# Fisheries and Marine Service Manuscript Report 1540

December 1979

THE OFFSHORE SEDIMENTS OF LAKE WINNIPEG

bу

G. J. Brunskill and B. W. Graham\*

Western Region

Fisheries and Marine Service

Department of Fisheries and the Environment

Winnipeg, Manitoba R3T 2N6

This is the 18th Manuscript Report from the Western Region, Winnipeg

<sup>\*</sup>Present Address: Inland Waters Directorate, Hydrology Research Division, 562 Booth Street, Ottawa, Ontario

© Minister of Supply and Services Canada 1979
Cat. no. Fs 97-4/1540 ISSN 0701-7618

BRUNSKILL, G. J., and B. W. GRAHAM. 1979. The offshore sediments of Lake Winnipeg. Can. Fish. Mar. Serv. MS Rep. 1540.

# **ERRATA**

p. 8, column 1, The reference should be to Brunskill and Schindler et al. 1979, Figs. 4-9.

p. 9, column 1, ...kurtosis values of 0-14 (graphic kurtosis range = 0.6 to 2.4).

p. 12, column 1, Should be: Brunskill, G. J., D. W. Schindler, S. E. M. reference 9 Elliott, and P. Campbell. 1979. The attenuation of light in Lake Winnipeg. Can. Fish. Mar. Serv. MS Rep. 1522: v + 79 p.

	TABLE OF CONTENTS		<u>Table</u>		Page
		<u>Page</u>	10	Water content, % loss on ignition, total carbon, carbonate carbon,	
	T/RESUME	v 1		organic carbon, total nitrogen and total phosphorus for sedi- ment samples from Lake Winnipeg,	
Sedim	ent sampling	1 2	11	3-12 October 1969	21
RESULTS		2		the labile fraction of Lake Winnipeg sedimentary phosphorus	22
Miner	alogy	3	12	Calculated values of organic carbon, carbonate carbon and total carbon	
	mistry of surface sediments	3 4		from ashing curves of Lake Winnipeg sediment samples collected	d
Rates	e elements in surface sediments . of supply of major and trace	5		3-12 October 1969, and their com- parison to wet chemical carbon	22
Lak	ments to the South Basin of the Winnipeg	5	13	analysis	2.2
Inter	of Lake Winnipeg sediments stitial water of surface	6 6		surface sediment samples from Lake Winnipeg collected 3-12 October 1969	23
DISCUSS	diments	0	14	Concentrations of major elements in Lake Winnipeg surface sediment	
SOL	rces	7	15	samples	24 s
ana	ilyses	8 9		in Lake Winnipeg surface sediment samples	25
Sedin	mentation rates and elemental	10	16	Relative atomic variation of trace elements in proportion to selected	
ACKNOWL REFEREN	EDGMENTS	11 12	• 7	conservative elements in Lake Winnipeg sediments	26
			17	Element concentration in suspended sediments from Red River drainage into the South Basin of Lake	
	LIST OF TABLES		18	Winnipeg, 2-9 May 1975 Suspended sediment chemistry for	26
<u>Table</u>		Page	10	Lake Winnipeg and Red River, by sodium metaborate fusion and	
1	Longitude, latitude and maximum depth of water at Lake Winnipeg			atomic absorption spectrophoto- metric determination	27
	sediment stations, 3-12 October 1969	15	. 19	Annual rates of supply of major and trace elements to the South Basin	
2	Particle size fractionation of Lake Winnipeg surface sediment		•	of Lake Winnipeg, and sedimenta- tion rates, derived from Red River suspended sediment chemistry, and	•
2	samples collected 3-12 October	15	20	discharge data over 1969-1973 .  Description of Lake Winnipeg sedi-	27
3	Quantitative X-ray diffraction carbonate mineralogy of Lake Winnipeg sediments collected		20	ment cores taken with a weighted benthos corer on 3-12 October	
4	3-12 October 1969	16	21	1969	28
7	fraction of Lake Winnipeg surface sediments collected			total carbon, carbonate carbon, organic carbon, total nitrogen,	
5	3-12 October 1969 Mineralogy of the fine silt	16		total phosphorus and molal ratios for selected Lake Winnipeg sedi-	
	fraction of Lake Winnipeg surface sediments collected	17		ment cores taken 3-12 October	31
6	3-12 October 1969 Mineralogy of the total clay	17	22	Major element chemistry of Lake Winnipeg sediment cores, taken 3-12 October 1969	34
	fraction of Lake Winnipeg surface sediments collected 3-12 October 1969	18	23	Trace element chemistry of Lake Winnipeg sediment cores, taken	
7	Mineralogy of coarse clay fraction of Lake Winnipeg surface sedi-		24	3-12 October 1969	36
	ments collected 3-12 October	19		and molybdenum in South Basin Lake Winnipeg sediment cores	20
8	Mineralogy of medium clay fraction of Lake Winnipeg surface sedi-		25	taken in July 1975	
_	ments collected 3-12 October 1969	19		water chemistry of sediment sample from Lake Winnipeg collected 3-12	<b>e</b> S
9	Mineralogy of suspended sediments from rivers draining into Lake		26	October 1969, squeezed February 1970	40 res
	Winnipeg, and a station from the South and North Basin of lake Winnipeg	20	בָט	applied to sediment particle size distributions	Α 1

<u>Table</u>		<u>Page</u>	<u>Figure</u>	<u> </u>	Page
27	Graphic parameters of the distri- bution of particle sizes of	42	19	The relationship between cobalt and iron for surface sediment	63
28	Lake Winnipeg sediment samples . Clay mineralogy of the total clay fraction of selected South Basin	43	20	samples from Lake Winnipegq The relationship between chromium and iron for surface sediment	64
29	Lake Winnipeg sediment samples . Comparison of present-day sedimen- tation rate data for Lake	43	21	samples from Lake Winnipeg The relationship between nickel and iron for surface sediment	65
30	Winnipeg and the St. Lawrence Great Lakes	43	22	samples from Lake Winnipeg The relationship between beryllium and iron for surface sediment	66
31	Supply rates Comparison of 1926 and 1969 sediment particle size data for Red River	44	23	samples from Lake Winnipeg The relationship between phosphorus and iron for surface sediment	00
	and Lake Winnipeg	44	24	samples from Lake Winnipeg Location of Lake Winnipeg sediment cores taken by Dr. R. J. Allan	67
	LIST OF FIGURES		25	in 1975	68
Figure		Page		lation in the North Basin of Lake Winnipeg	69
1 Igui c			26	Major regions of sediment accumu-	
1	Map of the watershed of Lake Winnipeg	],		lation in the Narrows and South Basin of Lake Winnipeg	7Ó
	its major tributaries, and the maximum extent of glacial Lake		27	Basin of Lake Winnipeg	
	Agassiz sediments	45		size and graphic standard deviation	
2	Locations of 1969 sampling stations in Lake Winnipeg	46		for Lake Winnipeg off-shore sedi- ment samples	71
3	Bathymetry of the South Basin and		28	The relationship between X and graphic	
_	Narrows of Lake Winnipeg	47		skewness of the particle"size dis-	
4	Bathymetry of the North Basin of Lake Winnipeg	48		tributions of Lake Winnipeg off- shore sediments	72
5	The distribution of particle sizes		29	The relationship between $\bar{X}_{a}$ and graphic	
	of Lake Winnipeg sediments	49		The relationship between $\bar{X}_{\varphi}$ and graphic kurtosis of the particle size dis-	
6	Isopleths of % silt in Lake Winnipeg sediments	50		tributions of Lake Winnipeg off- shore sediments	73
7	Isopleths of % clay in Lake Winnipeg	F1	30	The relationship of graphic skewness	
	sediments	51		to kurtosis for the particle size	
8	sediments	52		distributions of Lake Winnipeg off- shore sediments	74
9	The relationship between the sum of		31	The relationship of graphic_kurtosis	
	Ca and Mg and inorganic carbon for sediment samples from Lake			to standard deviation of X <sub>0</sub> for the particle size distributions of Lake	
	Winnipeg	53		Winnipeg off-shore sediments	75
10	The relationship between vanadium			, ,	
	and titanium for surface sedi- ment samples from Lake Winnipeg .	54			
11	The relationship between copper				
	and titanium for surface sedi-	55			
12	ment samples from Lake Winnipeg . The relationship between cobalt				
15	and titanium for surface sedi-	F.C			
3.7	ment samples from Lake Winnipeg . The relationship between chromium	56			
13	and titanium for surface sedi-				
	ment samples from Lake Winnipeg .	57			
14	The relationship between nickel and titanium for surface sedi-				
	ment samples from Lake Winnipeg .	58			
15	The relationship between beryllium				
	and titanium for surface sedi- ment samples from Lake Winnipeg .	59			
16	The relationship between phosphorus				
	and titanium for surface sedi-	60			
17	ment samples from Lake Winnipeg . The relationship between vanadium	ou			
11	and iron for surface sediment				
	samples from Lake Winnipeg	61	•		
18	The relationship between copper and iron for surface sediment				
	samples from Lake Winnipeg	62			

#### ABSTRACT

Brunskill, G. J. and B. W. Graham. 1979. The offshore sediments of Lake Winnipeg. Can. Fish. Mar. Serv. MS Rep. 1540: v + 75 p.

Samples of surface sediments of Lake Winnipeg were taken with dredges at 50 stations, and 1 to  $\bar{2}$ m cores were taken at 13 stations. Suspended and bottom sediments of the Red River were also obtained. Over 70% of off-shore sediment samples were silty clays or clayey silts. Coarser sediments near shore were not adequately sampled. The major minerals in sediment samples were quartz, feldspars, dolomite, calcite, illite, the montmorillonite group, and chlorite and kaolinite. Water content varied from 27% to 80%. Inorganic carbon ( $C_i$ , dolomite and calcite) varied from 83 to 3500  $\mu$ moles  $C_{1}/g$ , and organic carbon ( $C_{\rm O}$ ) varied from 166 to 3920  $\mu moles$  $C_0/g$ . Total phosphorus  $(\Sigma P)$  in sediment samples varied little (13 to 27 µmoles P/g) throughout the lake, whereas total nitrogen  $(\Sigma N)$  varied over a wider range (<7 to 286  $\mu$ moles N/g). Dilute acid or buffered NTA extracted 40% to 70% of EP in sediments, but the sediment sorbed orthophosphate from solutions at concentrations similar to those of Lake Winnipeg water. Ratios of  $C_0$ ,  $\Sigma N$ , and  $\Sigma P$ in sediments were not those expected for organic matter, and it is suggested that much of the  $\Sigma N$  and  $\Sigma P$  in Lake Winnipeg sediments is inorganic. Among the major elements of the sediments, Al was most strongly correlated with the abundance of the clay fraction, Ca and Mg were correlated with  $C_1$  (likely dolomite), and  $S_1$  was correlated with the large particle sizes (likely quartz and feldspars in sands and silts). Trace elements (V, Cu, Co, Cr, Ni, Be, Cd, Mo, As, Zn, Pb, Sr, Mn) in these sediment samples varied little with location or sediment depth, but small increases in Hg concentrations near the sediment surface of several cores of the south basin were noted. Major and trace element data are also given for Red River suspended sediment, and an estimate of elemental sedimentation rates is given for the south basin of Lake Winnipeg. Major element data are reported for interstitial water squeezed from dredge samples. Sulfate was usually depleted in these interstitial waters, and Ca, Mg, Na, K, CI, and Si were usually enriched, compared to overlying lake water.

Major basins of sediment accumulation in Lake Winnipeg are designated and described. The Saskatchewan, Cannibal, Mukutawa, Berens, and Black Bear Basins in the northern part of the lake are supplied with sediments primarily from shore and island erosion. The Red River Basin is provided with sediments from the Red River and shore erosion. Sedimentation rates for the Red River Basin ranged from 150 to 1970 g m<sup>-2</sup> yr<sup>-1</sup> over 1969-73, and sediment accumulation rates of 0.5 and 0.9 mm yr-1 were estimated from sonar penetration and Red River sediment supply rates. Sedimentation rates for  $C_0$ ,  $\Sigma N$ ,  $\Sigma P$  were computed, and were found to lie in the low part of the range for St. Lawrence Great Lakes sedimentation rates. Moment parameters for the particle size distributions of Lake Winnipeg sediment samples

were compared to St. Lawrence Great Lakes data. Lake Winnipeg sediments approach a normal distribution of particle sizes, compared to St. Lawrence Great Lakes sediments. This is because most of the sediments supplied to Lake Winnipeg are pre-sorted silts and clays derived from glacial Lake Agassiz sediments.

Key words: sediment geochemistry; particle size; cores; interstitial water; sedimentation rates; trace elements.

### RESUME

Brunskill, G. J. and B. W. Graham. 1979. The offshore sediments of Lake Winnipeg. Can. Fish. Mar. Serv. MS Rep. 1540: v + 75 p.

Dans le lac Winnipeg, des échantillons de la couche superficielle de sédiments ont été dragués à 50 stations et carottés à une longueur de 1 à 2 m à 13 stations. Ont aussi été prélevés des échantillons de sédiments et de matières en suspension dans la Red River. Plus de 70% des échantillons prélevés loin de la rive se composajent d'argile limoneuse ou de limon argileux. L'échantillonnage des sédiments plus gros situés près de la rive, n'a pas été convenablement fait. Les échantillons renfermaient surtout du quartz, du feldspath, de la dolomite, de la calcite, de l'illite, du groupe de la montmorillonite, de la chlorite et de la kaolinite. La teneur en eau a varié entre 27 et 80%. La concentration du carbone inorganique (Cj, dolomite et calcite) s'est située entre 83 et 3,500  $\mu$ moles C<sub>1</sub>/g et celle du carbone organique (C<sub>0</sub>) entre 166 et 3,920  $\mu$ moles C<sub>0</sub>/g. La concentration du phosphore total (EP) a été sensiblement la même dans tous les sédiments du lac (13 à 27 μmoles P/g) tandis que celle de l'azote total (ΣΝ) a varié fortement (7 à 286 μmoles N/g). Entre 40 et 70% du ΣP a été extrait à l'acide dilué ou au NTA tamponné, mais les sédiments ont absorbé les orthophosphates qui étaient à des concentrations comparables à celles que l'on trouve dans le lac Winnipeg. Les proportions de  $C_0$ , de  $\Sigma N$  et de  $\Sigma P$ n'ont pas été ce que l'on s'attendait à trouver dans de la matière organique ce qui laisse supposer que la majeure partie de <code>rN</code> et de <code>rP</code> des sédiments du lac Winnipeg est inorganique. Parmi les principaux éléments figurent Al, attribuable sans doute à l'importance de la fraction argileuse, Ca et Mg, liés au  $C_1$  (sans doute à dolomite), et Si, en provenance des grosses particules (sans doute le quartz et le feldspath du sable et du limon). Les échantillons renfermaient des éléments à l'état de trace (V, Cu, Co, Cr, Ni, Be, Cd, Mo, As, Zn, Pb, Sr, Mn) dans des concentrations analogues en dépit de leur provenance ou de la profondeur à laquelle ils avaient été prélevés. Dans certaines carottes du bassin du sud, de légères augmentations de la concentration de Hg ont cependant été observées près de la couche superficielle des sédiments. Des données sur les éléments pri cipaux et à l'état de

٧

trace présents dans les sédiments suspendus de la Red River sont aussi fournies, ainsi qu'une évaluation de la sédimentation de base dans le bassin du sud du lac Winnipeg. Les teneurs en éléments principaux sont celles de l'eau interstitielle des échantillons dragués et essorés. Par rapport à la couche d'eau supérieure, l'eau interstitielle renfermait moins de sulfates et plus de Ca, de Mg, de Na, de K, de Cl et de Si.

Les principals bassins d'accumulation des sédiments dans le lac Winnipeg sont recensées et décrites. Les bassins (Saskatchewan, Cannibal, Mukutaoua, Berens et Black Bear) situes dans la partie septentrionale du lac se chargent surtout de sédiments formés par l'érosion des rives et des îles. Quant au bassin Red River, les sédiments qu'elle renferme proviennent de l'érosion des rives et de cette rivière. Entre 1969 et 1973, les vitesses de sédimentation y ont varié entre 150 et 1970 g.m<sup>-2</sup>.an<sup>-1</sup>, et d'après des sondages et des données sur l'apport de matières par la Rouge, leur accumulation, entre 0.5 et 0.9 mm.an-1. Comparativement à celle des Grands lacs du Saint Laurent, la vitesse de sédimentation de  $C_0$ , de  $\Sigma N$  et de  $\Sigma P$  est faible. Si l'on compare la répartition granulometrique des sediments dû lac Winnipeg à celle des Grands lacs, il ressort que, dans le premier, cette répartition s'approche de la normale. Cela est dû au fait que la majeure partie des sédiments, en provenance du lac Agassiz glaciaire, sont constitués d'argile et de limon déjà répartis uniformément.

Mots-cles: géochimie des sédiments; granulométrie; carottes; eau interstitielle; vitesse de sédimentation' éléments à l'état de trace.

#### INTRODUCTION

Early explorers passing through the Lake Winnipeg watershed occasionally commented on the muddy waters of the lake and the Red River (Ross 1856, p. 9; Palliser's Journals of 1857-1860 in Spry 1968, p. 11, 88, 94; Upham 1890, p. 18E). The deposition and resuspension of this Red River sediment load in Lake Winnipeg were observed by Upham (1890), who was also the first to study rigorously the geomorphology of the glacial lake deposits around the south basin of Lake Winnipeg. Upham (1890) gave the name Lake Agassiz to this Late Wisconsinan and Holocene body of water, the sediments of which are now found over some  $518,000~{\rm km}^2$  of what is now Manitoba, Ontario, Saskatchewan, Minnesota, North and South Dakota (Elson 1967). Tamplin (1967) reviewed the history of Lake Agassiz research. Lake Agassiz clays and silts are now agricultural soils, and likely constitute the majority of the sediments carried to Lake Winnipeg by the Red River (and to a lesser extent, by the Winnipeg and Saskatchewan Rivers). These Lake Agassiz clays and silts appear to have been derived from Cretaceous shales and Quaternary glacial sediments to the south and west of the present Lake Winnipeg (Kushnir 1971) via rivers draining the Dakotas and as far west as the Rocky Mountains in Alberta (Teller 1976) during the Pleistocene. Sediment supply to the north basin of Lake Winnipeg in recent time via the Saskatchewan River appears to have been relatively less than the Red River sediment supply to the south basin of the lake. Upham (1890) noted that the north basin of Lake Winnipeg was relatively clear, compared to the south basin of the lake. Many early explorers commented on the trapping of Saskatchewan River sediment in Moose and Cedar Lakes (Mackenzie 1801, p. 115; Hopwood 1971, p. 85). Denis (1916, p. 123) indicated that the Saskatchewan River deposited much of its sediment load into Cedar Lake (about 80 mm/yr in some locations). Sediment supply to the north basin of Lake Winnipeg has been further decreased by the construction of numerous reservoirs on the Saskatchewan River from 1913 to the present (Report of the Saskatchewan-Nelson Basin Board, 1972). Lake Diefenbaker collects about 23 x  $10^6$  tonnes sediments yr<sup>-1</sup>, and Tobin Reservoir collects about  $15 \times 10^6$  tonnes yr<sup>-1</sup>). Franklin (1823, p. 43) found the waters of the north shore of Lake Winnipeg rather turbid, however, and told a Cree story which accounted for the muddiness of the lake and its name (Winnipeg = muddy or dirty water in Cree).

The literature on Lake Agassiz sediments is concerned largely with beaches, outlet structures, clay stability for road and building construction, and general studies concerned with the history of that expansive and oscillating glacial lake. Much of this literature is reviewed in Mayer-Oakes (1967). We have been unable to find any references to the thickness or characteristics of Lake Agassiz sediments in the present Lake Winnipeg basin. Teller et al. (1976) have mapped the thickness of glacial sediments and bedrock topography in southern Manitoba. Kushnir (1971) speculates that some laminated clay clasts

from dredged sediments from the south basin of Lake Winnipeg are subaqueous outcrops of glacial Lake Agassiz sediments, based partly on sonar reflections and clay mineralogy. The sediments carried by the Red and Assiniboine Rivers to the south basin of Lake Winnipeg are, of course, largely Lake Agassiz sediments.

Other studies on Lake Winnipeg sediments include Solohub and Klovan (1970), Solohub (1967), and Veldman (1969), who were concerned largely with nearshore and beach sediment movement, deposition, and particle size sorting. Wallace and Maynard (1924), Elson (1961), Ehrlich et al. (1955), Wicks (1965), Last (1974), and Teller (1976) gave chemical, mineralogical, and particle size data for Lake Agassiz sediments, which may be useful in studies of Lake Winnipeg sediments. Ward (1926) contributed an estimate of Red River sediment transport rates, particle size and chemical analyses of Red River suspended sediments, Red River bottom sediments, and Lake Winnipeg sediments near the mouth of the Red River. Bajkov (1930) briefly described and categorized surface sediments of Lake Winnipeg. There have been no synoptic chemical studies on Lake Winnipeg sediments, except for Allan and Brunskill (1977) which is part of this work.

We now give descriptive data for Lake Winnipeg sediment samples, interstitial water, and riverborn sediments. Our objectives were to provide a whole lake view of Lake Winnipeg off-shore sediment texture, chemistry, and sediment supply rates, as a background for future detailed studies on selected problems.

