
Mg/Ca		0.75		0.78		0.84		0.74		0.86		0.76		0.47		0.32		0.43
Analyst	M.N.		M.N.		M.N.		M.N.		M.N.		M.T.		M.T.		M.T.		M.T.	

Table 2—9. Chemical analyses of groundwaters from Eastern Osaka

Group																				
Well No.	Ka-29		Ka-30		F-36		F-37		F-38		F-39		Y-14		Y-16		Y-44		Y-48	
Depth of Perforation (m)	47~198		49~185		106~162		79~165		40~115		161~290		23~58		50~117		52~142		53~186	
Total Depth(m)	200		200		167		205		130		290		60		118		150		200	
Date of Collection	Nov. 20, '69		Nov. 20, '69		Nov. 13, '69		Nov. 13, '69		Nov. 13, '69		Nov. 13, '69		July. 21, '66		July. 21, '66		Aug. 17, '66		Aug. 17, '66	
Temp. (°C)	18.3		18.4		20.5		19.6		19.0		24.2		20.6		18.0		18.1		17.5	
EC. (μ U. 18°C)	433		450		288		302		365		289		440		410		250		253	
pH	6.8		7.1		6.9		6.9		6.9		7.1		6.6		6.8		6.8		6.8	
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
HCO ₃	289.6	4.748	292.1	4.789	196.3	3.218	213.6	3.502	252.6	4.141	201.1	3.296	139.6	2.289	142.7	2.340	145.4	2.384	147.0	2.410
Cl	20.17	0.569	25.40	0.716	9.54	0.269	6.00	0.169	11.18	0.315	4.36	0.123	48.56	1.370	60.01	1.692	14.78	0.417	15.67	0.442
SO ₄	0.8	0.017	0.8	0.017	<1		<1	.0	<1		<1		55.0	1.146	43.5	0.906	8.1	0.169	7.8	0.163
NO ₃ -N	0.041	0.003	0.043	0.003	0.032	0.002	0.04	0.003	0.043	0.003	0.04	0.003	1.50	0.107	0		1.5	0.108	1.3	0.093
PO ₄ -P	2.54	0.247	2.67	0.259	0.895	0.087	1.415	0.137	1.69	0.164	1.81	0.176	0.065	0.006	0.13	0.013	0.855	0.083	0.9	0.087
Total Anions(A)	5.584		5.784		3.576		3.811		4.623		3.598		4.918		4.951		3.161		3.195	
NH ₄ -N	7.02	0.501	7.43	0.530	2.46	0.247	4.17	0.297	4.75	0.339	5.25	0.375	0.054	0.004	0.115	0.008	1.69	0.121	2.04	0.146
Na	44.2	1.922	45.8	1.991	21.32	0.927	21.46	0.933	26.84	1.167	25.34	1.102	27.5	1.196	29.0	1.261	19.0	0.827	19.3	0.839
K	13.6	0.349	13.5	0.345	9.65	0.247	11.38	0.291	13.44	0.344	11.64	0.298	2.9	0.074	2.0	0.051	3.3	0.084	3.4	0.087
Ca	26.18	1.309	27.82	1.391	17.76	0.888	20.06	1.003	21.28	1.064	17.18	0.859	46.90	2.340	43.92	2.192	28.11	1.406	25.89	1.292
Mg	16.33	1.343	14.93	1.228	13.93	1.146	13.78	1.113	17.81	1.465	10.20	0.839	15.84	1.303	14.11	1.160	5.18	0.426	5.89	0.484
Fe II	4.46	0.160	4.40	0.157	2.26	0.081	2.02	0.072	4.08	0.146	1.02	0.036	0.92	0.033	3.05	0.109	5.06	0.181	6.18	0.222
Mn	0.69	0.025	0.695	0.025	0.60	0.022	0.51	0.018	0.675	0.025	0.395	0.014	4.65	0.169	2.70	0.098	0.86	0.031	0.97	0.035
Total Cations(C)	5.609		5.667		3.558		3.747		4.550		3.518		5.119		4.879		3.076		3.105	
SiO ₂	62		62		55		53.4		54.6		67.4		27.2		35.8		50.8		49.2	
(A)/(C)		0.995		1.02		1.005		1.02		1.02		1.02		0.96		1.015		1.03		1.03
Ca/Na		0.68		0.70		0.96		1.075		0.91		0.78		1.96		1.74		1.70		1.54
K/Na		0.18		0.17		0.27		0.31		0.295		0.27		0.062		0.040		0.10		0.105
Mg/Ca		1.03		0.88		1.29		1.11		1.38		0.98		0.56		0.53		0.30		0.375
Ma/Cl																				
Mg/Cl																				
Analyst	M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.		M. T.	