# METHODS

The CGS Bradbury (Department of Public Works, Selkirk, Manitoba) was utilized as a research vessel. Her draft of 1.8 meters prohibited sampling in many of the large, shallow water areas (5% to 10% of lake surface area) of Lake Winnipeg (Figs. 3 and 4). Sampling stations (Fig. 2, Table 1) were located with radar, radiocompass, sonar (20 khz), and ship speed from Canadian Hydrographic Service charts 6240, 6241, 6267 and 6251, which were accurate to a radius of 100 meters or better. Some stations were at sextant-located navigation buoys identified on the above navigation charts for the lake. Water and sediment samples from the Red, Assiniboine, and Winnipeg Rivers were taken from bridges near their mouths. Water and sediment samples from other rivers were taken near river mouths using small boats.

# SEDIMENT SAMPLING

Surficial lake sediments from 50 stations were taken 3-12 October 1969 with Ponar grabs, Shipek, or Ekman dredges, and transferred into polyethylene containers which were stored at  $4^{\circ}\text{C}$ . Cores were taken with a weighted 2 meter Benthos gravity corer (with a core catcher in soft sediments) from 13 locations (see Table 20), and were stored in an upright position on the deck of the

ship. The sediment was sealed in core tubes with overlying water while on the ship, and later stored at 4°C in a laboratory cold room. Analysis of surface sediments were done in 1969, but the cores were not extruded until early 1971. Cores were extruded with a piston, sliced into 5 cm intervals, dried and ground for storage in plastic vials. Core samples were also taken in 1975 by Dr. R. J. Allan using similar equipment, but the cores were sliced into 2 cm intervals immediately after arrival in the laboratory. Sonar transparent and sonar reflecting sediment layers were recorded on the instrument chart paper: thickness of sediment layers were read off the water depth calibrated scale. Comparison of sonar sediment thickness with core logs was not useful since sedimentation basin cores rarely penetrated more than 1 meter of sediment and cores would not penetrate the sediment in regions of sonar reflecting top sediment.

Suspended sediments in tributary rivers were taken with 10L van Dorn samplers, or with weighted 2L bottles which were allowed to fall through the water column. Bottom sediments were sampled with the above noted array of dredges. Suspended sediments in lake water were obtained by continuous-flow centrifugation of large (20-60L) volumes of surface water.

# ANALYTICAL METHODS

Water content of the sediment was determined by loss of weight upon drying at 110°C, and loss on ignition was determined by heating these samples at 900°C for 4-6 hours. For most Lake Winnipeg sediment samples, loss on ignition at 900°C includes decrepitation of calcite and dolomite, as well as combustion of organic matter and loss of hydration water. Heating sediment samples in 100°C increments over the range 100° to 1000°C allowed individual estimates of organic matter, carbonate minerals, and waters of hydration (see Dean 1974). More detailed differential thermal analyses of the clay fractions from these samples and other Agassiz clays are given in Kushnir (1971) and Wicks (1965).

Particle size composition of sediments (in terms of fraction of sand (2000-50  $\mu m$ ), silt (50-2  $\mu m$ ) and clay (2  $\mu m$  and smaller)) was determined by the sieve and pipette method of Jenning et al. (1922) and Robinson (1922). In a few samples, the centrifugation and settling method of Jackson (1956) was also used. We followed Folk and Ward (1957), Shepard (1954), and Carver (1971) to obtain sediment particle size graphic parameters.

Minerals in particle size fractionated and unfractionated sediments were identified by X-ray diffraction (Klug and Alexander 1954; Jackson 1956; Rex 1969) using a Philips diffractometer (PW 1010 Generator, PW 1352/10 Circuit panel, PW 1170/00 Automatic Sample Holder, AMR-3-202 Graphite Monochromator and Co or Cu radiation). Whole or fractionated sediment samples were cleaned with  $\rm H_2O_2$  and sonification, and pipetted onto 6 mm thick porcelain tiles for

X-ray diffraction. Known mixtures of quartz, feldspar and selected clay minerals were used for calibration to estimate semi-quantitative mineralogical composition of the samples. Major elements (Si, Ca, Mg, K, Na, Fe, Al and Mn) in sediments were determined with a Perkin Elmer 403 atomic absorption spectrophotometer following the sediment dissolution technique of Suhr and Ingamells (1966). Total carbon and nitrogen were determined on dried (110  $^{\circ}$ C), ground and well-mixed sediment samples with a Carlo Erba Model 1100 CHN-O analyzer (Hauser, 1973). Inorganic carbon (calcite and dolomite) was determined by the Stainton (1973) method as follows: a weighed quantity of dried ground sediment and the appropriate quantity of sulfuric acid were introduced into a glass ampoule which was then quickly sealed and autoclaved at 121°C for one hour. The liberated CO2 was subsampled and measured on a Fisher-Hamilton gas partition chromatograph. Carbonate mineralogy was determined according to Tennant and Berger (1957) and Dean (1974). Total phosphorus was determined according to Andersen (1976). Sulfur was determined on a few samples by the University of Manitoba, Department of Earth Sciences, using a Leco Analyzer. Color of moist sediment in cores was determined by comparison with a Munsell Soil Color Chart (1954).

Interstitial water from the surface (roughly 0-20 cm) sediment samples and cores was extracted within a month of sampling by squeezing samples or sections of the core for 2 to 4 hours at room temperature, under 20-80 p.s.i., using an apparatus modified from the design of Reeburgh (1967); pH was measured with a Radiometer PHM4 (glass and calomel electrodes), conductivity with a Radiometer CDM-2c, major cations by a Perkin Elmer 403 spectrophotometer and major anions according to Stainton et al. (1974).

Trace elements and some major elements were measured by radio frequency plasma emission spectroscopy (Allan and Brunskill 1977; see Tables 14 and 15). Sediments for this analysis were refluxed with HF,  $\rm HNO_3$ , and  $\rm HClO_4$ , evaporated to dryness, and redissolved in 4%  $\rm HNO_3$ . Mercury was determined by stannous sulphate reduction to arsine gas, complexation, and colorimetric measurement at 520 nm on a spectrophotometer.

Sediment phosphorus was extracted with 0.01 M nitrilotriacetic acid (NTA) according to Golterman's methods (Golterman 1977 and personal communication).

# RESULTS

# PARTICLE SIZE DATA

Sediment sampling station locations and water depth are shown in Table 1 and Fig. 2. Bathymetry of Lake Winnipeg is given in Figs. 3 and 4. Lake Winnipeg has a surface area of 23,750 km², a mean depth of 12 m, and a volume of 284 km³ (Brunskill, Elliott, and Campbell 1980). Table 2 gives particle size fractionation data for surface sediments at 50 stations in Lake Winnipeg. The sampling stations are biased toward the deeperwater localities because of the ship's draft, and therefore many of the sand-gravel-boulder sediments near shores, islands and reef localities

were not sampled. Coarse sand, gravel, and boulders were commonly found or observed at stations along the east side of the north basin of the lake (Stations 21, 17), near shore or islands (12, 31, 68), and at river mouths (1, 29). These coarse sediments were often composed of dolomitic and calcitic sands, gravels, and boulders on the north, west, south, and south-eastern shores of the lake, most islands, and shallow areas. Sands, gravels, and boulders from igneous rocks of the Precambrian Shield were common on the east margin of the lake, but were also observed all around the lake margin in lesser abundance. According to Kushnir (1971), these Shield-derived coarse sediments are largely composed of quartz, feldspars, hornblende, pyroxene, muscovite, weathered biotite, sericitic and chloritic schists. The high proportion of sand at Station 39 is likely related to the proximity and easterly continuation of Long Point (part of The Pas Moraine, see Davies et al. 1962) which is composed of mostly sand and gravel. Over 70% of the sediment samples are silty clays and clayey silts (Fig. 5). This distribution of silt- and clay-sized particles throughout the lake is given in Figs. 6 and 7. These data (Table 2, Figs. 6 and 7) are not directly comparable to the particle size data of Kushnir (1971), because Kushnir used different particle size boundaries for the clay and silt fractions (i.e., Kushnir used <3.9  $\mu m$  for clay size, where we used <2 μm).

# MINERALOGY

Table 3 gives the proportion of carbonate minerals found in selected Lake Winnipeg surface sediment samples. In general, sediments with >5% carbonate were near river mouths or near the south, west, and north shores. Dolomite was nearly always more abundant than calcite. No aragonite was detected, although we know this mineral occurs in pelecypod and gastropod shells of the lake. Larger amounts of dolomite occur in lake sediments along the north and west shore of the lake (Stations 33, 31, 27, 60B), where Ordovician (Red River, Stony Mountain, and Stonewall Formation) dolomites, limestones, and shales outcrop as cliffs on the lake shore and as islands or reefs in the lakes (Davies et al. 1962). Surprisingly large amounts of dolomite were also found in the sediments of Traverse Bay (Stations 7 and 8), under the low ionic strength water plume of the Winnipeg River, and Kushnir (1971) claims that these carbonates were from locally derived aeolian sandy silts which were deposited after glacial Lake Agassiz finally was drained. The estimates of the proportions of calcite and dolomite in Lake Winnipeg sediments derived from incremental heating to 950°C (Dean 1974) given in Table 12 appear to overestimate dolomite, compared to the X-ray diffraction data in Table 3. With the exception of the Sturgeon Bay area (Stations 64-66), where sediments rich in abraded mollusc shells were seen, carbonate minerals appear to be allochthonous in origin, and were most abundant in the silt and sand fraction of the sediment. Carbonate minerals were also present in the suspended sediments of the Red and Assiniboine Rivers, and in Lake Winnipeg suspended sediment (Table 9) during the ice-free season.

The non-carbonate fraction of our Lake Winnipeg surfacé sediment samples was composed largely of quartz, feldspar, illite, montmorillonite clays, chlorite and kaolinite (Tables 4 to 8). Quartz and feldspar were most abundant in the sand and silt fractions, and the clay mineral groups montmorillonite, illite, chlorite and kaolinite (undifferentiated) were more abundant in the fine silt and clay fractions. In a few samples we attempted to differentiate chlorite from kaolinite by heating glycerated slides, and we found that kaolinite usually exceeded chlorite in abundance. Kushnir (1971) studied many of our south basin Lake Winnipeg samples in more detail, and found a very homogeneous distribution of these clay minerals in the clay particle size fraction throughout the south basin of the lake (Table 28). Based upon X-ray diffraction and differential thermal analyses, he found that the montmorillonite clays were interstratified with illite, that the illite was being transformed into an abnormal montmorillonite, and that the dominant exchangeable cation on the montmorillonite was Ca. In the clay-size fraction of the surface sediments, this complex montmorillonite group was frequently the dominant identifiable mineral (Tables 7 and 8).

The Red River is likely the main source for all of these minerals in the south basin of Lake Winnipeg (Table 9), and samples of suspended sediment from the Red and Assiniboine Rivers yield X-ray diffractograms similar to the surface sediments of the south basin. Suspended sediments in the lake waters were also mineralogically similar to Red River suspended sediments and the lake sediments.

ORGANIC MATTER AND NUTRIENT ELEMENT CHEMISTRY OF SURFACE SEDIMENTS

Data on water content, loss on ignition, C, N, and P for Lake Winnipeg surface sediments are given in Table 10. Water content varied from a low of 27% to almost 80%. Sediment with high water content was usually composed of silts and clays, had relatively high concentrations of organic matter, and were usually found in the deeper depositional basins shown in Figs. 25 and 26.

The loss on ignition and total carbon ( $\Sigma C$ ) concentrations given in Table 10 are the sum of contributions by carbonate minerals, organic matter, and hydration water on clay minerals (for loss on ignition only). Since the ratio of inorganic (carbonate) carbon ( $C_1$ ) to organic carbon ( $C_0$ ) varies greatly,  $\Sigma C$  data are not directly meaningful. An independent gasometric measurement of  $C_1$  was obtained and subtracted from  $\Sigma C$  to estimate  $C_0$  in Lake Winnipeg sediments. Table 12 gives another estimate of the proportions of  $C_1$  and  $C_0$  in  $\Sigma C$  by the ignition method of Dean (1974), which agrees fairly well with Table 10.  $C_0$  varied from 166 to 3916  $\mu$ moles  $C_0$ /g of dry sediment. The lower values were associated with the largely inorganic sediments near the mouth of the Red River, and the northern and eastern margin of the north basin; higher values of  $C_0$  were obtained from sediment samples from near the mouths of rivers draining the Shield (Berens and Winnipeg) and in the narrows of the lake.

 $\text{C}_{\bar{1}}$  varied from 83 to 3500 µmoles  $\text{C}_{\bar{1}}/g$  (Table 10). Lower values of  $\text{C}_{\bar{1}}$  were found in sediments from the narrows of the lake and much of the deep water

areas of the north and south basin. Higher values of  $C_1$  were obtained from sediment samples from near the mouths of the Red and Winnipeg Rivers, and near the west and north shores of the lake, where dolomite and limestone cliffs occur (Baillie 1952).

Total phosphorus  $(\Sigma P)$  in Lake Winnipeg surface sediments varied little, from 13 to 27  $\mu moles$  P/g (Table 10). The lower values were found in sediments from stations along the northwestern shore of the north basin and at the mouth of the Winnipeg River. Higher values were found in sediments from stations along the northwestern shore of the north basin and at the mouth of the Winnipeg River. Higher values were found in sediments from the center of the north basin of the lake. The standard deviation of the mean  $\Sigma P$ concentration in Lake Winnipeg sediments is very small, considering the size of the lake and the differences in tributary drainages. These ΣP data in Table 10 are from acidification, combustion, and colorimetric methods described by Stainton et al. (1974) and Andersen (1976), whereas the  $\Sigma P$ data in Table 15 were determined by radio frequency plasma emission spectroscopy. Table 15  $\Sigma P$  data show the same limited variation in concentration, but the absolute values are lower, compared to  $\Sigma P$  data in Table 10. Concentrations of  $\Sigma P$  in suspended sediments of the Red and Assiniboine Rivers (Tables 17 and 18) are well within the range of the surface sediment concentrations (Table 10), but suspended sediments in Lake Winnipeg have higher concentrations of  $\Sigma P$  (Table 18) than the sediments.

Experiments were done in the laboratory to determine the labile or exchangeable fraction of P in sediments from the center of the north and south basins of Lake Winnipeg (Table 11). A gentle leach with NH $_4$ Cl released IO-12% of  $\Sigma$ P from these sediment samples, and dilute acid or NTA buffered to pH 7 removed 40-77% of  $\Sigma$ P.

These sediments sorbed orthophosphate from solutions at or above Lake Winnipeg PO $_4$  concentrations (see Brunskill, Schindler et al. 1980). Sorption of orthophosphate by sediments increased with increasing solution concentrations (1.6 to 9.6 µmoles/L) from 0.26 to 3 µmoles P/g sediment.

Total nitrogen ( $\Sigma N$ ) varied considerably from <7 to 286 µmoles N/g. Minimal values were obtained from sediment samples near the mouths of the Red and Winnipeg Rivers, and from stations on the eastern and northern margins of the north basin. Higher values occurred in samples from the deep water central and western areas of the north basin, Berens River mouth, Pigeon Bay, and the Narrows.

The mole ratio  $C_0\colon\Sigma N$  (Table 10) varied from 1.6 to 48. The extremely low  $C_0\colon\Sigma N$  ratios of 1.6-4 are partially due to analytical difficulties with samples containing a high proportion of  $C_1$  compared to  $C_0$ , and possibly NH, sorbed to clay minerals (Kemp and Mudrochova 1972; Byrnes et al. 1972).  $C_0\colon N_0$  ratios between 6 and 15 are representative of recently dead or living algae (Healey 1975), and  $C_0\colon N_0$  ratios >15 probably represent an increasing proportion of organic matter derived from terrestrial soils.  $C_0\colon\Sigma P$  ratios varied from 30 to 156, which is low compared to  $C\colon P$  ratios for other lake sediment

or seston data (Brunskill et al. 1971; Schindler 1976). The mole ratio  $\Sigma N:\Sigma P$  varied from 1.0 to 11.5, which is extremely low. Sewage has a N:P ratio of =11, and living plants have N:P of 10-40. As shown in Table 11, over 50% of sediment P cannot be extracted by acid, and likely is an allochthonous mineral form of P. There were no statistically significant linear relationships between  $C_0$  and  $\Sigma N$  (r=0.22, n=44),  $C_0$  and  $\Sigma P$ , (r=0.26, n=46), or  $\Sigma N$  and  $\Sigma P$  (r=0.24, n=47) for the sediment data in Table 10.

### MAJOR ELEMENTS IN SURFACE SEDIMENTS

Major element composition of selected Lake Winnipeg surface sediment samples is given in Tables 13 and 14. The data in Table 13 are results of metaborate fusion and flame atomic absorption determination of the elements, whereas the data in Table 14 are from acid refluxing of sediments and determination by radio frequency plasma emission spectroscopy. The sediment analyzed by these two methods are from the same station and dredge sample, but are different subsamples. The differences between the results are likely due to the sum of subsampling, different sample preparation (fusion v.s. reflux), and different analytical determination. In most cases, we have used the data set in Table 14 because it is larger.

The most abundant elements are Si and Al, which are components of the clay fraction of the sediments, and also feldspar and quartz minerals. Al was significantly (at  $\alpha=0.05$ ) and linearly correlated with the abundance of clay in the sediment [Log Al = 0.0244 + 0.2529 Log (% Clay),  $r=0.84,\;n=26$ ], but Si was inversely and weakly correlated with the clay fraction [Si = 12.9325 - 1.7352 Log (% Clay),  $r=-0.45,\;n=26$ ]. This latter finding implies that quartz and feldspars in the silt and sand fractions are an important control on the abundance of Si in these sediment samples.

Variations in sediment concentrations of Ca and Mg appear to be related to the abundance of carbonate minerals in these sediment samples. Ca from Table 14 was significantly related to  $C_i$  (carbonate carbon, in Table 10) according to the equation:  $C_i = 0.2792 + 0.5361 \ C_i$ ,  $C_i = 0.88$ , Csediments, we might expect Ca + Mg to be even better correlated with Ci. Figure 9 confirms this, and the obtained slope of the relationship is near unity for the molar relationship. The intercept of Fig. 9 gives some indication of the proportion of Ca and Mg that may be fixed in clay minerals and plagioclase feldspars. There was no statistically significant correlation (at  $\alpha = 0.05$ ) between Ca and % montmorillonite (Kushnir 1971; Table 28), which indicates that the relatively small carbonate fraction (Table 10) controls the variation of concentrations of Ca and Mg in these sediments, despite the much larger quantities of clay. was an inverse correlation between Ca and log (% Clay), (r = -0.42, n = 50), which implies that  $CaCO_3$  was more abundant in the silt and sand fractions of the sediment than in the clay fraction.

Na and K are most abundant in plagioclase and K-feldspar minerals, and illite (Tables 4 to 8) will likely carry some of the K. Iron, Mn and P are

components of both the detrital mineral phase and the flocculent rusty-orange colored surface layer (0-1 cm) of the sediment. Fe was significantly correlated with the abundance of clay in the sediment [Log Fe = -0.7246 + 0.3792 Log (% Clay), r = 0.75, n = 26], and was also correlated with  $\Sigma P$  (r = 0.89 for south basin sample and r = 0.83 for north basin, n = 21 and 18).  $\Sigma P$  was not correlated with the abundance of the clay size fraction (r = 0.31, n = 26). Total sulfur determinations were done on sediments from 4 stations in the central south basin, which gave an average of 11  $\mu moles$  S g<sup>-1</sup> dry weight (range = 7.2 - 13.8), and one station in the Narrows (Station 55) had 20  $\mu moles$  S g<sup>-1</sup>.

Most of these major element concentrations for Lake Winnipeg surface sediments (Tables 13 and 14) were similar to concentrations in Red River suspended sediment samples (Tables 17 and 18), except for an apparent small enrichment of Si in the lake sediments compared to river sediments. The ratio Ca:Mg in Red River suspended sediments was <1 (Table 18), whereas this ratio in the lake sediments varied considerably (Table 13), and suspended lake sediments have Ca:Mg ratios >1 (Table 18). The ratio Na: K in river suspended sediments was generally <1 (Table 18), whereas lake sediments varied greatly (Table 13), and lake suspended sediment Na:K ratios were >1 (Table 18).

# TRACE ELEMENTS IN SURFACE SEDIMENTS

Table 15 gives the results of analyses of Lake Winnipeg surface sediments for selected trace elements. With the exception of a few inexplicably high values (e.g. 1.34 µmoles Pb/g at Station 25) the variations in trace element concentrations were small. Sediment samples from near the mouth of the major rivers draining watersheds with agricultural and industrial activities were not detectibly enriched in trace elements (see Stations 1-4, 7 and 8, 25-27). In a following section on sediment cores, however, Hg was found to be above background levels in some cores (Table 24).

The variation of trace element concentrations is better viewed in proportion to some conservative parameter of the sediment. We chose Fe, Ti, Si, and Al as conservative elements in our work, because they are abundant in the sediment phase, relatively immobile in the well-oxygenated surface sediments, and they are unlikely to have been greatly affected by pollution sources. In Table 16 we give the average ratio of selected trace elements to conservative elements, or the slope of linear regressions of each trace element against each conservative element for 21 Lake Winnipeg sampling stations. This analysis indicates that V and Zn are more abundant, relative to Fe, Ti and Al, in south basin sediments, and that Ni, Cu, and P are more abundant, relative to Fe and Ti, in the north basin of the lake. Trace elements listed in Table 15 but not shown in Table 16 were not significantly correlated with the conservative elements used here. Some of these correlations were highly significant (see Figs. 10-16 for relationships with Ti) and can be improved by subdividing the data set into the major lake basins (south basin, narrows,

and north basin). Similar relationships are shown for trace elements against Fe (Figs. 17-23) in which it appears that Ti is a better predictor of P (Fig. 16) than is Fe (Fig. 23).

Trace element concentrations in Lake Winnipeg sediments (Table 15) are similar to comparable data on Red River suspended sediments (Table 17). As and V are higher in concentration in the river sediments, and Sr and Ni are lower, compared to Lake Winnipeg sediments (Table 15).

RATES OF SUPPLY OF MAJOR AND TRACE ELEMENTS TO THE SOUTH BASIN OF LAKE WINNIPEG

The chemical data for suspended sediments from the Red River downstream from the City of Winnipeg (Tables 17 and 18) are used to estimate the composition of the mass of sediments carried to the south basin of Lake Winnipeg in a year. The samples of river suspended sediments in Tables 17 and 18 were taken during high spring discharges, when suspended sediment concentrations were relatively high (see Table 9). From this rather small data base, we calculated the annual rate of supply of major and trace elements in particulate form to the south basin of Lake Winnipeg (Table 19), utilizing complete seasonal 1969-1974 water discharge and suspended sediment concentration data from Water Survey of Canada (1969-1974).

The annual supply of each element was divided by the area of the 8 m contour (our estimate of the zone of fine sediment accumulation) of the south basin of Lake Winnipeg (see Fig. 3, and Brunskill, Elliott and Campbell 1980) to estimate an average sediment supply rate for each element. This elemental rate of supply is an estimate because only the Red River supply is considered, and supply from the Winnipeg River, other small rivers, nearshore and shore erosion is not estimated. Although annual discharge of the Winnipeg River is 4 to 10 times greater than the discharge of the Red River, its annual suspended sediment mass is usually less, due to its largely Precambrian Shield watershed (Brunskill, Elliott and Campbell 1980). During high discharge years (1969, 1974), the Red River transports nearly an order of magnitude more sediment ( $\simeq 3 \times 10^6$  tonnes yr $^{-1}$ ) than the Winnipeg River, but in low discharge years (1973) the two rivers transport nearly equal amounts of suspended sediment (=0.2 x 106 tonnes  $yr^{-1}$ ). The large range for the estimated sediment supply rates is due to variations in annual discharge of water and suspended sediments. If this sediment has an average density of 2.5 tonnes m $^{-3}$  the sedimentation rate of 1170 gm $^{-2}$  yr $^{-1}$  is equivalent to about 1 mm yr $^{-1}$  in the area of the 8 m contour of the south basin of the lake. This sedimentation is primarily comprised of clay and silt particles which contain the major elements Si, Al, Mg, Ca, K, Fe, Na, and Ti, in order of abundance. Among the trace elements, sediment supply rates of Zn, Sr, Cr, and V are an order of magnitude greater than other trace elements (Table 19). Much of the coarse sediment near the mouth of the Red River is transported by bedload movement, and is not represented in this calculation.

# CORES OF LAKE WINNIPEG SEDIMENTS

Location and general description of 14 cores of Lake Winnipeg sediments are given in Table 20. Despite the long storage time, many of the vertically-stored cores still had lake water over the sediment surface, and had about the same % H<sub>2</sub>O in the surface 15 cm as the dredged samples (see Table 10). Maximum water content in surface sediments was 70-75%, whereas 25-50% water was found below 90 cm sediment core depth, or in association with silty sands in cores from Traverse Bay (Station 8) and Pigeon Bay (52). Many of the core bottom sediments were composed of conchoidally fractured or granular clay, which fell apart on the extrusion trough, as if the sediments were dry, yet these clays were 30-50% water.

Data for inorganic  $(C_1)$  and organic  $(C_0)$ carbon,  $\Sigma N$ , and  $\Sigma P$  in 5 cm intervals for these cores are given in Table 21. Data for the surface 0-20 cm of the core samples can be compared to data from the dredged sediment samples in Table 10. Although some of the data in Table 10 and 21 are similar, some show great differences, which seem likely to be due to sample location and sediment variation.  $C_i$  (as dolomite or calcite, see Table 3) was highest in Traverse Bay (Station 8) and lowest in the central north basin (Station 35) and narrows (Stations 54, 14) of the lake. Intermediate values were found in sediments from near the mouth of the Red and Saskatchewan Rivers (Stations 4 and 25).  $C_1$ variation with depth was small in most cores, but a two- to five-fold variation occurred in the cores from the narrows (Stations 14 and 54) and at the center of the north basin (Stations 35-2).

Co in the core profiles showed little geographic variation throughout the lake (with exception of Station 59 core, which seems aberrantly low), and about a three-fold variation occurred with depth. Cores in the narrows (Stations 15 and 54) and Traverse Bay (Station 8) indicated a decrease in  $C_0$  with depth.  $\Sigma N$  was generally higher in sediments from the center of the north basin (Station 35) and the northern narrows (Station 54). Variation with depth for EN was usually less than two-fold, with only slight indications of a decrease with depth. The variation of P was small (two-fold) throughout the lake, and there was no trend in  $\Sigma P$  concentration with depth. The highest average  $\Sigma P$ concentration occurred off the mouth of the Saskatchewan River (Station 25).  $C_0:\Sigma N$ ,  $C_0:\Sigma P$  and  $\Sigma N:\Sigma P$  were less variable than the same data from dredged surface sediments (Table 10), but still very low values occurred. The often chaotic variation with depth seems to be imposed on a slight decrease in these ratios with increasing depth.

Major element chemistry for selected Lake Winnipeg cores is given in Table 22. These data are in general agreement with similar data for dredged sediments (Tables 13 and 14). Ca and Mg are high in Traverse Bay core (8), and are nearly equimolar, reflecting their association as dolomite (see Table 3). Little variation with core depth occurred for Ca and Mg, except in cores from the mouth of the Red and Winnipeg Rivers

(Stations 04 and 08), where these elements increased with core depth. There was little variation of Na and Al with depth. Fe was low in sediments from the Traverse Bay core (08), and high in cores from the north basin and central south basin. Fe tended to decrease in concentration with sediment depth at core stations 08 and 61, but opposite trends were also evident (Table 22). Mn increased with sediment depth at core stations 04 and 25, and decreased with depth at stations 08 and 35. Ti varied little throughout the lake and with sediment depth. As in Table 21,  $\Sigma P$  varied little throughout the lake and with sediment depth.

Trace element data for Lake Winnipeg cores are given in Tables 23 and 24, which are in reasonable agreement with the dredged sediment data in Table 15. The variation of Cd, Co, Ni, Sr, Cr, and Zn in core samples throughout the lake and with sediment depth was less than a factor of three. Lead varied four-fold in these samples, but no pattern could be discerned. Beryllium and Cu were found in lower concentrations in Traverse Bay (08), and higher values were found in the central north and south basin cores. Vanadium was found in lower concentrations in Traverse Bay (Station 08), and higher values were found in the south basin (Stations 04, 10, 57, 59, 61). Arsenic was higher in concentration in the central south basin (Stations 10, 61, 57, 59) and northern narrows (Station 54) compared to other cores. The variation of the Mo values seems chaotic. Because the variation in element concentrations is small, the few trends with depth are likely not meaningful without more detailed sampling. Another set of cores were taken in 1975 by R. J. Allan and staff (see Fig. 24 for core locations, and Table 24), which were sampled at 2 cm intervals. As and Mo show concentrations and depth profiles similar to those in Table 23. The concentrations of Hg in the sediments at the top of these cores are about two to three-fold greater than at the bottom of the cores (Table 24, cores 1, 3-5, 8 and 9 in Fig. 24).

# INTERSTITIAL WATER OF SURFACE SEDIMENTS

Interstitial water from 4°C stored dredged surface sediments was squeezed from homogenized subsamples on the ship or in the laboratory at room temperatures within 2 months of sample collection. The data are given in Table 25, where interstitial water element concentrations are compared to concentrations of the element in lake water (usually from 1 m water depth, see Brunskill, Campbell and Elliott 1979). In most cases, concentrations of Ca, Mg, Na, K, Cl, and Si were greater in interstitial water than in overlying water. Si was 5-10 fold enriched in interstitial water, compared to lake water. Na was found in very high concentration in interstitial water from Station 31 in the northwest sector of the north basin.

# DISCUSSION

### SEDIMENTATION BASINS AND SEDIMENT SOURCES

Based upon bathymetry (Figs. 3 and 4), sediment particle size (Figs. 6, 7, 8), qualitative  $\frac{1}{2}$ interpretation of sonar penetration into sediments, and textural stratigraphy of cores, we designate the major basins of fine sediment accumulation in Lake Winnipeg (Figs. 25 and 26). The large, northerly Saskatchewan Basin receives sediment from the Saskatchewan River, and shore and island erosion of the north and northwesterly shores of this part of the lake (Fig. 26, see also Back 1836, p. 60). Sonar profiles from the center of the Saskatchewan Basin to the north, west or to the east revealed a gradual transition from a 4-15 m thickness of sonar-transparent sediment overlying a more dense sonar-reflecting layer (the center of the sedimentation basin), to a thin, irregular or absent sonar-transparent sediment overlying irregular ("spiky" or "stepped") sonar-reflecting sediment or bedrock (nearshore and off shore areas of erosion of sediment transport). Surface sediments of the central and northern regions of the Saskatchewan Basin are clayey silts, whereas the southern part of the Saskatchewan Basin has largely silty sediments (see Figs. 6 and 7). The boundaries of the Saskatchewan and other basins proposed here are approximate, and require verifification by increased sample coverage.

The Saskatchewan Basin is separated from the smaller Cannibal and Mukutawa Basins by a chain of sand and gravel islands (Georges, Little Georges, Big Sandy, Little Sandy, Cannibal Islands, see Hydrographic Chart 6241, or Fig. 25), shallow areas, and a slight sandy rise between Long Point and this island-shallows barrier (Fig. 8). The delineation of the Cannibal and Mukutawa Basins is based on sonar records and bathymetry alone. The major source of sediment to these basins must be island and shore erosion because there are no large rivers in this area. Sonartransparent sediment thins gradually to the north, east and west of the Cannibal Basin. The Mukutawa Basin may reveal a curious sediment texture when it is more extensively sampled, as sediments to the east (Stations 19 and 20) are bedrock, gravels and sands, with fragments of a compact (sometimes laminated and/or vesicular) silty clay. The vesicular (honeycomb) silty clay samples often had adult beetles in the cavities, but it is unlikely that they formed the hole, because hole diameters varied from 1 mm to over 1 cm. Sonar profiles east and north of the Mukutawa Basin showed a very "spiky" bottom with little and irregular amounts of sonar transparent sediment. We interpret this condition to represent areas of sediment erosion and transport, and little recent sediment accumulation. George's Island sediments (Station 17, see Table 2) are mostly sand and silts, and are probably transported to both basins. The Mukutawa and Popular Rivers may contribute some sediment to the Mukutawa Basin, but the watershed is well-forested Precambrian Shield, and their water discharges are small.

The Berens Basin (Fig. 25) is supplied with sediment largely from erosion of Reindeer, Georges, and Berens Island, and possibly from shallow water sediments in Sturgeon Bay (Figs. 25 and 26). No large, sediment laden rivers enter the lake in this region. Sonar transects to the northeast, east, and southeast of the Berens Basin thin out abruptly to a hard, spiky, or stepped sonar-reflecting sediment which was either sand and gravel, or a hard, compact, often laminated and vesiculate, silty clay and clay (Stations 50, 50B, 50C, 50D). The 98% clay sediment samples from Station 50D (see Table 2, Fig. 7) are an example of this latter sediment, and it is probably a Lake Agassiz deposit being eroded and transported into the Berens Basin. Berens Basin surface sediments are largely silty clays (Figs. 6 and 7).

The Black Bear Basin (Fig. 26) probably receives most of its sediment from shore and nearshore erosion to the east, and island and shallowwater sediment erosion from the west. Sonar profiles within this basin indicate a variety of sediment types, from silty sand sonar-reflecting sediment, to sonar-transparent clayey silts with relatively high amounts of organic matter (Figs. 6 and 7, Table 10, Stations 15, 52, 53, 54, 55; and Table 2). Spiky, irregular sonar profiles occurred when approaching the east shore, indicating bedrock, coarse glacial till, or erosional surfaces. Black Bear Basin sediments, and most of the narrows south to Hecla and Black Island, are probably subjected to some sorting due to currents generated by wind-generated seiches and storm surges (Einarsson and Lowe 1968; Hamblin 1976; Kenney 1979). Based on bathymetry alone, sediments of the Black Bear Basin may be transported into the Berens Basin via the narrow channel between Berens Island and Berens River. Other small depositional basins occur in the narrows (such as in Washow (Humbug) Bay, to the west of Station 14), but most of the deeper areas of the narrows south of the Black Bear Basin are areas of erosion by wind-generated currents. Stations 13, 13B, and 13C have hard, sonarreflecting sediments, gravels, or sediments that could not be sampled with our equipment (see Kushnir 1971; Table 3, Station 713). Net suspended sediment transport should be from south to north, from the south basin to the north basin.

The Red River Basin (Fig. 26) is supplied with sediment from the Red River (see Table 19), shore erosion (Veldman 1969), and to a lesser extent, the Winnipeg River. Kushnir (1971, Fig. 11) delineated two basins in this area, based upon clay content. Large amounts of sediments are also deposited in the Red River Delta (Netley Marsh) and off the mouth of the Red River. Sonar transects in the Red River Basin showed an apparent thickness of 7-8 m of sonar-transparent sediments (60-80% clay, see Fig. 7) in the center of the Basin, to essentially zero thickness (sand and gravel) in the nearshore zones to the north, east, and west. At nearshore stations 12, 58, 62, 60B, and 9, we found either sands and gravels, or a compact, laminated and/or vesicular silty clay (Figs. 6, 7, 8; Table 2; see also Kushnir 1971). Veldman (1969) describes south basin shore erosion and littoral sand transport, and Solohub (1967) and Solohub and Klovan (1970) studied the

usefulness of computed sediment particle size parameters as indicators of deposition environments in a Grand Beach lagoon, bay-mouth bar, and nearshore zone of the southeast shore of Lake Winnipeg. These latter authors found that graphic and moment parameters of grain size distributions failed to identify known environments of deposition (beach, aeolian dunes, channels, deltas, off-beach deepwater, reef, and lagoon), whereas factor analyses (Klovan 1966) did sort the samples into different types and energies of deposition.

With some exceptions, most of the lake bottom outside the depositional basins named here (Figs. 25 and 26) are areas of sediment erosion and transport. Much of the eastern margin of the lake bottom has sediments characteristic of old glacial lake sediments which are now being eroded and transported into the deeper basins. With exception of the southeast shore of the south basin, the east shoreline and terrestrial watershed of the lake is of igneous bedrock and does not appear to be contributing large amounts of sediment to the lake basins. The more erodible northern, western, and southern margins of the lake, however, are likely contributing a fairly large amount of sediment to the lake basin. If most of the Saskatchewan River's sediment load is deposited in Cedar and Moose Lakes (and recently the Grand Rapids Reservoir), which are part of the Saskatchewan River Delta (see Franklin 1823, p. 46-47; Mackenzie 1801, p. 115; Hopwood 1971, p. 85; and Denis 1916, p. 123), then much of the inorganic sedimentation in the Saskatchewan, Cannibal, and Mukutawa Basins must be derived from shore and island erosion. In the north basin, wind fetch is >200 km, wind velocities reach 30 km/hr, and we observed >1 m waves. Under these conditions, and by comparison to shore erosion studies at South Indian Lake (Newbury et al. 1978), erosion on the north shore of the north basin could be  $10-20\ \mathrm{m}^3$  of fine sediment per meter of shoreline. We estimate that sedimentation rates of 140 to 2200 g  $\rm m^{-2}~\rm yr^{-1}$  could occur over the area of the 16 m depth contour from erosion of the northern and western exposed shorelines of the north basin (based on the data and methods of Newbury et al. 1978, and Kachugin 1966). The northern and western margins of the north basin of Lake Winnipeg were often very turbid, from the shoreline out to 8-10 km offshore, in contrast to the relatively clear water in the center of the basin (see Figs. 4-9 in Brunskill, Schindler, et al. 1980). Considerable shore erosion in the south basin has been observed by Veldman (1969), but the dominant sediment source for the Red River Basin is likely the Red River (Table 19). The effect of the wind in controlling the path of the sediment-laden Red River plume was often observed on the ship and from the air. Strong on-shore winds also resuspended shallow water sediments and/or transported fine sediments into deeper waters (Sheng and Lick 1979).

# IMPLICATION OF PARTICLE SIZE ANALYSES

Dickas (1970), Upchurch (1970), Sly (1977), and many others have successfully used grain size analysis of sediments to deduce the mode and energy of deposition. Solohub and Klovan (1970)

found that the graphic and moment parameters (see Table 26) of sediment particle size distributions were not useful in distinguishing known environments of deposition in their study area (a lagoon and beach area near Grand Beach, on the east shore of the south basin of Lake Winnipeg). We also found that these parameters for the whole of Lake Winnipeg (Table 27) were of limited use in interpreting the energy associated with sediment transportation or deposition, at least in our set of samples. The inclusion of more samples of coarse sediments from nearshore environments might improve the range of these computed parameters, and facilitate more general comparisons. Our sediment samples ranged from about 2 mm sands to very fine clays, whereas many beaches, shorelines and sublittoral zones of Lake Winnipeg (which we did not sample) are composed of sorted and unsorted pebbles, cobbles, and boulders. Therefore, the following comments about particle size distributions and computed graphic measurements are relevant to offshore sediment areas only.

Mean particle size ranged from 3.7 to 10.8  $\phi$  with a south basin average of 8.13  $\phi$ , and the narrows and north basin averaged 7.5  $\phi$ . The standard deviation of  $X_{\varphi}$  was moderate to low (0.8-1.5) for the clay sediment samples, quite variable for the silty sediments (=2), and moderate for the sands (1.5-2) (see Fig. 27). Although the scatter at 6-7  $\phi$  (in Fig. 27) is great, the line indicated is similar to the shape of the same plot for Lake Erie and Ontario (Thomas et al. 1976; Thomas et al. 1972), except that our range of  $\phi$  is restricted (>3  $\phi$ ). Similar values of standard deviation were found in Lake Winnipeg, compared to the Erie and Ontario data. Figure 27 is interpreted as evidence for better sorting of the clays and sands, and lesser sorting of some of the silty sediments. The data were segregated into the major areas of the lake (south basin, narrows, north basin), and the same trend was observed as in Fig. 27.

Figure 28 shows the relationship between skewness of the sediment particle size distributions and mean particle size for 50 Lake Winnipeg sediment samples. Clayey sediments tended to be negatively skewed, and sandy sediments were positively skewed. This suggests that the greatly variable silt fraction of the sediments (Fig. 6) is present in both the sandy (near shore) and clayey (deeper) sediment zones. The trend of the data in Fig. 28 is similar to data plots for the St. Lawrence Great Lakes (Thomas et al. 1976; Thomas et al. 1972; Thomas et al. 1973; Sly 1977; Thomas and Jaquet 1975) but the values of skewness are closer to zero for the Lake Winnipeg sediments. Maximum positive values of skewness were for sandy sediments from near the mouth of the Red River (Station 1), near Hecla Island (12), near Georges Island (17), and near Eagle Island (31). Maximum negative values of skewness were found in clayey sediments from the center of the Red River Basin (Stations 59, 59B, 60, 60B), the Berens Basin (48, 64), the western margin of the Saskatchewan Basin (25), and the northern part of the Black Bear Basin (52) (see Figs. 25 and 26).

The distribution of particle sizes in Lake Winnipeg sediments showed a slight trend toward graphic leptokurtosis, and most values were near +1 (Table 27, Fig. 29) over the range 3-12 φ. Moment kurtosis of Lake Erie and Ontario sediments varied from -2 to +20 (Thomas et al. 1972, 1976),

whereas Lake Winnipeg samples gave moment kurtosis values of 0 to 14. The relationship of graphic kurtosis to skewness is shown in Fig. 30, which indicates that the Lake Winnipeg offshore sediment sample particle size distribution is closer to the normal distribution curve (Carver 1971) than sediments of the St. Lawrence Great Lakes (cf. Thomas et al. 1972, 1976). The three samples with anomalously high values of kurtosis in Figs. 29 and 30 are from Station 12 (near Hecla Island, a source of uniform sands), Station 01 (mouth of the Red River, likely bedload sands) and Station 31 (near the NW shore of the north basin, having limestone and dolomite cliffs). Sediment particle size distributions with larger standard deviations had generally lower values of kurtosis, compared to sediments having lower standard deviations (Fig. 31).