Table 2-10. Chemical analyses of groundwaters from test wells

Group	Test Well, Niwakubo						Test Well, Konoike					
Well No.	N-50m		N-100m		N-250m		K-50m		K-100m		K-200m	
Depth of Perforation (m)	31.5~45		59~82		208~238.5		45~50		92~97		175~193	
Total Depth (m)	50		100		250		51		100		200	
Date of Collection	Mar. 11, '68		Mar. 15, '68		Mar. 20, '68		Dec. 22, '67		Jan. 24, '68		Dec. 7, '67	
Temp. (°C)	16.5		17.4		21.0		18.3		19.9		23.3	
EC. (μ S · 18°C)	700		810		250		575		440		909	
pH	7.1		7.4		7.4		6.8		7.4		6.8	
	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm	ppm	epm
HCO ₃	163.7	2.684	153.8	2.521	151.3	2.480	380.0	6.229	293.6	4.813	251.5	4.123
Cl	160.0	4.513	184.6	5.206	3.30	0.093	40.86	1.152	5.73	0.162	212	5.979
SO ₄	2.94	0.051	2.34	0.049	1.17	0.024	1.35	0.028	3.64	0.076	1.19	0.025
NO ₃ -N	0		0.04	0.003	0.04	0.003	0.057	0.004	0.066	0.005	0.025	0.002
PO ₄ -P	0.01	0.001	0.046	0.045	0.30	0.029	1.95	0.188	2.38	0.231	1.60	0.155
Total Anions (A)		7.249		7.824		2.629		7.601		5.287		10.28
NH ₄ -N	2.60	0.186	5.18	0.370	2.60	0.186	9.95	0.711	10.9	0.779	12.4	0.885
Na	94	4.087	92	4.000	17.2	0.748	46.0	2.000	28.0	1.217	57.0	2.478
K	10.4	0.266	19.7	0.504	9.0	0.230	19.9	0.509	21.1	0.537	20.0	0.512
Ca	24.44	1.222	24.86	1.243	14.76	0.738	41.28	2.064	24.75	1.237	67.26	3.363
Mg	17.90	1.472	19.06	1.567	7.30	0.600	26.06	2.142	17.24	1.420	33.80	2.775
Fe II	5.90	0.211	4.12	0.148	0.97	0.035	5.82	0.208	1.19	0.043	3.10	0.111
Mn	0.77	0.028	0.75	0.027	0.95	0.035	0.97	0.035	0.99	0.036
Total Cations (C)		7.472		7.849		2.572		7.669		5.233		19.16
SiO ₂	35.5		56.7		55		49.8		45		62.5	
Residue	470		533		173		407		247		655	
KMnO ₄ cons.	5.60		4.12		3.35		13.57		8.50		11.37	
CH ₄ ml/l	4.2		9.7		0.69		41.0		21.2		24.6	
Tritium T.U.	7.3		2.7		1.2		12.7		3.9		1.9	
Analyst	M. T.		M. T.		M. T.		M. T.		M. T.		M. T.	

Table 2-11. Chemical analyses of surface waters

Group	Surface Water				
	Yodo R.		Yamato R.		Ishi R.
Local Description	Hira-kata	Kema	Kashi-hara	Fujii-dera	Fujii-dera
Date of Collection	Feb. 1956	Nov. 1956	Feb. 14, 1968		
Temp. (°C)	5.0	8.3	5.1	4.6	4.0
pH	7.1	7.0	7.2	7.2	7.1
	ppm	ppm	ppm	ppm	ppm
HCO ₃			141.3	131.9	82.2
Cl	7.8	37.8	37.55	47.35	77.31
SO ₄	5	8.6	28.3	43.0	19.9
NO ₃ -N			0.09	0.06	1.02
PO ₄ -P	0.10	0.01	0.71	0.55	0.10
NH ₄ -N	0	0.39	6.6	3.12	0.22
Na			32.0	44.0	45.6
K			9.4	7.0	2.65
Ca	8.0	10.2	29.12	28.61	28.95
Mg	1.6	4.1	5.88	6.15	6.45
Fe (Total)	0.03	2.55	1.66	1.14	0.98
Mn			0.45	0.34	0.06
SiO ₂	16.4	16.4	26.8	24.8	17.4
Analyst	Geol. Surv. Japan		M. Tsurumaki		