The significance of the results of the calculation of the four graphic parameters of Lake Winnipeg sediment particle size distributions is not immediately obvious. In comparison to the St. Lawrence Great Lakes sediment data (Thomas et al. 1972, 1973; Damiani and Thomas 1974), Western Lake Geneva (Vernet et al. 1972), and Lake Chad (Mothersill 1975), the Lake Winnipeg sediments appear to have small values of skewness and leptokurtosis. This implies bimodal populations of coarse silt and fine clay, with a deficiency of fine silt-sized particles. Yet the standard deviations about mean of the samples are similar to the above cited data, which would be interpreted as an indication of poor sorting in a lower energy environment. Due to Lake Winnipeg's shallowness (Z = 12 m), large surface area ( $A_0 = 12 \text{ m}$ ) 23,745 km) and fetch, and exposure to strong prairie winds, the entire lake bottom is likely subjected to rather energetic water movements resulting from seiches, wind-driven current, and river plumes (Hamblin 1976; Einarsson and Lowe 1968). We propose that the apparent low sorting (high standard deviation about  $\overline{X}_{\Phi})$  of Lake Winnipeg sediments is due to the lack of a low energy (deep) sedimentation basin to act as a reservoir for the sorted sediments. There is likely more than adequate energy to sort sediment particle sizes, but there is an inadequate gradient of depth to allow the products of sorting to be segregated. Despite this high standard deviation, the sediments still approach a normal distribution of particle sizes. seems likely due to a similarity of source materials, i.e. the surrounding glacial Lake Agassiz sediments. Last (1974) and Teller (1976) indicate that offshore Lake Agassiz sediment, south of Lake Winnipeg, averages 79% clay, 20% silt, and 1% sand, with the proportions of silt and sand increasing with proximity to glacial lake deltas, beaches, and glacial moraines and eskers. Wallace and Maynard (1924), Johnston (1934), Elson (1961, 1967), Wicks (1965), Smith et al. (1967), Last (1974) and Teller (1976) describe these fertile and erodible Agassiz sediments, glacial deposits and soils.

# REACTIVITY OF THE SEDIMENTS

Kushnir (1971) and Wicks (1965) describe in detail the mineralogy of glacial Lake Agassiz sediments in the Red River watershed and in the present Lake Winnipeg. Much of the clay fraction

of both these sediments was shown to be montmorillonite and a randomly interstratified, dioctahedral illite-montmorillonite (Table 28). The illite appears to be in transition to a montmorillonite structure, and the montmorillonite group of clays has Ca as the dominant exchangeable cation in the lake sediment (Kushnir 1971), whereas the water-shed Agassiz sediments carry Ca or Mg as exchange-able cations. Suspended and bottom illitic clays in Lake Winnipeg are thus likely to take up Ca from lake or interstitial water. Computation of the appropriate ion activity products (according to Garrels and Christ 1965) indicates that the detrital dolomite and calcite should be dissolving in lake water and interstitial water (Brunskill, Campbell and Elliott 1979). Table 3 indicates that carbonate minerals were abundant near their source (rivers, island or shore erosion), but most deeper water sediments had little or undetectible amounts of carbonate minerals. Either the carbonates (probably mostly in the silt and sand sizes) are not transported to the major depositional basins (Figs. 26 and 27) or they are dissolved in these deeper sediments. Waters of the Winnipeg River are greatly undersaturated with respect to calcite and dolomite, and the relatively large amounts of these minerals in Traverse Bay surface sediments should be dissolving. The presence of carbonate minerals in Traverse Bay sediments under dilute water suggests that the rate of supply of carbonate minerals is greater than the rate of dissolution. In the higher ionic strength waters of Lake Manitoba, sedimentary calcite may be authigenic, according to Last (1979).

Lake Winnipeg bottom sediments (and likely suspended sediments) appear to act as a sink for phosphorus (Table 11), and about 50% of  $\Sigma P$  in the sediment is labile (leachable by dilute acid). pP may be sorbed or precipitated with the relatively abundant exchangable Ca of the sediment similar to soil reactions studied by Racz and Soper (1967). With increasing concentrations of P in solution, the sediment sorbed more P from the solution. Similar results were obtained by Harter (1968) for a eutrophic lake sediment in Connecticut. The sediment did not appear to be saturated with sorbed P, according to these experiments. Despite both sediment and algal uptake of P from the lake water, molybdate-reactive phosphorus remained in high concentration (0.5-3 μmoles TDP/L) throughout most of the lake waters (Brunskill, Schindler et al. 1980). In the unlikely event that P supply to Lake Winnipeg were greatly decreased, these sediments could act as a source of P to the extent of perhaps 10% of EP in clay-rich sediments (NH<sub>4</sub>Cl extract in Table 11). Numerous studies on the movement and binding of P in Red River Valley soils (Racz and Soper 1967; Lewis and Racz 1969; and references therein) have been done, and they indicate that P is firmly and inorganically bound with Ca, Al, and Fe in soil profiles.

The proportions of organic carbon ( $C_0$ ), total nitrogen ( $\Sigma N$ ) and phosphorus ( $\Sigma P$ ) in Lake Winnipeg sediments are not those expected for degradation products of terrestrial and aquatic biota (Table 10). The variation of  $C_0$ ,  $\Sigma N$ ,  $\Sigma P$  with sediment depth in the core samples (Table 21) does not show the usual decrease with depth attributed to diagensis of organic matter (e.g. see Kemp (1971)). We propose that many of the very low  $C_0$ :N ratios are due to sorption of NH4 and other N species on

the abundant clay fraction of the sediment. In the south basin of Lake Winnipeg, all sediment samples with low  $C_0:N$  ratios were from the center of the Red River Basin (where the sediment is 60-80% clay, see Fig. 7), whereas  $C_0:N$  ratios >10 were usually found within 10 km from shore. This relationship was completely reversed in the north basin of the lake, however, as all low  $C_0:N$ sediment samples were from nearshore areas, and  $C_0:N$  ratios >30 were found in the sediments of the Saskatchewan Basin. Kemp and Mudrochova (1972) found that a fraction of EN in Lake Ontario sediments was exchangeable and fixed NH<sub>4</sub>, and that illite (see Table 28) was the clay mineral likely to immobilize NH4 (Stevenson and Dhariwal 1959). In a similar manner, much (=50%) of the Lake Winnipeg sediment P appears to be in a tightly bound, possibly crystalline form (Table 11). Co: P and EN: P ratios for sediment samples depend largely upon variations in concentrations of  $C_0$  and  $\Sigma N$  , because  $\Sigma P$  is relatively uniform throughout the lake. There is insufficient Co and EN for the P concentration of these sediments, relative to the composition of plant material (where C:P  $\approx 85-260$ , N:P  $\approx 10-40$ ). Williams et al. (1976) partitioned Lake Erie sediment phosphorus into allochthonous apatite-P (in nearshore silts), nonapatite inorganic-P, and organic-P (in deep water clays).

# SEDIMENTATION RATES AND ELEMENTAL SEDIMENT SUPPLY RATES

Our estimate of sedimentation rate (Table 19) for the Red River Basin (Fig. 26) is in the same range as estimated basin sedimentation rates for Lakes Erie and Ontario (Table 29). It seems likely that the Red River Basin sedimentation rate is at least twice the values given in Tables 19 and 29, since lake shore erosion, atmospheric (Teller 1972) and Winnipeg River contributions are not represented. If post-Agassiz or early Lake Winnipeg sedimentation began about 8000 years B.P. (Teller 1976; Teller and Fenton 1979), and if the sonar-transparent Red River Basin sediment thickness of 7-8 m represent this period of the lake's history (see also Kushnir 1971), then the average sedimentation rate in this basin would be 1.0 mm yr-1. The southern and western lobes of the Saskatchewan Basin (Fig. 25) had sonartransparent sediment depths of 12-15 m, which represents about 1.5 mm yr-1 accumulation rate.

Ringrose (1975, and personal communication) provided evidence indicating that the south basin of present-day Lake Winnipeg was dry during 7000-8000 yrs B.P., just before the final draining of Lake Agassiz to the north. Our Shipek sediment samples from nearshore stations along the northeastern shore of the south basin (Station 9, 6.4 m water depth, see Fig. 2) were compact, laminated, and sometimes vesicular silty clays and sandy clays. Some of the larger holes or tubes (1-10 mm in diameter) in the hard sediment were occupied by living adult beetles and fine organic matter. These samples could represent desiccated Lake Agassiz sediments with fossil root casts from terrestrial vegetation. Similar compact vesicular silty clays were found at Stations 50, 50B, 50C, and 19, along the eastern margin of the north basin, ranging in water depth from 5 to 16 m (Fig. 2)

which would indicate a very low water stage for the north basin during this time. In any case Ringrose's (1975) hypothesis suggests that Lake Agassiz and post-Agassiz aquatic sedimentation may not have been continuous in the present Lake Winnipeg basin. Last and Teller (1979a, b) and Last (1979) interpret episodes of lake desiccation from sediment cores of adjacent Lake Manitoba, and they have radiocarbon dates of 9500, 8500, and 4500 yrs B.P. for these events. Their description of desiccated "marker horizon" (low % water) sediments is similar to our observations of "conchoidally fractured" or granular clay found at the bottom of most of our cores (Table 20). In Lake Winnipeg cores, this relatively dry sediment occurred at sediment depths of 50 to 110 cm in the south basin, narrows, and north basin, and represents sediments deposited 600 to 1200 yrs B.P. based on our speculative, indirect sediment chronology (1000-2000 g m $^{-2}$  yr $^{-1}$ ) mentioned above. In the Lake Manitoba cores, the top-most horizon of desiccated sediment occurs at 3 m sediment depth, and  $^{14}\mathrm{C}$  sedimentation rates are  $100-750 \text{ g m}^{-2} \text{ yr}^{-1}$  (Last and Teller 1979).

Ward (1926) gave an estimate of annual suspended sediment transport via the Red River to the south basin of Lake Winnipeg. Based on records from 1916 to 1926, he estimated that about  $1.712\ x$  $10^{12} \ \text{g yr}^{-1}$  of sediment was added to the south basin. If this sediment mass was deposited over the area of the 8 m contour in the south basin, it would be 986 g m $^{-2}$  yr $^{-1}$ , which is within the range of the 1969-73 data in Table 19. Ward (1926) also analyzed suspended and bottom sediments from the Red River in the City of Winnipeg and 4 miles off the mouth of the Red River in Lake Winnipeg. His major element data for these samples are in fair agreement with our data in Tables 17 and 18. A comparison of our 1969 data and Ward's particle size data for Red River and Lake Winnipeg sediments (Table 31) indicates that the river sediments in 1969 had more clay, and the 1969 Lake Winnipeg Station 4 has less clay and more sand, compared to the 1926 data. This change is consistent with the aggradation and procession of the Red River Delta. The Red River Delta is being slowly drowned because of the slow (≈15 cm/100 yrs, see Penner 1974; Johnston 1946) increase in elevation of land at the lake's outlet. This process is significant over geologic time, but probably would not affect sediment distributions over 50 years time.

Lake Winnipeg elemental sediment supply rates (Table 30) were usually on the lower side of ranges of sedimentation rates for Lakes Erie and Ontario, usually greater than the sedimentation rates of Lakes Superior and Huron, and surprisingly comparable to rates for California coastal sedimentation basins. We found a small increase in Hg concentration in surface sediments (relative to older sediments) from the Red River Basin, the mouth of the Red River, and Traverse Bay near the mouth of the Winnipeg River (Table 24). Armstrong and Hamilton (1973) found high (5-35 µmoles Hg/g dry weight) concentrations of Hg in bottom and suspended sediments of Clay Lake and the Wabigoon River system (downstream from a chlor-alkali plant at Dryden, Ontario). These waters are tributary to the Winnipeg River and Lake Winnipeg, and are a likely source for the higher Hg concentrations observed in Traverse Bay sediments. Derksen (1973) found slightly elevated Hg concentrations in waters and fish from the Assiniboine and Souris Rivers, which

are tributary to the Red River and the Red River Basin. It is also known that municipal sewage is a source of Hg (van Loon 1974; F.A.J. Armstrong, personal communication), so it would appear that the City of Winnipeg and the Assiniboine River are also sources for the higher Hg values observed in surface samples of Red River Basin cores (Table 24). Industrial and municipal sources of Hg are also known on the Saskatchewan River (Wobeser et al. 1970, Uthe et al. 1973; Patterson and Nursall 1975), but we have no Hg data for the sediments of the north basin of Lake Winnipeg.

Hg concentrations began to increase in the top 14-22 cm of sediment cores from the Red  $\,$ River Basin (Table 24). If the sedimentation rate for this basin is  $\approx 1$  mm yr<sup>-1</sup> (see above), then this depth of increasing Hg concentration is associated with 100-200 year old sediments. Perhaps this represents sediment mixing of technologically derived Hg into pre-settlement age sediments. Wolery and Walters (1974) used sediment depth of Hg increase to calculate sedimentation rates for western Lake Erie, based upon a known history of chlor-alkali industrial activity beginning in 1939. Thomas (1972) and Kemp et al. (1974) studied the distribution of Hg in pollen dated Lake Ontario cores. They found that sedimentary Hg profiles were in general agreement with industrial history, which indicates that initial Hg additions to the lake began in 1900. Some interstitial water samples did not show great enrichments of major ions or depletion of SO<sub>4</sub>, relative to overlying lake water (Table 25), and for these stations, relatively complete mixing between surface sediments and lake water probably occurs. We plan to measure <sup>210</sup>Pb and <sup>137</sup>Cs on some of these cores, which will give us independent estimates of sediment mixing depth and sedimentation rates.

Brunskill (1973) gave a nutrient budget for Lake Winnipeg for 1969, and he reported the fraction of river supplied N and P which was retained in the lake (input - outflow) as 60% and 25%, respectively. If this mass of nutrient elements annually retained in the whole lake was sedimented into the Red River Basin (the 8 m contour in the south basin, Figs. 3 and 26) and the Saskatchewan, Cannibal, Mukutawa, and Berens Basins (the 16 m contour in the north basin, Figs. 4 and 25), the sediment supply rates would be 291  $\mu$ moles N m-2 yr^1 and 4.6  $\mu$ moles P m-2 yr^1. This rough estimation indicates that N and P sediment supply rates in the Black Bear Basin, Berens Basin, Cannibal Basin, Mukutawa Basin, and Saskatchewan Basin (see Figs. 25 and 26) must be much less than in the Red River Basin (Table 30).

Allan and Brunskill (1977) discussed the variation of trace element concentrations in Lake Winnipeg sediments, based on some of the data given here in Tables 15-17. The enrichment of V (relative to Fe and Ti) in south basin sediments was thought to be related to small amounts of hydrocarbons and humic organic material in Cretaceous and Agassiz sediments of the Red River Valley, according to these authors. Although concentrations of some trace elements (Zn, Cr, V) were rather high compared to other lake sediment data, the co-variation of most of these trace

elements with conservative elements (Fe, Ti, Al) (see Table 16) seems to indicate a relatively natural rate of supply of these elements. Linear correlation matrices of data in Tables 2, 10, 15, 27 showed that the abundance of Co was not correlated with any of the trace elements determined. but that Cu, Zn, V, As and P were positively correlated with mean particle size  $(\phi)$ . Lacking individual sedimentation rates for each core, it is not feasible to identify sites of accelerated trace element deposition from Table 23. In most cases (with exception of Hg and Pb), there is little difference between trace element concentrations at core tops and bottoms. This is in marked contrast to the results of many sediment core studies on the St. Lawrence Great Lakes (Kemp and Thomas 1976a, b; Kemp et al. 1976; Kemp et al. 1978; Kemp and Dell 1976; Lineback and Gross 1972; Frye and Shrimp 1973; Walters et al. 1974; and others referenced therein) which demonstrate authropogenic enrichments of many trace elements in the surface 0-10 cm of deep sediments.

# **ACKNOWLEDGMENTS**

We thank Capt. Kris Thorsteinsen and the crew of the Bradbury for their assistance and interest in our work. R. Sandilands, D. W. Kushnir, and J. F. Flannagan got very muddy and wet helping with shipboard operations. Dr. P. Sly (Canada Centre for Inland Waters) and Mr. A. Kirby (Department of Public Works, Selkirk) are thanked for support. Dr. R. J. Allan provided us with most of the trace element data. Forbes and Coleman (Young & Forbes Diving & Marine Co., Toronto) allowed us to calibrate our sediment sampling gear against their diver cores. F.A.J. Armstrong and M. P. Stainton, of the Freshwater Institute analytical laboratories, did some of the analytical determinations. C. Anderson, A. Prefontaine, and D. Laroque typed the manuscript. K. E. Marshall provided excellent support in the library, and S. T. Zettler, L. Davies and L. Taite did the drafting. S.E.M. Elliott guided our intuitions to statistically acceptable conclusions, and P. Campbell helped to find lost files of data. Dr. J.R.W. Vallentyne provided spiritual guidance. P. Sly, J. T. Teller, W. Last, R. Hecky, T. A. Jackson, and M. J. Capel reviewed the manuscript.

# REFERENCES

ALLAN, R. J., and G. J. BRUNSKILL. 1977. Relative atomic variation (RAV) of elements in lake sediments: Lake Winnipeg and other Canadian Lakes, p. 108-120. In H. L. Golterman (ed.) Interactions between sediments and freshwater. W. Junk B. V. Publishers, The Hague. 473 p.

The Hague. 473 p.

ANDERSON, J. M. 1976. An ignition method for determination of total phosphorus in lake sediments. Water Res. 10: 329-331

sediments. Water Res. 10: 329-331.

ARMSTRONG, F.A.J., and A. L. HAMILTON. 1973.

Pathways of mercury in a polluted northwestern Ontario lake, p. 131-155. In Trace metals and metal-organic interactions in natural waters. Ann Arbor Science Publishers, Ann Arbor, Michigan.

BACK, G. 1836. Narrative of the Arctic land expedition to the mouth of the Great Fish River . . . in 1833, 1834, and 1835. John Murray, London. 663 p.

BAILLIE, A. W. 1952. Ordovician geology of Lake Winnipeg and adjacent areas in Manitoba. Manit. Dep. Mines Nat. Resour., Mines Branch Publ. 51-6: 1-64.

BAJKOV, A. 1930. Biological conditions of Manitoban lakes. Contrib. Can. Biol. Fish. N.S. 5(12):

383-422.

BRULAND, K. W., K. BERTINE M. KOIDE, and E. D. GOLDBERG. 1974. History of metal pollution on southern California coastal zones. Environ. Sci. Technol. 8: 425-432.

BRULAND, K. W., M. KOIDE, C. BOWSER, L. J. MAHER, and E. D. GOLDBERG. 1975. Lead-210 and pollen geochronologies on Lake Superior

sediments. Quat. Res. (N.Y.) 6: 89-98. BRUNSKILL, G. J. 1973. Rates of supply of nitrogen and phosphorus to Lake Winnipeg, Manitoba, Canada. Int. Ver. theor. angew. Limnol. Verh. 18: 1755-1759.

BRUNSKILL, G. J., P. CAMPBELL, and S. E. M. ELLIOTT. 1979. Temperature, oxygen, conductance and dissolved major elements in Lake Winnipeg. Can. Fish. Mar. Serv. MS Rep. 1526: v + 127 p. BRUNSKILL, G. J., S. E. M. ELLIOTT, and P. CAMPBELL.

1980. Morphometry, hydrology, and watershed data pertinent to the limnology of Lake Winnipeg. Can. Fish. Mar. Serv. MS Rep. (in prep.).

BRUNSKILL, G. J., D. POVOLEDO, B. W. GRAHAM and M. P. STAINTON. 1971. Chemistry of surface sediments of sixteen lakes in the Experimental Lakes Area, northwestern Ontario. J. Fish.

Res. Board Can. 28: 277-294.
BRUNSKILL, G. J., D. W. SCHINDLER, S. K. HOLMGREN, H. KLING, P. CAMPBELL, F. A. J. ARMSTRONG, and M. P. STAINTON. 1980. Nutrients, chlorophyll, phytoplankton, and primary production in Lake Winnipeg. Can. Fish. Mar. Serv. MS Rep. (in prep.).

BYRNES, B. H., D. R. KEENEY, and D. A. GRAETZ. 1972. Release of ammonium-N from sediments to waters. Proc. Conf. Int. Assoc. Great

Lakes Res. 15: 249-259.

CARVER, R. E. 1971. Procedures in sedimentary petrology. Wiley-Interscience, Toronto. 653 p.

DAMIANI, V., and R. L. THOMAS. 1974. The surficial sediments of the Big Bay section of the Bay of Quinte, Lake Ontario. Can. J. Earth Sci. 11: 1562-1576.

DAVIES, J. F., B. B. BANNATYNE, G. S. BARRY, and H. R. McCABE. 1962. Geology and mineral resources of Manitoba. Manit. Dep. Mines Nat. Resour., Mines Branch. 190 p.

DEAN, W. E. 1974. Determination of carbonate and organic matter in calcareous sediments and sedimentary rocks by loss on ignition: comparison with other methods. J. Sediment. Petrol. 44: 242-248.

DENIS, L. G. 1916. Water-powers of Manitoba, Saskatchewan, and Alberta. Warwick Bro's. and Rutter, Ltd., Toronto. 334 p. DERKSEN, A. J. 1973. Mercury monitoring of

fishes and waters from the Assiniboine drainage system in Manitoba, 1970-71. Manit. Dep. Mines Resour. Environ. Manage. Res. Branch MS Rep. 73-32. 49 p.

DICKAS, A. B. 1970. Depositional environments of Lake Superior sands through grain size analyses. Proc. Conf. Int. Assoc. Great Lakes Res. 13: 227-232.

EDGINGTON, D. N., and J. A. ROBBINS. 1976. Patterns of deposition of natural and fallout radionuclides in the sediments of Lake Michigan, and their relation to limnological processes, p. 705-729. *In J. O. Nriagu (ed.) Environmental biogeochemistry. Vol. 2. Ann Arbor Science* Publishers, Ann Arbor, Michigan.

EHRLICH, W. A., H. M. RICE, and J. H. ELLIS. 1955. Influence of the composition of parent materials of soil formation in Manitoba. Can. J. Agric.

Sci. 35: 407-421.

EINARSSON, E., and A. B. LOWE. 1968. Seiches and set-up on Lake Winnipeg. Limnol. Oceanogr. 13: 257-271.

ELSON, J. A. 1961. Soils of Lake Agassiz, p. 51-79. In R. F. Legget (ed.) Soils in Canada. R. Soc. Can. Spec. Publ. 3.

1967. Geology of glacial Lake Agassiz, p. 37-95. In W. J. Mayer-Oakes (ed.) Life, land and water. University of Manitoba Press, Winnipeg.

FOLK, R. L., and W. C. WARD. 1957. Brazos River Bar: a study in the significance of grainsize parameters. J. Sediment. Petrol. 27: 3-26.

FRANKLIN, J. 1823. Narrative of a journey to the shores of the Polar Sea in the years 1819, 1820, 1821, and 1822. John Murray, London, Reprinted

by M. G. Hurtig, Edmonton, 1969. 768 p. FRYE, J. C., and N. F. SHRIMP. 1973. Major, minor, and trace elements in sediments of Late Pleistocene Lake saline compared to those in Lake Michigan. III. State Geol. Surv. Environ. eol. Notes 60: 1-14.

GARRELS, R. M., and C. L. CHRIST. 1965. Solutions, minerals, and equilibria. Harper and Row,

New York, 450 p.

GOLTERMAN, H. L. 1977. Sediments as a source of phosphate for algal growth, p. 286-293. H. L. Golterman (ed.) Interactions between sediments and freshwater. W. Junk B. V. Publishers, The Hague. 473 p.

HAMBLIN, P. F. 1976. Seiches, circulation and storm surges of an ice-free Lake Winnipeg. J. Fish. Res. Board Can. 33: 2377-2391.

HARTER, R. D. 1968. Adsorption of phosphorus by lake sediments. Soil Sci. Soc. Am. Proc. 32: 514-518.

HAUSER, B. W. 1973. Modifications to Carlo-Erba elemental analyzer for rapid determination of carbon and nitrogen in suspended matter of natural water. Fish. Res. Board Can. Tech. Rep. 412: iii + 11 p. HEALEY, F. P. 1975. Physiological indicators of

nutrient deficiency in algae. Can. Fish.

Mar. Serv. Tech. Rep. 585: 30 p. HOPWOOD, V. G. 1971. David Thompson. in western North America 1784-1812. Macmillan

of Canada, Toronto. 342 p.
JACKSON, M. L. 1956. Soil chemical analysis advanced course. Dep. Soil. Sci., Univ. of Wisconsin, Madison, Wisconsin. 894 p. JENNING, D. S., M. D. THOMAS, and W. GARDINER.

1922. A new method of mechanical analysis

of soil. Soil Sci. 14: 485-499. JOHNSTON, W. A. 1934. Surface deposits and ground water supply of the Winnipeg map-area, Manitoba. Geol. Surv. Can. Mem. 174.

1946. Glacial Lake Agassiz, with special reference to the mode of deformation of the beaches. Geol. Surv. Can. Bull. 7: 1-20.

KACHUGIN, E. G. 1966. The destructive action of waves on the water reservoir banks. In IASH Symposium Garda 1: 511-517

KEMP, A. L. W. 1971. Organic carbon and nitrogen in the surface sediments of Lakes Ontario, Erie and Huron. J. Sediment. Petrol. 41: 537-548.

KEMP. A. L. W., T. W. ANDERSON, R. L. THOMAS, and A. MUDROCHOVA. 1974. Sedimentation rates and recent sediment history of Lakes Ontario, Erie, and Huron. J. Sediment. Petrol. 44: 207-218.

KEMP, A. L. W., and C. I. DELL. 1976. A preliminary comparison of the composition of bluffs and sediments from Lakes Erie and Ontario. Can. J. Earth Sci. 13: 1070-1081.

KEMP, A. L. W., and A. MUDROCHOVA. 1972. Distribution and forms of nitrogen in a Lake Ontario sediment core. Limnol. Oceanogr. 17: 855-867.

KEMP, A. L. W., and R. L. THOMAS. 1976a. Cultural impact on the geochemistry of the sediments of Lakes Ontario, Erie, and Huron. Geosci. Can. 3: 191-207.

1976b. Impact of man's activities on the chemical composition

in the sediments of Lakes Ontario, Erie, and Huron. Water Air Soil Pollut. 5: 469-490.

KEMP, A. L. W., R. L. THOMAS, C. I. DELL, and J. M. JAQUET. 1976. Cultural impact on the geochemistry of sediments in Lake Erie. J. Fish.

Res. Board Can. 33: 440-462.

KEMP, A. L. W., J. D. H. WILLIAMS, R. L. THOMAS, and M. L. GREGORY. 1978. Impact of man's activities on the chemical composition of the sediments of Lakes Superior and Huron. Water Air Soil Pollut. 10: 381-402. KENNEY, B. C. 1979. Lake surface fluctuations

and the mass flow through the narrows of

Lake Winnipeg. J. Geophys. Res. 84: 1225-1235. KLOVAN, J. E. 1966. The use of factor analyses in determining depositional environments from grain size distributions. J. Sediment. Petrol. 36: 115-125.

KLUG, H. P., and L. E. ALEXANDER. 1954. X-ray diffraction procedures for pure crystalline and amorphous materials. John Wiley and Sons Inc., New York. 716 p.

KUSHNIR, D. W. 1971. Sediments in the south basin of Lake Winnipeg. Unpublished M.Sc. Thesis, Dep. Geology, Univ. Manitoba, Winnipeg, Man. 92 p.

LAST, W. M. 1974. Clay mineralogy and stratigraphy of offshore Lake Agassiz sediments in southern Manitoba. Unpublished M.Sc. Thesis, Dep. Geology, Univ. Manitoba, Winnipeg, Man. 183 p.

1979. Recent sediment of Lake Manitoba. Geol. Soc. Am. 13th Ann. Meeting (No. Central Sect.) 11: 233 (abstract).

LAST, W. M., and J. T. TELLER. 1979a. The Lake Manitoba stratigraphic record: a preliminary summary. Univ. Manit. Field Stn Rep. 13: 43-49.

1979b. Holocene and Late Pleistocene sedimentation history of Lake Manitoba, Canada. Geol. Soc. Am. 15th Ann. Meeting (No. Central Sect.) 11: 234 (abstract).

LEWIS, E. T., and G. J. RACZ. 1969. Phosphorus movement in some calcareous and non-calcareous

Manitoba soils. Can. J. Soil Sci. 49: 305-312. LINEBACK, J. A., and D. L. GROSS. 1972. Depositional patterns, facies, and trace element accumulation in the Waukegan member of the Late

Pleistocene Lake Michigan Formation in Southern Lake Michigan. Ill. State Geol. Surv. Environ. Geol. Notes 58: 1-25.

MACKENZIE, A. 1801. Voyages from Montreal on the River St. Lawrence, through the continent of North America, to the frozen and Pacific Oceans . . . T. Cadell, Tun., and W. Davies, London. 412 p. (A good modern reprint is edited by W. Kay Lamb, Macmillan of Canada, Toronto, 1970)

MAYER-OAKES, W. J. (ed.). 1967. Life, land, and water. University of Manitoba Press, Winnipeg.

414 р.

MOTHERSILL, J. S. 1975. Lake Chad: geochemistry and sedimentary aspects of a shallow polymictic Take. J. Sediment. Petrol. 45: 295-309.

MUNSELL SOIL COLOR CHARTS. 1954. Munsell Color

Company, Inc. Baltimore, Maryland. NEWBURY, R. W., K. G. BEATY, and G. K. McCULLOUGH. 1978. Initial shoreline erosion in a permafrost affected reservoir, Southern Indian Lake, Canada, p. 833-839. In Proc. Third Int. Conf. on Permafrost, July 1978, Edmonton, Alberta. V. I.

PATTERSON, C. G., and J. R. NURSALL. 1975. The effects of domestic and industrial effluents on a large turbulent river. Water Res. 9: 425-435.

PENNER, F. 1974. Lake Winnipeg, Churchill, and Nelson River study board, hydrologic, hydraulic, and geomorphologie studies. Vol. 1, Appendix 2, Part B, Annex 3. p. 105-109.

RACZ, G. J., and R. J. SOPER. 1967. Reaction products of orthophosphates in soils containing varying amounts of calcium and magnesium. Can.

J. Soil Sci. 47: 223-230. REEBURGH, W. S. 1967. An improved interstitial water sampler. Limnol. Oceanogr. 12: 163-230.

REPORT OF THE SASKATCHEWAN-NELSON BASIN BOARD. 1972. Water supply for the Saskatchewan-

Nelson Basin. Section E, 439-468. REX, R. W. 1969. X-ray mineralogy studies - Leg. 1, p. 354-367. In M. Ewing et al. Initial Reports of the Deep Sea Drilling Project. Vol. 1. U.S. Gov. Printing Office, Washington, D.C.

RINGROSE, S. M. 1975. A re-evaluation of Lake Agassiz shoreline data from north-central

Manitoba. Albertan Geogr. 11: 33-41. ROBINSON, G. W. 1922. A new method for the mechanical analysis of soil and other dispersions. J. Agric. Sci. 12: 306-311.

ROSS, A. 1856. The Red River settlement . . . Smith, Elder & Co., 65 Cornhill, London, Reprinted by Ross & Haines, Inc., Minneapolis,

Minnesota, 1957. 416 p. SCHINDLER, D. W. 1976. Biogeochemical evolution of phosphorus limitation in nutrient-enriched lakes of the Precambrian Shield, p. 647-664. In J. O. Nriagu (ed.) Environmental biogeochemistry. Vol. 2. Ann Arbor Science
Publishers, Ann Arbor, Michigan.
SHENG, Y. P., and W. LICK. 1979. The transport

and resuspension of sediments in a shallow lake. J. Geophys. Res. 84: 1809-1826.

SHEPARD, R. P. 1954. Nomenclature based on sandsilt-clay ratios. J. Sediment. Petrol. 24: 151-158.

SLY, P. G. 1977. Sedimentary environments in the Great Lakes, p. 76-82. In H. L. Golterman (ed.) Interactions between sediments and freshwater. W. Junk B. V. Publishers, The Hague. 473 p.

SLY, P. G., and R. L. THOMAS. 1974. Review of geological research as it relates to an

understanding of Great Lakes limnology. J. Fish. Res. Board Can. 31: 795-825.

SMITH, R. E., W. A. EHRLICH, and S. C. ZOLTAI. 1967. Soils of the Lac du Bonnet Area. Manitoba and Canada Department Agriculture, Soils Report No. 15: 1-118.

SOLOHUB, J. T. 1967. Grand Beach - a test of grain-size distribution statistics as indicators of depositional environments. Unpublished M.Sc. Thesis, Dep. Geology,

Univ. Manitoba, Winnipeg, Man. 118 p. SOLOHUB, J. T., and J. E. KLOVAN. 1970. Evaluation of grain-size parameters in lacustrine

environments. J. Sediment. Petrol. 40: 81-101. SPRY, I. M. (ed.) 1968. The papers of the Palliser Expedition in 1857-1860. The

Champlain Society, Toronto. 694 p. STAINTON, M. P. 1973. A syringe gas-stripping procedure for gas-chromatographic determination of dissolved inorganic and organic carbon in fresh water and carbonates in sediments. J. Fish. Res. Board Can. 30: 1441-1445.

STAINTON, M. P., M. J. CAPEL, and F. A. J. ARMSTRONG. 1974. The chemical analysis of fresh water. Fish. Res. Board Can. Misc. Spec. Publ. 25. 125 p.

STEVENSON, F. J., and A. P. S. DHARIWAL. 1959. Distribution of fixed ammonium in soils.

Soil Sci. Soc. Am. Proc. 23: 121-125. SUHR, N. H., and C. O. INGAMELLS. 1966. Solution techniques for analysis of silicates. Anal. Chem. 38: 730-734.

TAMPLIN, M. J. 1967. A brief summary of glacial Lake Agassiz studies, p. 27-36. In W. J. Mayer-Dakes (ed.) Life, land and water. University of Manitoba Press, Winnipeg.

TELLER, J. T. 1972. Aeolian deposits of clay sand. J. Sediment. Petrol. 42: 684-686. 1976. Lake Agassiz deposits in the main offshore basin of southern Manitoba.

Can. J. Earth Sci. 13: 27-43.

TELLER, J. T., B. B. BANNATYNE, P. LARGE, and S. RINGROSE. 1976. Quaternary sediment, bedrock, topography and geology of southern Manitoba. Manit. Mines Resour. Environ. Manag. Mineral Res. Div., Superficial Map Series 76-1 to 76-4.

TELLER, J. T., and M. M. FENTON. 1979. Late Wisconsinan glacial stratigraphy and history of southeastern Manitoba. Can. J. Earth

Sci. 16 (in press).

TENNANT, C. B., and R. W. BERGER. 1957. X-ray determination of dolomite-calcite ratio of

a carbonate rock. Am. Mineral. 42: 23-29. THOMAS, R. L. 1972. The distribution of mercury in the sediments of Lake Ontario. Can. J. Earth Sci. 9: 636-651.

THOMAS, R. L., A. L. W. KEMP, and C. F. M. LEWIS. 1972. Distribution, composition, and characteristics of the surficial sediments of Lake Ontario. J. Sediment. Petrol. 42: 66-84.

1973. The surficial sediments of Lake Huron. Can. J. Earth Sci. 10: 226-271.

THOMAS, R. L., and J. M. JAQUET. 1975. The surficial sediments of Lake Superior. Abst. Int. Sediment Congress, Nice (France). 11: 7 p.

THOMAS, R. L., J. M. JAQUET, A. L. W. KEMP, and D. F. M. LEWIS. 1976. Surficial sediments of Lake Erie. J. Fish. Res. Board Can. 33: 385-403.

UPCHURCH, S. B. 1970. Mixed population sediment in nearshore environments. Proc. Conf. Int. Assoc. Great Lakes Res. 13: 768-778.

UPHAM, W. 1890. Report on the exploration of the glacial Lake Agassiz in Manitoba. Ann. Rep. Geol. Nat. Hist. Surv. Can. (New Series): 4(E): 1-121.

UTHE, J. F., F. M. ATTON, and L. M. ROYER. 1973. Uptake of mercury by caged rainbow trout (Salmo gairdneri) in the South Saskatchewan

River. J. Fish. Res. Board Can. 30: 643-650. VAN LOON, J. C. 1974. Mercury input to the environment resulting from products and effluents from municipal sewage treatment plants. Environ. Pollut. 7: 141-147.

VELDMAN, W. M. 1969. Shoreline processes on Lake Winnipeg. Unpublished M.Sc. Thesis, Dep. Civil Engineering, Univ. Manitoba, Winnipeg, Man. 87 p.

VERNET, J. P., R. L. THOMAS, J. M. JAQUET, and R. FRIEDLI. 1972. Texture of the sediments of the Petit Lac (Western Lake Geneva). Eclogae Geol. Helv. 65: 591-610.

WALLACE, R. C., and J. E. MAYNARD. 1924. The clays of the Lake Agassiz Basin. Trans. R.

Soc. Can. Sect. IV, 18: 9-39.
WALTERS, L. J., JR., T. J. WOLERY, and R. D. MYSER. 1974. Occurrence of As, Cd, Co, Cr, Cu, Fe, Hg, Ni, Sb, and Zn in Lake Erie sediments. Proc. Conf. Int. Assoc. Great Lakes Res. 17: 219-234.

WARD, G. 1926. Seasonal variation in the composition of the Red River. Unpublished M.Sc. Thesis, Dep. Geology, Univ. Manitoba, Winnipeg,

Man. 24 p.

WATER SURVEY OF CANADA. 1969-1974. Sediment data for Canadian Rivers. Inland Waters Directorate,

Environment Canada, Ottawa.

WICKS, F. J. 1965. Differential thermal analysis of the sediments of the Lake Agassiz basin in metropolitan Winnipeg, Manitoba. Unpublished M.Sc. Thesis, Dep. Geology, Univ. Manitoba, Winnipeg, Man. 232 p.

WILLIAMS, J. D. H., J. M. JAQUET, and R. L. THOMAS. 1976. Forms of phosphorus in the surficial sediments of Lake Erie. J. Fish. Res. Board

Can. 33: 413-429.

WOBESER, G., N. O. NIELSEN, R. H. DUNLOP, and F. M. ATTON. 1970. Mercury contamination in tissues of fish from the Saskatchewan River. J. Fish. Res. Board Can. 27: 830-834.

WOLERY, T. J., and L. J. WALTERS, JR. 1974. Pollutant mercury and sedimentation in the western basin of Lake Erie. Proc. Conf. Int. Assoc. Great Lakes Res. 17: 235-249.

Winnipeg	Water Depth (m)	2 8 8 2 2 7 7 7 8 2 11 8 2 1 1 1 8 2 1 1 1 1 1 2 1 1 1 1
of water at Lake See also Fig. 2	Latitude (N)	88898888888888888888888888888888888888
, latitude and maximum depth stations, October 3-12, 1969	Longitude (W)	98 98 98 98 98 98 98 98 98 98 98 98 98 9
Table 1. Longitude, sediment st	Sediment Station	88889000000000000000000000000000000000

Particle size fractionation [sand (>50 µm), silt (50-2 µm), clay (<2 µm)] of Lake Winnipeg Surface sediment samples (homogenized, 0-20 cm sediment depth), expressed as % of mineral fraction <2 mm in size, collected 3-12 October 1969. Table 2.

Station	Puas #		
	™t 05<	# 5116 50-2 µm	λ Clay <2 μm
10	70.7	24.5	4.8
05	1.4 4.1	57.3	41.3
750	.u. ←	£ 7. 7.	4.05
030	2.0	51.5	2,45
04	19.6	36.6	43.00
048	8.0	51.3	47.9
- S	4.0	40.5	59.1
000	7.81	23.7	5.0°
80	200	40.00	70.0
10	0.9	28.1	71.0
12	76.5	14.9	8.6
± 1.	10.2 2.03	57.1	26.7
17	49.4	23.7	17.3
18	3.5	69.7	26.8
21	40.7	24.1	35.2
23H	0.0	43.4 R1 F	26.0
28	0.5	34.1	4.4
26	2.5	0.69	28.5
27	3.7	74.3	22.0
33.	, m	10.1 60.2	m 52
34	3.3	65,3	31.4
35	ກຸດ	67.6	31.9
41	1.0	30°C	24.8 40.5
47	0	62,9	37.1
48	æ. ;	26.0	72.2
-10-1 -10-1	10.3	42,1	47.6
21.	18.3	20.05	3.6
52	9.1	22.9	68.6
53	17.1	38.6	44.3
÷ 1:	1.5	49.0	47.9
3 45	1.4	. n	18.6 12.7
22	0.5	41.9	57.6
100 m	т. С	30.5	67.7
50		18./	80.8
90g	2.1	31.8	66.1
50C	6.0	45.6	53.5
50U	- a	41.1	57.8
77	9.0	7 100	2.5
	47.3	29.2	23.5
65	24.1	32.7	43.2
Red River in	3.6	44.9	51.5

surface	
(20-5 µm) of Lake Winnipeg surface	1969.+
F Lake 1	3-12, 1969.+
5 pm) o	ctober
Mineralogy of the medium silt fraction (20-5	0-20 cm homogenized) sediments collected O
t fracti	its coll
ium sil	sedime
the med	enized)
ogy of	m homog
Mineral	(0-20 cl
Table 4.	

Mineral Station	Quartz	Feldspar	Illite	Chlorite plus Kaolinite	Mantmarillonite
04	*5	s	ĸ	S	s
08	Σ	Σ	ស	Ŋ	ı
10	'n	ı	s	и	ហ
14	æ	Σ.	s	v.	'n
25	æ	æ	Σ	Σ	I
35	<b>_</b>	٠.	Σ	E	ı
52	Σ	Z	s	и	1
54	ب	_	Ŋ	S.	נעו
27	£	ı	s	ın	И
59	Ŋ	υ	Z	E	Σ
90	S	'n	ις	и.	E

\* L = 8-20 times background trace, a large  $\langle >50\% \rangle$  component of the crystalline mineral phase of the sediment.

M = 5-8 times background trace

(-) = none detected

S = 2-5 times background trace, a small (<10%) component of the crystalline mineral phase.

† The method of Jackson (1958) was used. Calcite and dolomite were identified separately by X-ray diffraction on an unacidified sample.

Table 3. Quantitative X-ray diffraction carbonate mineralogy (according to Tennant and Berger, 1957) of Lake Winnipeg 0-20 cm homogenized sediments collected October 3-12, 1969.

Station	% Dolomite	% Calcite	Ratio Dolomite: Calcite	Station	% Dolomite	% Calcite	Ratio Bolomite: Calcite
01	13	11	1.2	33	10	29	0.4
ŌS	12	7	1.7	34	8	< 1	> 8
03	8	7	1.1	35	1	< 1	> 1
03C	10	6	1.7	39	3	< 1	> 3
030	7	6	1.2	41	4	< 1	> 4
04	5	4	1.3	47	6	< 1	> ti
04B	8 3	< 1	> B	48	3	< 1	> 3
05		< 1	> 3	50	< 1	< i	-
06	4	< 1	> 4	50b	< 1	< 1	-
07	19	2	9.5	51	< 1	< 1	-
80	13	< 1	>13	52	6	< 1	> 6
10	1	< 1	> 1	53	< 1	< I	-
12	į.	< 1	> <u>l</u>	54	< l	< 1	-
14	<del>-</del>	< l	> 1	55	< 1	< 1	-
15	1	< 1	> <u>1</u>	57	< 1	< 1	-
17	1	< 1	> 1	59	< 1	< 1	-
18 21	1	< l	> 1	59B	< 1	< 1	-
21 23C	2	< 1	> 1	60	.3	< 1	> 3
23E	2	< 1 < 1	> 2	60B	15	< 1	>15
23E 23H	. 7	< 1 < 1	> 1 > 7	60C	′.	< 1	> /
25n 25		< 1	· <u>-</u>	60b	4 4	< 1	> 4
26	5 9	< 1	_	61 64	4 7	< 1	> 4
27	15	< 1 4	> 9 3.B	68	′	< 1	> /
27 31	20	< 1-	>20	69	< 1	< 1	-

Table 5. Mineralogy of the fine silt fraction (5-2  $\mu m$ ) of Lake Winnipeg surface (0-20 cm homogenized) sediments collected October 3-12, 1969.+

Mineral Station	Quartz	Feldspar	Illite	Chlorite plus Kaolinite	Montmorillonite
04	<u></u>	-	M	M	_
08	L	M	S	S	<del>-</del>
10	L	М	S	S	-
14	L	L	•••	S	-
25	М	S	S	S	~
35	М	M	М	М	S
52	L	М	\$	S	-
54	М	М	S	S	-
57	М	S	S	S	S
59	М	S	S	S	S
60	M	S	М	М	S

<sup>\*</sup> L = 8-20 times background trace

M = 5-8 times background trace

S = 2-5 times background trace

<sup>(-) =</sup> none detected

t The method of Jackson (1958) was used. Calcite and dolomite were identified separately by X-ray diffraction on an unacidified sample.

Table 6. Mineralogy of the total clay fraction (<2  $\mu m$ ) of Lake Winnipeg surface (0-20 homongenized) sediments collected October 3-12, 1969.

Mineral	Montmorillonite	Kaolinite plus Chlorite	Illite	Quartz	Feldspar
tation				М	
01	L	L	L S S	M	-
02	S	S S	Š	М	-
03	S		Ľ	M	-
03C	Ļ	1	Ī.	M	-
03D	Ľ.	S	S	М	<del>-</del> ,
04	Ş	Ĺ	L.	M	-
04B		· <u>Ē</u>	L	M	_
05	S	S	Ş	M	_
06		L	Ļ	M	_
07	ī	L	L	M M	М
08	Ĺ	M	r L	M	-
10 12	Š	· §	S S	M	
14	S	, <u> </u>	j I	M	_
15	L	L	S	M	M M
17	М	S M	М	M	М
18	M	S	Š	M	M
18 21	M	M	М	М	W
23C	М	ï	L,	S	S M
23E	- М	Š	М	M	M
23H	M M	М	M	M	M
25	M	М	M	M S	M
26	S	S	5	M M	М.
27 31	M	W	М	l l	Ĺ
33 31	-	Ŀ.	_ M	M	М
34	M	M	171 1	M	- S M
33 34 35	L	L M	м	M	М
39	Ņ		Ĺ	М	М
A1	L.	Ĺ	L.	M	-
47 48 50 50D	L f	<del>M</del>	L	M	M
48	<u>L</u> I	M	l.	М.	M 5
50	1	M	Ļ	М	J
500	ī	L	L	M	M
21	M	M	M 1	L	Ĺ
51 52 53 54 55 57	L	<u> </u>	M	M	M
54	М	M	M	S	S
55	М	M	i.	L	L
57	Ŀ.	M	M	S	М
59	M	1	L	L M	L
59B	L 88	М	М		M
60	M C	; · S	S	M	M M
60B	M S S M	5 S M	S S M	M M M	M M
60C	э М	_ M			
600			L.	L	L. M
61	M	M	M M	Ņ M	M
60 60B 60C 60D 61 64 68	L M M S	L M M S	M	M M S	L M M S
58 53	5	S	S	2	ų.

<sup>\*</sup> L = 8-20 times background trace

M = 5-8 times background trace

S = 2-5 times background trace

<sup>(-) =</sup> none detected

Mineralogy of suspended sediments from rivers draining into Lake Winnipeg, and a station from the South and North Basin of Lake Winnipeg. Table 9.

Location	Date	Suspended sediment mg/L	Montmor- illonite	Kaolinite	Illite	Illite Quartz	Feldspar	Dolomite	Calcite
Red River at Lockport (St. 00C)	11 May 1970	322	*	Σ	Σ	S	S	X	Σ
Red River at South Perimeter Bridge (St. 00A)	27 April 1970	555	. <b>.</b>	Σ	Σ	S	S		_
Assiniboine River at West Perimeter Bridge (St. 00B)	14 April 1969	High	Σ	Σ	Σ	Σ	S	Σ	Σ
Lake Winnipeg (St. 01)	3 Oct. 1969	85	v	ν	S	S	1	ı	J
Lake Winnipeg (St. 35)**	8 Oct. 1969	21	ı,	1	1	S	Σ	t .	Σ
			:				-		

L = 8-20 times background trace

M = 5-8 times background trace

S = 2-5 times background trace

(-) = none detected

\*\* Very small amount of sediment sample

Table 7. Mineralogy of coarse clay fraction (2-0.2 µm) of Lake Winnipeg surface (0-20 cm homogenized) sediments collected October 3-12, 1969.†

		Kaoline pus Kaolini pus M M M M	Montmorillonite  S  M  M  M  M  M  M  M  M  M  M  M  M
S	s	S	ь
v	t	=	2

\* L = 8-20 times background trace

† The method of Jackson (1958) was used. Calcite and dolomite were identified separately by X-ray diffraction on an unacidified sample.

Table 8. Mineralogy of medium clay fraction (0.2-0.8 µm) of Lake Winnipeg surface (0-20 cm homogenized) sediments collected October 3-12, 1969.+

Station	Quartz	Feldspar	Illite	Chlorite plus Kaolinite	Mantmorillonite
	•	1		•	E
	*5:	•	ı	ιs	Ŋ
	1	1	ı	ı	Σ
	Ŋ	•	s	и	Σ
	ហ	t	S	s	Σ
	1	ι	1	ı	Σ
	1	1	•	ı	Σ
	ŧ	1	1	ı	Σ
	1	•	s	s	

<sup>\*</sup> L = 8-20 times background trace

S = 2-5 times background trace

M = 5-8 times background trace

S = 2-5 times background trace

<sup>(-) =</sup> none detected

M = 5-8 times background trace

<sup>(-) =</sup> none detected

<sup>†</sup> The method of Jackson (1958) was used. Calcite and dolomite were identified separately by X-ray diffraction on an unacidified sample.

Table 10. Water content ( $H_2O$ ), expressed as % loss of weight of wet sample after drying to  $110^{\circ}C$ ; % loss on ignition (L.O.I.) is after ignition of dry sediment at  $900^{\circ}C$ ; total carbon ( $\Sigma C$ ), carbonate carbon ( $C_i$ ), organic carbon ( $C_0$ ), total nitrogen ( $\Sigma N$ ), total phosphorus ( $\Sigma P$ ) are expressed as  $\mu$ moles/g dry weight, with ratios on a molal basis for sediment samples (homogenized, 0-20 cm sediment depth) from Lake Winnipeg, October 3-12, 1969.

Station	% H <sub>2</sub> O	% L.O.I.	ΣC	c <sub>i</sub>	co	ΣN	ΣΡ	C <sub>O</sub> :ΣN	C <sub>o</sub> :ΣP	ΣΝ:ΣΡ
01 02	37.5	8.0	2000	1000	1000		20		50.0	-
02	60.0 59.2	12.5	3167 2333	1500	1667	64	25	25.9	67.0 39.5 47.0	2.6 2.7 2.5 3.7
03C	59.2 61.1	14.6 13.7	2333	1500 1500	83 <b>3</b> 1083	57 57	21 23	14.6 19.0	39.5	2.7
03D	71.0	13 7	2583 2167 1583	1167	1000	86	23	11.7	43.5	2.5
04	71.0	10.8 13.9 11.9	1583	833	750	79	23 25	9.6	30.0	3.2
04B	70.0	13.9	2083	1333 417 583 2250	750	79 107	21 23	7.0	30.0 35.7	5.1
05	73.5	11.9	1833	417	1416	114	23	12.4	61.7	5.0
06	68.4	19 //	1583 3583 3833	583	1000 1333	114 93 71	21 17	10.7	47.6	4.4 4.2
07	44.5	15.1 13.2 12.1 2.7 8.8 11.8	3583	2250 1417	1333 2416	71 50	1/	18.7	78.2	4.2
08 10	57.3 76.6	13.2	1500	83	1417	20	21	48.4 15.3	115.2 67.6	2.4 4.4
12	27.3	2 7	583	- 03	1417	93 < 7 79	21 21 22 24	15.5	07.0	4.4
14	60.9	8.8	2333	250	2083	79	24	- 26.5	- 86.7	3.3
15	60.9 66.7	11.8	2500	167	2333	114	23	20.4	101.3	5.0
17	56.8	5.2	583 2333 2500 1000	167 83	917	< 7	20		46.0	
18	76.9	10.8	1916	167 333	1749	157 93 28 136	24	11.1	72.9	- 6.5
21	52.0	4.6	1167 1333	333	834 1166	93	20	9.0	41.5	4.7
23C	77.4	8.5	1333	167 167	1166	28	26	40.8	45.0	1.0
23E 23H	78.9 77.8	8.2 11.8	1583 2666	1000	1416 1666	156 157	25 24	10.4 10.6	56.8 69.6	5.4 6.5
25n 25	77.0 76.7	11.0	2000	500	1500	157 150	24 24	10.0	62 5	6.3
26	76.7 71.7	11.5 11.7	2417	417	2000	150 93 157	24	21.6	62.5 83.3 78.9	3.9
27	59.8	21.2	4000	2500	1500	157	19	9.6	78.9	8.3
31	28.2	10.3	4083	3500	583	150	13	1.6	44.6	11.5
33	35.3 72.5 79.0 77.8	15.6 13.2	4083 3750 2333	3000	750 1083 1250	50 136	16	1.6 7.9	44.1 46.9 48.1 55.4 61.7 52.1 61.7 68.7 8.9 156.8	2.9 5.9 5.8
34 35	72.5	13.2	2333	1250	1083	136	23	7.9	46.9	5.9
35 39	/9.U	8.8 11.6	1500 1667	250 333	1250	150 179	26	8.3 7.5	48.1 EE 1	5.8 7.5
41	77.0 78.5	10.0	2000	533 583	1334 1417 1250	164	24 23	7.5 8.6	55.4 61.7	7 1
47	79.7	10.0 11.8	1917	667	1250	164	24	8.6 7.6	52.1	6.8
48	78.5 79.7 75.2 70.9 40.5 70.5	9.5 11.2	2000 1917 1833 1917 333	583 667 417 333 167 250 167 167	1416	150	24 23	9.4 10.1 3.4	61.7	6.8 6.5 6.2 2.6
50	70.9	11.2	1917	333	1584	143 50 286 171	23 19 25	10.1	68.7	5.2
50D	40.5	4.7 15.4	333	167	166 3916	50	19	3.4	8.9	2.6
51 52	70.5	15.4	4166 2417	250	3916	286	25	13.8	156.8	11.4 7.4
52 53	55.1 69.0	7.4 11.5	2417 1750	10/ 167	2250 1583	1/1	23	13.2 11.7	97.8	7.4 6.2
53 54	69.8	12.3	3250	167	3083	171	23	18.0	133 9	7.4
54 55	69.8 72.7	7.4 11.5 12.2 13.4	1750 3250 1750	167 167	3083 1583	136 171 93 136 121	23 22 23 23 25 25	17.0	71.8 133.9 69.1	4.0
57	79.5	9.5 9.4 11.2 9.2	1333 1250 1500 1500	167 83 167 333	1166	136	25	8.6	46.8	5.4
59	76.0	9.4	1250	83	1167	121	25	4.1	46.8 53.2	4.8 5.7
59B	74.2	11.2	1500	167	1333	143	25	9.3 9.1	53.2	5.7
<b>6</b> 0	74.8	9.2	1500	333	1167	129	27	9.1	43.3	4.8
60B	66.1 68.6	15.8	2583 2000	1583	1000 1333	100	19	18.3 13.3	96.3 60.5	5.3 4.5
60C 60D	71.7	13.5 13.1	1833	667 333	1500	100 129	22 24	11.7	62.5	5.4
61	71.6	9.6	2417	500	1917	214	23	9.0	83.4	9.3
54	74.0	10.4	2167	500	1667	164	24	10.1	69.6	6.8
68	65.9	9,3	2500	500	2000	136	19	14.7	105.3	7.2
69	65.8	12.2	2000	167	1833	121	20	15.0	91.5	6.1
ζ, 1σ	65.7 ±	11.1	2144 ±	732 ±	1412	117	23	13.3	65.5	5.4
	13.5	± 3.2	8 <del>5</del> 0	± 790	± 633	± 52	± 3	± 8.6	± 26.8	2.2

Table 11. Results of experiments to determine the labile fraction of Lake Winnipeg sedimentary phosphorus.+

Sediment Station	Extractant Solution	Sediment Concentration	Total P (EP) in sediment	P released	from sediment	P absorbed	d to sediment	Final p
	301661011	concent de lon	μmoles P/g	umoles P/g	% of EP	µmoles P/g	% of solution P	4.5
35	Distilled water	0.5 g/L	27 (±1.7)	1.10	4.1	-	_	6.8
	1M NH <sub>4</sub> C1	II	и	2.71	10.0	-	_	6.0
NORTH	0.01 N HCT	н	n	11.7	43.3	-	_	2,3
BASIN	D.O1 M NTA	n	D	17.1(±0.3)	63.3(±1.1)	-	_	2,5
	0.01 M NTA*	u	li .	10.9(±0.9)	40.5(±3.1)	-	-	7.0
	1.6 µM PO <sub>4</sub> -P	ti .	*1	-	-	0,26	8.0	6.8
	3.2 µM PO <sub>4</sub> -P	It	11	-	-	1.23	19.0	6,B
	6.4 µM PO4-P	п	ĮI.	-	-	2.06	16.0	6.9
	9.6 µM PO <sub>4</sub> -P	0	(I	-	-	2.97	15,3	6.8
60 <sup>°</sup>	Distilled water	0.5 g/L	23 (±0.1)	1.29	5.6	_		6.9
	1 M NH <sub>4</sub> C1			2.90	12.6	-	_	6.1
SOUTH	0.01 N HC1	п	u	10.2	44.3	_	-	2.2
BASIN	0.01 M NTA	n	u	18.4	80.0	_	_	2.5
	0.01 M NTA *	ь	,n	17.7	77.0	-	_	7.0
	1.6 µM PO <sub>4</sub> -P	и	H	-	-	0.52	16.0	6.9
	3.2 µM PO4-P	u	и	_		1.23	19.0	5.8
	6.4 µM PO <sub>4</sub> -P	' a	н	-	-	2.15(±0.23		6.8
	9.6 uM POu-P	п	Ħ	_	-	2.52	13.0	6.8

t Dry, homogenized surface sediment was added to the solutions listed below. The solution was stirred manually on occasion over a 68 hr period. Release or uptake of P by sediment was calculated from molybdate-reactive phosphorus measurements done on the filtered (Sartorius glass fibre) solution (minus experimental blank values). The PO $_{\rm t}$  solution was made from KH $_{\rm 2}$ PO $_{\rm t}$ . Numbers in parentheses are standard deviations for triplicate experiments.

\* The solution was buffered to pH 7.

Table 12. Calculated values, according to the method of Dean (1974), for organic carbon  $\{C_0\}$ , carbonate carbon (calcite + dolomite =  $C_1$ ), and total carbon (CC) from ashing curves of Lake Winnipeg sediment samples collected October 3-12, 1969, and their comparison to wet chemical carbon analysis (see Methods) as percentages of whole sediment dry weight at 100°C.

Station	Location	r% loss on ignition at 950°C	% wt loss of total wt loss 100-500°C	% wt loss of total wt loss 550~800°C	% wt loss of total wt loss 800-950°C	Col. ix2 % Co	Col. Ix3 % C as Dolomite	Col. 1x4 % C as Calcite	Col. 6+7 % C <sub>i</sub>	Co1. 5+8 % C
	Column	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	(9)
02	South Basin	12.50	44.17	53.33	2,50	1.5(2.0)*	1,81	0,09	1.9(1.8)	3.4(3.8)
04	South Basin	10.80	74.68	21.52	3.80	2.2(0,9)	0.68	0.12	0.8(1.0)	3.0(1.9)
12	Narrows	2.70	73.21	23,21	3.58	0.5	0.17	0.03	0,2(-)	0.7
14	Narrows	8,90	88.98	7.87	3.15	2.1(2.5)	0.21	0,09	0,3(0,3)	2.4(2.8)
26	North Basin	11.70	53.85	43.96	2, 19	1.7(2.4)	1.43	0.07	1.5(0.5)	3.2(2.9)
35	North Basin	9.80	81.04	13.79	5.17	1.9(1.5)	0.29	0.11	0.4(0.3)	2.3(1.8)
41	North Basin	10.00	76,92	19,23	3.85	2.1(1.7)	0.49	0.11	0.6(0.7)	2.7(2.4)
48	North Basin	9.50	82.47	14.43	3.10	2.1(1.7)	0.41	0.09	0.5(0.5)	2.6(2.2)
54	Narrows	12,20	93.01	3.50	3,49	3.1(3.7)	0.10	0.10	1.2(0.2)	3,3(3,9)
59	South Basin	9.40	88.37	8.53	3.10	2.2(1.6)	0.22	0.08	0.3(0.2)	2.5(1.8)

In column 5, 8 and 9, the values (% dry weight) in parenthesis are from wet chemical analysis given in Table 10. Organic carbon ( $C_0$ ) is from ( $EC-C_1$ ).

Elemental composition (in millimoles/g dry weight, 110°C) of selected surface sediment samples (homogenized, 0-20 cm sediment depth) from Lake Winnipeg. Collected October 3-12, 1969. Also given is % of total oxides that these elements represent. Table 13.

Station	Si	<u>a</u>	A1	Ca	Mg	Na	×	T. a	Mn	% Total
				1	}	1		1		
01	ي م	٠.		0	œ	ഹ	m.	m.	٥.	~
02	0.0	٠.		C.	$\sim$	ო.		7	0	,_i
90	8	9	•	œ.	٥.	~	Ξ.	æ	0	ď
07	9.	٠.		πż	<u></u>	ഹ	4.	4.	٥.	~
88	9.7	٠.	•		4.	₹.	r.	9	0.	ο.
12	3.7	٠.		4	ω.	4.	$\sim$	$\sim$	0	į.
14	ო.	٠.		4.	τċ	<u></u>	4.	9.	۰.	ij.
21	ω. Ο	٠.		ι.	9.	۲.	rt.	ഹ	٥.	~
230	0.	0.	•	u.	σ.	4.	۲.	0	٥.	۲.
23E	0	٠.		m.	Q.	4.	۲.	ο.	0.	Ġ.
26	0	٠.		Q.	₹.	4.	ø.	ω.	0	ά
33	~	0		9	۲.	5	9	9	0.	ъ.
35	0	0		സ	Q.	۲.	9	o.	o	o.
39	o. ⊘.	О.		φ.	0.	4.	۲.	$\infty$	٥.	~
41	സ	٥.		4.	o.	۷.	9.	α	o.	ς.
48	0	0		4.	Q.	ഹ	9.	o.	o.	, i
20D	9.2	0.		ന	ο.	<u></u>	ထ	ο.	٥.	o.
51	9	٥.	•	7.	9.	9.	9.	ω	0.	o,
52	0.0	0		ĸ.	0	r.	9	$\infty$	o.	
53	0.7	0		₹.	9	9	ന	۲.	٥.	o.
54	0.3	О.		₹.	~	ō.	സ	۲.	0.	o,
27	0.3	0.		က္	. 7	സ	Τ.	Q	0.	σ,
29	ლ. О	0		ო.	~	സ		Q.	0	ď
90	0.7	٥.		'n.	$\overline{\omega}$	ď	Ξ.	ο.	0.	ď
19	0.4	٥.		₹.	œ	٧.	-	φ	0	ď
64	10.46	0.02	3.56	0.46	0.50	0.88	0.58	0.54	0.01	95.64
69	u,	۰.		· .	0	₹.	4.	<b>ب</b>	o.	

Table 14. Concentrations of major elements in Lake Winnipeg surface sediment samples (0-20 cm, homogenized).

	•	mMo1	les/gram dry we	ight		
ion Number	Ca	Mg	Al	Fe	Mn	Ti
01	1.12	0.70	1.52	0.25	0.009	0.044
02	1.12	1.07	2.45	0.50	0.019	0.081
03	1.15	1.07	2,56	0.50	0.016	0.075
03C	1.02	1.03	2.22	0.43	0.014	0.069 0.073
03D	1.02	0.95	2.41	0.48	0.019	0.073
04	0.82	0.95	2.82	0.61	0.021	0.084
04B	0.62	0.99 0.78	2.71	0.57 0.65	0.019 0.020	0.088
05	0.42	0.78	2.93	0.57	0.016	0.088
06	.0.55	0.86	2.82 1.96	0.22	0.007	0.050
07	1.45	1.48	2.33	0.32	0.012	0.067
08	1.00	1,15	3.11	0.54	0.027	0.088
10	0.27 0.27	1.15 0.62 0.25	1.52	0.16	0.011	0.027
12	0.50	0.49	2.63	0.36	0.016	0.071
14 15	0.45	0.49	2.74	0.38	0.015	0.067
17	0.57	0.49	3.74	0.39	0.014	0.067
18	0.37	0.70	3.00	0.52	0.013	0.088
21	0.67	0,66	2.78	0.39	0.012	0.069
23C	0.35	0.78	3.22	0.63	0.021	0.096 0.098
23E	0.37	0.82	3.22	0.59	0.017	0.084
23H	0.70	1.07	2.85	0.54	0.020 0.021	0.086
25	0.62	0.99	3.04	0.56	0.017	0.081
26	0.92	1.23	2.82	0.50 0.36	0.016	0.071
27	0.70	1.85	2.37 1.59	0.18	0.016	0.031
31	2.32	2.14 1.28	2.11	0.18	0.011	0.065
33	2.64	1.11	2.59	0.45	0.011	0.079
34	0.87 0.37	0.78	3.15	0.61	0.015	0.094
35 39	0.42	0.78	2.93	0.52	0.025	0.084
41	0.47	0.86	3.00	0.54	0.024	0.084
47	0.55	0.91	2.89	0.52	0.012	0.081
48	0.37	0.70	2.71	0.47	0.013	0.077 0.088
50	0.45	0.66	3.15	0.52	0.012 0.015	0.090
50D	0.50	0.86	3.37	0.57	0.013	0.069
51	0.40	0.49	2.67 2.82	0.45 0.32	0.010	0.063
52	0.50	0.45	2.74	0.41	0.012	0.079
53	0.40	0.54 0.58	2.89	0.41	0.014	0.084
54 55	0.40 0.40	0.58	2.74	0.45	0.020	0.079
57	0.30	0.66	3.22	0.57	0.024	0.094
59	0.27	0.58	2.89	0.50	0.018	0.081
59B	0.30	0.66	3.08	0.56	0.030	0.092
60	0.35	0.66	2.96	0.54	0.023	0.088
60B	1.12	1.40	2.56	0.41	0.019	0.073 0.081
60C	0.70	0.91	2.74	0.45	0.017 0.022	0.090
60D	0.42	0.74	2.89	0.52 0.52	0.022	0.030
61	0.42	0.74	2.96 2.93	0.52	0.011	0.084
64	0.72	0.91 0.41	2.08	0.30	0.011	0.048
68 69	0.37 0.32	0.62	2.82	0.45	0.011	0.084
Average	$\bar{x} = 0.67$	0.86	2.73	0.46	0.016	0.077
Standard Deviation	± 0.48	± 0.35	± 0.45	± 0.12	± 0.005	± 0.015

<sup>+</sup> These data are from acid refluxing of the sediment, and analysis by radio frequency plasma emission spectroscopy. The data were provided to us by Dr. R. J. Allan.

Table 15. Concentrations of some trace elements in Lake Winnipeg surface sediment samples (0-20 cm, homogenized).+

						<del></del>	<del></del>					
Station					οMμ	les/gram d	ry weight o	of total	sediment			
Number	Сд	Pb	Ве	Co	Си	in	Sr	Cr	Zn	٧	As	p
01 02 03 03C 04 04B 05 06 07 08 10 12 145 17 18 21 23E 23E 24 27 31 33 41 47 48 50D 51 52 53 54 55 57 59 60C 60C 60C 60C 60C 60C 60C 60C 60C 60C	0.009 0.018	0.08 0.15 0.11 0.09 0.12 0.09 0.27 0.06 0.13 0.14 0.08 0.15 0.14 0.12 0.06 0.07 0.06 0.07 0.06 0.17 0.13 0.14 0.12 0.06 0.07 0.14 0.15 0.14 0.15 0.17 0.14 0.17 0.18 0.19	0.12 0.22 0.21 0.22 0.21 0.22 0.22 0.22	0.34 0.54 0.46 0.54 0.554 0.554 0.559 0.554 0.559 0.554 0.554 0.556 0.554 0.55	0.25 0.654 0.5558 0.	0.49 0.80 0.75 0.80 0.75 0.92 0.95 0.92 0.63 1.04 0.92 0.87 1.04 1.18 0.94 1.33 1.23 1.19 0.58 0.92 1.16 0.89 0.90 0.90 0.90 0.90 0.90 0.90 0.90	2.10 1.72 1.76 1.56 1.57 1.56 1.59 2.07 2.17 2.19 2.19 2.19 2.19 2.11 2.21 2.21 2.21	1.44 2.56 2.606 3.885 1.657 2.123.4775 2.123	1.88 3.20 3.15 3.14 3.30 44 3.31 2.38 3.30 2.46 3.30 2.31 2.31 3.30 2.31 3.30 2.31 3.31 2.31 3.31 3.31 3.31 3.31 3.31	1.73 3.83 3.98 3.91 4.55 4.28 4.50 2.45 2.45 2.45 2.45 3.61 3.47 3.20 4.20 3.31 2.20 3.55 4.20 3.22 3.55 3.08 3.22 3.55 3.08 4.50 3.40 4.53 3.40 4.53 3.40 4.53 3.40 4.53 3.40 4.53 4.50 3.55 4.10 4.50 3.60 5.50 4.10 4.50 6.60 6.60 6.60 6.60 6.60 6.60 6.60 6	0.040 0.040 0.040 0.040 0.040 0.040 0.040 0.040 0.027 0.027 0.027 0.027 0.040 0.027 0.040 0.027 0.040 0.027 0.040 0.027 0.040 0.027 0.040 0.027 0.040 0.027 0.040 0.027 0.040 0.053 0.040 0.053 0.040 0.053 0.067 0.053 0.067 0.053 0.040 0.053 0.040 0.053 0.040 0.040 0.053 0.040 0.053 0.040 0.040 0.053 0.040 0.053 0.040 0.040	14.2 20.7 19.4 18.4 19.0 21.0 20.3 19.0 20.3 19.0 20.3 17.8 21.6 21.6 21.6 21.6 21.6 21.6 21.6 21.6
						±0.20	2.04 ±0.32	2.65 ±0.64	2.90 ±0.50	3.32 ±0.90	0.040 ±0.013	

<sup>+</sup> These data are largely from determinations by radio frequency plasma emission spectroscopy and were provided to us by Dr. R. J. Allan.

Table 16. Relative atomic variation of trace elements in proportion to selected conservative elements (Fe, Ti, Al) in Lake Winnipeg sediments.\*

21 12 1 21 12 1 21 12 1 0 Ni C	Co 5 4 4 Co	Cr 38 38 Cr Cr 305	10 Co Cr V 11	68 68 70 270	Zn 35 28 Zn Zn 276
21 12 1 21 12 1 0 Ni C 21 82 6	5 4 4 CO CO 45 45	38 38 305 343	73 55 65 7 7 7 8 7 8 7 8 8 8	68 68 70 270	35 28 Zn 276 249
21 12 1 n Ni C 21 82 6 21 111 10	4 co Co Co 45	38 Cr Cr 305	55 fi x 10 V 559 484	F 570	28 Zn 276 249
n Ni C 21 82 6 21 111 10	ce Ele Co 42 45	Cr Cr 305	ri x 10 v 559 484	270	Zn 276 249
n Ni C 21 82 6 21 111 10	CO CO 42	Cr Cr 305 343	ri x 10 V 559 484	270	Zn 276 249
n Ni 21 82 21 111 1	l	Cr 305 343	V 559 484	ł	Zn 276 249
82		305	559 484		276
21 111		343	484		249
Trace	Elem	ent/A]	Trace Element/Al x 10 <sup>-4</sup>	.≄	
n Ni Cu	8	ڻ	>	_	Zn
South Basin 21 3 2	1	11	20	41	σı
North Basin 21 3 2		7	11	33	ស

n = number of sediment stations sampled and analyzed. The numbers given are the slopes (x10^4) of linear regressions of each trace element against the indicated conservative element, and can be thought of as the average ratio of trace: conservative element. All regressions are significant at the 95% level.

Table 17. Element concentration in suspended sediments from Red River drainage into the South Basin of Lake Winnipeg, May 2-9, 1975.†

22 266 266 27 1 13 266 28 28 29 29 29 29 29 29 29 29 29 29 29 29 29	of Winnipeg 975 864 174 430 18.5 2444 73 0.080 0.222 0.017 0.492 0.709 2.50 0.767	Station 008, Station 008, Assimboine River Red River upstream upstream from City of from City of from City of from City of from ipeg 675 850 1 864 823 174 130 430 448
	3.97	266 7 7 7 7 7 7 7 92 22 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
10°	3 05	.5 1 266 3 3 1 .222 .017 .492 .709 .50 .189 .767 .72

<sup>†</sup> These data were supplied by Dr. R. J. Allan, and the elements were determined by radio frequency plasma emission spectroscopy.

Table 19.	Annual rates of supply basin of Lake Winnipeg Red River suspended sed 1969-1973.	Annual rates of supply of major and trace elements to the south basin of Lake Winnipeg and sedimentation rates, derived from Red River suspended sediment chemistry and discharge data over 1969–1973.	to the south ived from le data over
Elements	Range of Annual Supply, Tonnes yr-1	Average sedimentation rate µMoles m <sup>-2</sup> yr <sup>-1</sup>	Sedimentation rate range (1969-1973)
N.	11-147	0.8	0.1-1.4
ð	8.2-109	8.2-109	0.07-1.0
ဝဌ	6.7-89	0.5	0.0-70.0
ŗ.	33-446	2.9	0.4-4.9
As	1,8-24	0.1	0.01-0.2
>	51-682	4.6	0.6-7.7
Be	0,46-6.2	0.2	0.03-0.4
Zn	47-631	3,3	0.4-5.6
3	0.46-6.2	0.02	0.002-0.03
Pb	2.6-34	0.06	0.007-0.1
72	38-507	2	0.3-3.3
短	260-3460	21	2.7-36
<b>5</b> _	148-1980	22	3-37
Τį	900-1200	86	11-144
œ.	5900-79,000	480	61-810
A1	19,400-258,900	3280	413-5,526
5;	64,700-866,260	10,530	1,326-17,760
S.	8,210-108,900	940	118-1,580
Mg	6,850-91,640	1,290	162-2,170
Na	2,350-31,520	470	60-790
<b>×</b> 4	4,910-65,660	570	72-970
Total Dry (110°C) Sediment		1170 g m <sup>-2</sup> yr <sup>-1</sup>	150-1970 g m <sup>-2</sup> yr <sup>-1</sup>

Table 18. Suspended sediment chemistry for Lake Winnipeg and Red River, by sodium metaborate fusion and atomic absorption spectrophotometric determination. Si and P were determined by coloremetric methods of Stainton et al. 1974.

Location	Date					mMo l e	es/g dry	weight		
Description	(1969)	51	A1	Ca	Mg	Na	K	Fe	Mn	þ
Composite of all Lake Yinnipeg stations sampled	Sept. 2-10 Oct. 3-13 Oct. 27-31	10.18 10.47 10.15	1.31 1.37 1.37	0.459 0.459 0.459	0.156 0.156 0.156	2.55 2.64 2.55	0.563 0.573 0.567	0.090 0.010 0.090	0.005 0.004 0.005	0.034
ake Winnipeg Station 701 Mouth of Red River	Oct. 3	2,57	0.437	4.84	0.424	0,339	0.123	0.095	0.009	0.032
ake Winnipeg Station 735 Center of North Basin	Oct. 8	4.64	1.23	0.961	0.395	0.631	0.315	0.251	0.010	0,068
Red River south of Jinnipeg (Station OOA)	Apr. 13 Apr. 27	6.50 9.31	1.99 3.22	1.44 0.614	1.73 0.913	1.05 0.244	0.425 0.517	0.440 0.68B	0.033 0.010	0.023
Red River north of Winnipeg (Station OOC)	Apr. 21 June 8	9.04 8.94	3.00 2.59	0.671 0.921	0,995 1,25	0.365 0.439	0.509 0.473	0.609 0.591	0.015 0.022	0.026

Table 20. Description of Lake Winnipeg sediment cores taken with a weighted Benthos corer on 3-12 October 1969.

Core No.	Location Long./Lat.	Date Taken	Date Extruded	Depth in Sediment	Munsell Color	Texture	% H <sub>2</sub> O of Dry Wt.	Remarks
04	South Basin 98°48'00" W 50°28'30" N	3 Oct 1969	11 Jan 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105 110-115	5Y3/1 Bluish- black	Fine plastic clay  Conchoidally fractured clay	75.4 70.0 64.8 57.5 57.0 59.8 57.0 55.9 51.9 45.0	Water above core. A few cm of flocculent orange sediment at surface. Sewage and hydrocarbon smell.
				120-125 130-135	5Y2/2	Granular clay	42.7 42.9	
08	Traverse Bay 96 <sup>0</sup> 28'30" 50 <sup>0</sup> 44'45"	4 Oct 1969	22 Jan 1971	0-5 cm 10-15 20-25 30-35 40-45	5Y3/1 Bluish black 5Y3/1	Fine plastic clay Conchoidally	60.9 57.8 50.5 48.3 48.0 43.1	Water above core.
				50-55 60-65 70-75 80-85 90-95	5Y3/1	fractured clay Sandy silt	40.6 35.5 32.9 30.6	· .
10	South Basin	4 Oct 1969	8 June 1971	0-5 cm 10-15 20-25	5Y2/1	Fine plastic	70.7 65.9 65.3 66.2	No water above core but surface sediment was wet
	96 <sup>0</sup> 33'30" 50 <sup>0</sup> 55'30"			30-35 40-45 50-55 60-65 70-75	5Y2/2	Conchoidally fractured	68.3 63.7 49.4 49.0	
				80-85	<u>5y2/2</u>	clay Granular clay	48.2	
14	Narrows 96 <sup>0</sup> 38'30" 51 <sup>0</sup> 26'30"	5 Oct 1969	13 May 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95	5Y2/2 5Y2/1	Fine plastic clay  Conchoidally fractured	64.0 65.2 64.2 65.7 63.7 61.8 46.1 58.1 58.4 45.4	Water above core.
				110-115 120-125	Black	clay Granular clay	- 48.7 42.9	
25	North Basin 98 <sup>0</sup> 53'30" 53 <sup>0</sup> 12'45"	6 Oct 196 <b>9</b>	1 Feb 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75	2.5Y3/1 Bluish grey	Fine plastic clay Slightly conchoidally fractured	69.3 55.4 70.4 70.0 69.4 68.7 69.7	No water above core. Dry surface Bluish blebs in surface sediment.
				80-85		clay Conchoidally fractured	57.3	
				90-95 100-105 110-115	5Y3/1 Very dar grey	clay rk Granular clay	- 48.3 48.4 45.9	

Table 20. Cont'd.

Core No.	Location Long./Lat.	Date Taken	Date Extruded	Depth in Sediment	Munsell Color	Texture	% H <sub>2</sub> O of Dry Wt.	Remarks
35-1	North Basin 98 <sup>0</sup> 34'45" 53 <sup>0</sup> 39'15"	8 Oct 1969	20 May 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85	5Y4/1 Bluish black 	Fine plastic clay  Conchoidally fractured clay Granular clay	72.2 72.3 74.3 73.8 73.1 72.4 71.1 -	No water above sediment, oxidation to 22 cm depth
35-2	North Basin 98 <sup>0</sup> 34'45" 53 <sup>0</sup> 30'15"	8 Oct 1969	7 Jan 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105	Brown (oxidized 5Y4/1 Bluish black	Conchoidally fractured clay	70.2 69.4 72.2 72.4 73.1 70.0 68.9 56.4 48.0 48.5	No water over core. Some oxidation to 30 cm. Gas trapped in sediment, $\rm H_2S$ smell.
39	North Basin 98 <sup>0</sup> 11'30" 53 <sup>0</sup> 02'30"	6 Oct 1969	4 June 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55	5Y2/2 Black	clay Fine plastic clay	70.1 71.4 73.4 72.5 72.0 71.7	
				70-75 80-85 90-95 100-105 110-115	572/1 (Black)	Conchoidally fractured clay Granular clay	68.7 70.1 72.5 70.8 52.5	
52	Narrows (Pigeon Bay) 97 <sup>0</sup> 08'00" 52 <sup>0</sup> 17'30"	5 Oct 1969	31 May 1971	0-5 cm 10-15 20-25 30-35 40-45	5Y3/1 Very dark grey	Silty sand Granular clay	54.1 49.1 26.3 26.4 37.3	Water above core.
54	Narrows 97°03'00" 52°00'00"	5 Oct 1969	27 May 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105 110-115	5Y4/1 Dark grey 5Y2/1 Black	Fine plastic clay  Conchoidally fractured clay Granular clay	72.1 70.9 69.1 66.7 65.2 63.8 63.2 59.9 57.7 40.8 43.7 52.5	Water above core.

Table 20. Cont'd.

Core No.	Location Long./Lat.	Date Taken	Date Extruded	Depth in Sediment	Munsell Color	Texture	% H <sub>2</sub> O of Dry Wt.	Remarks
57	South Basin 96°47'00" 50°51'30"	4 Oct 1969	3 June 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85	Marbled 5Y3/1 and 5Y2/1 Very dark grey and black	Fine plastic clay	75.6 75.3 73.0 71.7 71.3 70.2 69.4 69.4	Water above core.  Noticeable oxidation.
				90-95 100-105	5Y2/1 Black	Conchoidally fractured clay Granular clay	50.7 46.8	
59	South Basin 96°44'45" 50°42'00"	4 Oct 1969	2 May 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95	5Y3/2 	Fine plastic clay  Conchoidally fractured clay Granular clay		No water above core, Oxidized to 60 cm.
60	South Basin 96 <sup>0</sup> 47'15" 50 <sup>0</sup> 34'30"	3 Oct 1969	20 May 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85	5Y3/2	Fine plastic clay	67.1 - 54.2 - 50.7 - 44.1	Very hard to extrude. Compressed during extrusion
61	South Basin 96°48'15" 50°31'00"	3 Oct 1969	13 May 1971	0-5 cm 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105	5Y3/2 Dark olive grey 5Y2/2 Black	Fine plastic clay  Conchoidally fractured clay Granular clay	75.3 70.9 68.1 67.8 67.2 67.1 69.0 60.9 49.7 46.5 37.0	Water above core.

Table 21. Percent loss on ignition at  $900^{\circ}\text{C}$  (L.O.I.), µmoles/gm dry weight ( $110^{\circ}\text{C}$ ) of total carbon ( $\Sigma$ C), carbonate carbon ( $\Sigma$ C), organic carbon ( $\Sigma$ C), total nitrogen ( $\Sigma$ N), total phosphorus ( $\Sigma$ P), and molal ratios of selected Lake Winnipeg sediment cores taken October 3-12, 1969.

Core	Depth	L.O.I.	ΣC	c <sub>i</sub>	c <sub>a</sub>	ΣΝ	ΣΡ	C <sub>o</sub> :ΣN	C <sub>O</sub> :ΣP	ΣΝ: ΣΡ
04	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105 110-115 120-125 130-135	10.09 9.82 9.70 10.00 9.33 9.09 8.94 9.18 9.80 8.88 9.33 9.74 10.70 10.43	2083 1583 1500 1667 1867 1833 1417 2000 1750 2000 1583 1500 1917 1667	475 491 525 809 700 750 650 683 733 592 641 600 592 767	1608 1092 975 858 967 1083 767 1317 1017 1408 942 900 1325 900	136 93 78 93 71 78 64 64 29 100 64 86 71	22 20 20 18 20 19 22 22 20 18 23 19 21 20	11.8 11.7 12.5 9.2 13.6 13.9 12.0 20.6 35.1 14.1 14.7 10.5 15.4 12.7	73.1 54.6 48.8 47.7 48.4 57.0 34.9 59.9 50.9 78.2 41.0 47.4 63.1 45.0	6.2 4.7 3.9 5.2 3.6 4.1 2.9 2.5 5.6 2.8 4.5 4.1 3.6
08	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95	12.70 11.88 12.69 11.87 12.54 13.99 12.97 13.23 13.04 12.80	2792 2775 2783 2800 2525 3150 2850 2967 2725 2800	1542 1517 1283 1717 1767 1900 2092 2209 1967	1250 1258 1500 1083 758 1250 758 758 758	86 64 57 57 57 71 57 64 57 36	13 14 14 13 13 12 12 13 13	14.5 19.7 26.3 19.0 13.3 17.6 13.3 11.8 13.3	96.2 89.9 107.1 83.3 58.3 104.2 63.2 58.3 58.3	6.6 4.6 4.1 4.4 5.9 4.8 4.9 4.4 2.6
10	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85	20.17 19.71 10.26 12.48 10.03 10.34 9.96 11.97 12.03								
14	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105 110-115 120-125	7.41 8.81 10.18 8.69 8.59 8.49 8.99 8.06 9.52 8.62 6.21 8.39 8.30	1833 2083 2167 1583 1583 1500 1333 1667 1333 13417 1333	41 83 109 100 100 100 83 83 109 116 33 92	1792 2000 2058 1483 1483 1400 1250 1250 1558 1217 1300 1325 1242	100 129 121 100 93 86 86 107 100 100	20 22 22 19 22 21 24 23 22 22 23 22	17.9 15.5 17.0 14.8 15.9 20.6 14.5 14.5 14.6 12.2 13.0 13.3	89.6 90.9 93.5 78.1 67.4 63.6 59.5 52.1 67.7 55.3 59.1 57.6 56.5	5.0 5.5 5.3 4.2 3.9 4.1 4.5 4.5 4.5
25	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105 110-115	11.22 10.02 10.71 10.78 9.94 10.46 9.13 9.97 9.67 10.28 9.23 10.23	1583 1500 1667 1833 1500 1833 1417 1750 1500 1583 1667 1750	450 450 642 675 983 533 392 542 450 591 592 617	1133 1050 1025 1158 517 1300 1025 1208 1050 992 1075 1133	100 93 79 93 79 100 121 100 129 79 71	25 25 25 29 32 27 32 24 25 25 25	11.3 11.3 13.0 12.5 6.5 13.0 8.5 12.1 8.1 12.6 15.1	45.3 42.0 41.0 39.9 16.2 48.1 32.0 50.3 42.0 39.7 43.0 45.3	4.0 3.7 3.2 3.5 3.7 3.8 4.2 5.2 3.2 2.8 3.2

Table 21. Cont'd.

Core	Depth	L.O.I.	ΣC	C <sub>i</sub>	C <sub>o</sub>	ΣΝ	ΣΡ	C <sub>0</sub> :ΣN	C <sub>o</sub> :ΣP	ΣΝ: ΣΡ	
35-1	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95	8.27 8.08 7.49 7.79 7.70 6.25 7.64 -7.73									
35-2	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105	12.36 9.95 8.13 7.75 7.31 7.62 7.50 7.34 7.44 7.83 7.77	1833 1333 1417 1250 1000 1167 1167 6000 1083 1167 1000	41 66 59 50 42 84 25 42 100 117 75	1792 1267 1358 1200 958 1083 1042 958 983 1050 925	164 121 136 114 114 110 107 121 121	22 19 17 17 21 14 16 19 17 20 23	10.9 10.5 10.0 10.5 8.4 9.5 10.8 9.0 8.1 8.7 8.1	81.5 66.7 79.9 70.6 45.6 77.4 65.1 50.4 57.8 52.5 40.2	7.5 6.4 8.0 6.7 5.4 8.1 6.3 5.6 7.1 6.1	
39	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105 110-115	12.39 11.27 9.58 10.28 10.32 10.64 9.21 10.28 9.58 9.97 9.91 8.60 8.83									
52	0-5 10-15 20-25 30-35 40-45	48.00 10.82 5.22 5.39 9.87									
54	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105 110-115	11.52 10.24 20.19 10.40 8.97 8.00 17.31 6.93 8.34 6.45 6.51 6.50	2758 2067 1900 2433 2008 1508 1342 942 1600 1158 950 1067	75 50 58 33 41 25 25 25 25 25 25 34	2683 2017 1842 2400 1967 1483 1317 917 1575 1133 908 1033	164 136 114 129 129 79 121 129 114 79 93	22 22 26 20 28 24 25 24 22 29	16.4 14.8 16.2 18.6 15.2 18.8 10.9 7.1 13.8 14.3 9.8 14.5	122.0 91.7 83.7 92.3 98.4 53.0 54.9 36.7 65.6 51.5 31.3	7.5 6.2 5.0 6.5 2.8 5.0 5.2 4.8 3.6 3.7	
57	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105	11.77 12.17 11.09 10.25 10.85 13.13 8.71 9.74 9.38 9.72 10.61									

Table 21. Cont'd.

Core	Depth	L.0.I.	ΣC	c <sub>i</sub>	Co	ΣΝ	ΣΡ	Co:EN	C <sub>o</sub> :ΣP	ΣΝ:ΣΡ	
59	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95	9.57 8.52 9.83 8.20 7.95 7.79 7.74 7.70 7.65 7.68	917 750 750 708 667 708 667 667 750 917	100 83 83 116 92 109 67 100 75	817 667 667 592 575 617 600 567 675 842	79 71 71 64 64 71 64 71 57	23 22 21 22 21 22 21 22 20 22 23	10.3 9.4 9.4 9.3 9.0 8.7 9.4 8.0 11.8 14.8	35.5 29.0 30.3 28.2 26.1 29.4 27.3 28.4 30.7 36.6	3.4 3.1 3.2 3.1 2.9 3.4 2.9 3.6 2.6 2.5	
60	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85	8.99 - 8.63 - 8.96 - 8.71	1000 - 1083 - 1083 - 1000			100 - - 79 - - 71 - 71					
61	0-5 10-15 20-25 30-35 40-45 50-55 60-65 70-75 80-85 90-95 100-105	11.26 9.38 9.66 10.61 9.70 10.29 9.37 9.58 9.52 9.49 9.26	1667 1250 1250 1166 1166 1083 1083 833 1000 1083 1083			100 79 71 71 71 64 64 57 64 71					

Table 22. Major element chemistry of Lake Winnipeg sediment cores, taken 3-12 October 1969. These data were provided to us by Dr. R. J. Allan. Total element concentrations were determined by radio frequency plasma emission spectroscopy.

Station and Location	Depth in Core (cm)	Ca	Mg	Na	Al mMoles/	Fe 'gram	Ti	Mn	Р
04 South Basin, Near Red River	0-5 10-15 20-25 30-35 50-55 70-75	0.55 0.57 0.57 0.75 0.70 0.70	0.78 0.82 0.82 0.86 0.90 0.82	0.087 0.130 0.130 0.130 0.174 0.130	2.71 2.78 2.63 2.52 2.52 2.52	0.45 0.47 0.45 0.41 0.39 0.41	0.079 0.081 0.081 0.084 0.084 0.084	0.015 0.016 0.018 0.019 0.022 0.026	0.019 0.022 0.017 0.017 0.018 0.017
08 Traverse Bay, Near Winnipeg River	0-5 10-15 20-25 30-35 50-55 70-75	1.07 1.00 1.27 1.20 1.37 1.50	1.19 1.11 1.32 1.23 1.40 1.52	0.304 0.304 0.348 0.348 0.304 0.348	2.22 2.19 2.03 2.08 1.89 1.96	0.32 0.30 0.27 0.27 0.25 0.25	0.071 0.067 0.059 0.059 0.056 0.059	0.011 0.011 0.010 0.009 0.009 0.007	0.016 0.016 0.015 0.015 0.014 0.015
10 South Basin, North End	0-5 10-15 20-25 30-35 50-55 70-75	0.32 0.32 0.32 0.32 0.32 0.35	0.66 0.70 0.66 0.66 0.70 0.66	0.174 0.217 0.217 0.217 0.217 0.217	3.19 3.30 3.15 3.19 3.30 3.11	0.57 0.61 0.57 0.56 0.59 0.54	0.094 0.102 0.100 0.100 0.107 0.098	0.023 0.029 0.026 0.025 0.025 0.021	0.020 0.021 0.019 0.019 0.019 0.019
14 Narrows South End	0-5 10-15 20-25 30-35 50-55 70-75	0.45 0.45 0.42 0.45 0.42 0.42	0.45 0.45 0.49 0.53 0.53 0.49	0.565 0.565 0.478 0.565 0.522 0.565	2.67 2.67 2.74 2.93 2.85 2.93	0.38 0.38 0.41 0.41 0.43 0.39	0.075 0.077 0.084 0.086 0.088 0.083	0.014 0.016 0.018 0.016 0.016 0.015	0.019 0.019 0.019 0.019 0.018 0.019
25 North Basin Near Saskatchewan River mouth	0-5 10-15 20-25 30-35 50-55 70-75	0.62 0.62 0.78 0.80 0.72 0.75	0.99 1.03 1.07 1.11 1.07	0.261 0.304 0.261 0.304 0.304 0.304	2.93 2.97 2.82 3.00 3.04 3.19	0.56 0.56 0.56 0.57 0.59 0.61	0.086 0.088 0.086 0.094 0.096 0.102	0.013 0.018 0.023 0.027 0.027 0.022	0.019 0.020 0.020 0.023 0.022 0.021
35 North Basin	0-5 10-15 20-25 30-35 50-55 70-75	0.37 0.37 0.27 0.40 0.37	0.70 0.78 0.58 0.86 0.86 0.80	0.348 0.348 0.217 0.348 0.391 0.348	2.93 3.15 2.26 3.34 3.37 3.34	0.57 0.63 0.59 0.63 0.63 0.64	0.090 0.098 0.073 0.107 0.104 0.111	0.025 0.020 0.012 0.016 0.017 0.018	0.024 0.023 0.017 0.022 0.022 0.021
39 North Basin	0-5 10-15 20-25 30-35 50-55 70-75	0.47 0.42 0.37 0.42 0.42	0.82 0.82 0.82 0.86 0.86 0.82	0.348 0.348 0.304 0.304 0.348 0.304	3.19 3.22 3.15 3.22 3.34 3.04	0.56 0.57 0.57 0.61 0.59 1.54	0.093 0.098 0.093 0.098 0.102 0.090	0.011 0.014 0.015 0.015 0.015 0.014	0.020 0.021 0.020 0.020 0.020 0.020
54 Narrows North End	0-5 10-15 20-25 30-35 50-55 70-75	0.40 0.40 0.37 0.42 0.42 0.40	0.58 0.66 0.66 0.62 0.62 0.58	0.435 0.391 0.391 0.391 0.435 0.565	2.74 2.78 2.89 3.19 3.19 3.08	0.43 0.50 0.52 0.50 0.50 0.45	0.086 0.094 0.102 0.100 0.096 0.093	0.012 0.013 0.012 0.012 0.011 0.011	0.018 0.019 0.019 0.019 0.019 0.020
57 South Basin	0-5 10-15 20-25 30-35 50-55 70-75	0.30 0.30 0.30 0.30 0.30 0.30	0.66 0.66 0.70 0.66 0.66	0.217 0.174 0.174 0.174 0.174 0.174	3.26 3.26 3.30 3.30 3.30 3.41	0.41 0.36 0.57 0.57 0.56 0.59	0.098 0.094 0.098 0.098 0.102 0.100	0.025 0.023 0.025 0.026 0.025 0.025	0.019 0.020 0.020 0.019 0.019 0.020

Table 22. Cont'd.

Station and Location	Depth in Core (cm)	Ca	Мg	Na	Al mMoles	Fe /gram	Ti	Mn	Р
59 outh Basin	0-5 10-15 20-25 30-35 50-55	0.32 0.30 0.32 0.35 0.35	0.78 0.74 0.78 0.74 0.82	0.174 0.217 0.217 0.217 0.217	3.30 3.11 3.19 3.00 3.41	0.63 0.61 0.63 0.57 0.66	0.100 0.094 0.100 0.096 0.106	0.014 0.021 0.018 0.012 0.019	0.024 0.024 0.023 0.022 0.022
61	70-75 0-5	0.30	0.74 0.82	0.174	3.11	0.61	0.096	0.020	0.022
outh Basin	10-15 20-25 30-35 50-55 70-75	0.40 0.47 0.40 0.37 0.37	0.82 0.86 0.82 0.82 0.82	0.174 0.130 0.174 0.174 0.174	2.89 2.85 2.74 2.71 2.67	0.52 0.54 0.50 0.50 0.47	0.088 0.094 0.084 0.084 0.077	0.021 0.022 0.023 0.022 0.022	0.022 0.021 0.021 0.020 0.021

Table 23. Trace element chemistry of Lake Winnipeg sediment cores, taken 3-12 October 1969. These data were provided to us by Dr. R. J. Allan, and the total element concentrations were measured by radio frequency plasma emission spectroscopy.

		_								_			
Station and Location	Depth in Core (cm)	Cd	Pb	Ве	Со	Cu	Ni µMoles	Sr /gram	Cr	Zn	٧	As	Мо
04 South Basin Near Red River	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.018 0.018 0.018 0.018 0.018	0.17 0.12 0.05 0.08 0.12 0.08	0.27 0.26 0.23 0.27 0.26 0.24	0.49 0.49 0.49 0.49 0.49 0.49	0.55 0.50 0.46 0.46 0.46	1.06 0.83 0.85 0.78 0.82 0.83	1.50 1.52 1.55 1.62 1.70 1.60	3.29 2.85 2.87 2.48 2.67 2.73	3.66 3.30 3.11 3.27 3.01 2.89	4.69 4.44 4.34 3.97 4.12 4.38	- - - -	0.052 - - - - -
08 Traverse Bay Near Winnipeg River	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.018 0.018 0.009 0.009	0.13 0.07 0.06 0.04 0.13 0.09	0.21 0.18 0.15 0.15 0.14 0.12	0.46 0.46 0.42 0.42 0.42 0.37	0.46 0.44 0.36 0.39 0.35 0.27	0.73 0.73 0.68 0.65 0.66 0.61	2.04 2.10 2.18 2.21 2.13 2.25	2.27 2.42 2.23 2.10 2.15 1.87	2.59 2.40 2.19 2.11 2.08 2.05	2.77 2.81 2.26 2.20 2.14 1.90	0.013 0.013 0.013 0.013 0.027 0.013	<0.021 0.042 0.021 0.021 0.042 0.021
10 South Basin North End	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.018 0.018 0.018 0.018 0.018	0.13 0.16 0.14 0.11 0.09 0.14	0.30 0.32 0.30 0.30 0.30 0.30	0.61 0.64 0.61 0.61 0.66 0.59	0.66 0.69 0.63 0.58 0.61 0.60	1.04 1.12 1.07 1.06 1.12 1.04	1.81 1.95 1.93 1.99 2.03 2.02	3.40 3.63 3.56 3.42 3.58 3.27	3.30 3.50 3.18 3.14 3.30 3.18	4.79 5.08 4.73 4.87 4.93 4.63	0.080 0.053 0.053 0.080 0.067 0.053	0.084 0.042 <0.021 0.063 0.021 0.021
14 Narrows South End	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.018 0.018 0.018 0.018 0.018	0.08 0.07 0.06 0.11 0.10	0.23 0.22 0.23 0.23 0.26 0.26	0.51 0.49 0.51 0.53 0.51 0.49	0.58 0.69 0.65 0.69 0.63 0.63	0.99 0.89 1.00 0.97 1.00 0.89	2.40 2.40 2.39 2.40 2.40 2.40	2.71 2.58 2.77 2.75 2.85 2.65	2.36 2.33 2.46 2.52 2.54 2.45	2.96 3.02 3.36 3.36 3.38 3.28	0.040 0.040 0.040 0.040 0.053 0.040	0.021 0.021 <0.021 0.021 0.042 0.021
25 North Basin Near Saskat- chewan River	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.018 0.018 0.018 0.018 0.018	0.10 U.05 0.04 0.08 U.03 0.09	0.23 0.27 0.23 0.28 0.27 0.29	0.54 0.59 0.59 0.63 0.61 0.64	0.66 0.65 0.66 0.68 0.69 0.71	1.09 1.14 1.12 1.19 1.16 1.23	2.09 2.17 2.07 2.19 2.17 2.20	3.13 3.38 3.27 3.48 3.35 3.44	3.12 3.27 3.09 3.23 3.14 3.29	3.38 3.55 3.47 3.67 3.67 3.75	0.053 0.053 0.053 0.040 0.040 0.067	0.042 0.042 0.042 0.042 0.042 0.042
35 North Basin	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.018 0.018 0.018 0.018 0.018	0.11 0.09 0.10 0.14 0.13 0.08	0.30 0.32 0.22 0.30 0.30 0.32	0.66 0.66 0.46 0.68 0.73 0.68	0.79 0.76 0.56 0.79 0.79 0.77	1.23 1.29 0.99 1.33 1.38 1.28	2.26 2.26 1.84 2.27 2.27 2.26	3.56 3.75 2.79 3.58 3.65 3.67	3.43 3.53 2.51 3.33 3.41 3.41	3.91 4.12 3.00 4.14 4.38 4.24	0.067 0.040 - - 0.027	0.052 0.052 - - - -
39 North Basin	0-5 10-15 20-25 30-35 50-55 70-75	0.027 0.018 0.027 0.018 0.018 0.018	0.10 0.12 0.08 0.08 0.08 0.13	0.28 0.29 0.28 0.28 0.30 0.27	0.59 0.70 0.66 0.63 0.66 0.59	0.72 0.83 0.83 0.76 0.83 0.79	1.12 1.29 1.24 1.26 1.24 1.14	2.23 2.26 2.26 2.23 2.29 2.27	3.38 3.63 3.58 3.79 3.77 3.73	3.18 3.30 3.37 3.40 3.43 3.23	3.67 4.06 3.99 4.06 4.14 3.91	0.040 0.053 0.080 0.027 0.027 0.013	<0.021 0.042 0.063 - -
54 Narrows North End	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.018 0.018 0.018 0.018 0.018	0.08 0.07 0.10 0.06 0.10 0.14	0.27 0.28 0.29 0.30 0.27 0.29	0.54 0.54 0.61 0.61 0.63 0.53	0.79 0.72 0.74 0.74 0.63 0.66	1.04 1.04 1.09 1.12 1.04 0.97	2.36 2.31 2.32 2.34 2.36 2.41	3.06 3.29 3.31 3.33 3.21 3.00	2.80 2.95 3.04 3.03 2.88 2.91	3.51 3.93 4.20 4.18 3.95 3.73	0.107 0.080 0.067 0.080 0.080 0.053	0.042 0.084 0.063 0.042 0.042 0.021
57 South Basin	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.018 0.018 0.018 0.018 0.018	0.10 0.15 0.12 0.10 0.04 0.06	0.32 0.33 0.34 0.31 0.32 0.34	0.63 0.59 0.63 0.63 0.61 0.63	0.72 0.66 0.71 0.61 0.61 0.65	1.03 1.02 1.09 1.11 1.09 1.09	1.77 1.77 1.83 1.81 1.80 1.81	3.54 3.19 3.60 3.31 3.15 3.42	3.43 3.35 3.52 3.44 3.35 3.40	5.10 4.81 5.42 5.24 5.08 5.30	0.067 0.067 0.053 0.080 0.067	0.042 0.063 0.042 0.042 0.084 0.042

Table 23. Cont'd.

Station and Location	Depth in Core (cm)	Cd	Pb	Ве	Со	Cu	Ni μMoles	Sr s/gram	Cr	Zn	٧	As	Мо
59 South Basin	0-5 10-15 20-25 30-35 50-55 70-75	0.018 0.026 0.018 0.018 0.018 0.018	0.13 0.10 0.07 0.08 0.09 0.09	0.30 0.26 0.26 0.24 0.26 0.26	0.59 0.63 0.59 0.56 0.64 0.59	0.72 0.66 0.61 0.60 0.61 0.60	1.12 1.14 1.12 1.09 1.16 1.11	1.68 1.77 1.68 1.78 1.83 1.69	3.48 3.37 3.48 3.04 3.50 3.17	3.30 3.15 3.12 2.97 3.15 3.11	4.97 4.99 4.93 4.61 5.08 4.99	0.053 0.053 0.080 0.067 0.040 0.080	0.063 0.021 <0.021 <0.021 0.084 0.021
61 South Basin	0-5 10-15 20-25 30-35 50-55 70-75	0.027 0.018 0.018 0.018 0.018 0.018	0.12 0.12 0.10 0.08 0.08 0.10	0.22 0.24 0.22 0.26 0.26 0.26	0.49 0.53 0.51 0.54 0.54	0.77 0.58 0.55 0.58 0.83 0.58	1.02 1.04 0.99 1.02 1.04 1.06	1.50 1.60 1.53 1.54 1.54	3.19 2.79 3.00 2.65 2.88 2.79	3.21 3.17 2.95 3.00 2.97 3.01	4.69 4.67 4.57 4.73 4.73	0.093 0.053 0.053 0.053 0.053 0.053	<0.021 0.021 0.021 <0.021 <0.042 0.021

Table 24. Conce	Concentrations of mercury, basin Lake Winnipeg sedimen	arsenic, ot cores	and molybdenum taken in July 19	ybdenum in south July 1975.+
Station and Location	Depth in Core (cm)	Hg nMoles/gram	As µMoles	Mo es/gram
1 South Basin '	0-2 2-4 4-6 6-8 8-10 10-14 16-12 22-30 22-30 40-50 60-70	0.47 0.75 0.97 0.98 0.98 0.38 0.38 0.38	0.027 40.013 0.040 0.093 0.093 0.093 0.093 0.093 0.080 0.087 0.027	0.042 0.021 0.021 0.042 0.042 0.031 0.031 0.021 0.021 0.021 0.021
2 South Basin	0-2 4-6 6-8 6-8 8-10 10-14 11-18 18-22 22-30 30-40 50-60 60-70	0.48 0.60 0.75 0.75 0.75 0.67 0.30 0.30 0.30	0.013 0.093 0.080 0.040 0.040 0.067 0.013 0.023 0.027 0.050	0.021 0.021 0.021 0.021 0.021 6.021 0.021 0.021 6.021 6.021 6.021 6.021
3 South Basin	0-2 4-6 6-8 8-10 10-14 11-18 18-22 22-30 30-40 50-60 60-70	0.52 0.52 0.52 0.50 0.50 0.37 0.31 0.37	0.067 0.040 0.040 0.093 0.067 0.067 0.027 0.027 0.027	60.021 60.021 60.021 60.021 60.021 60.021 60.021 60.021 60.021
4 South Basin	0-2 2-4 4-6 6-8 8-10	0.75 0.47 0.47 0.50 0.47	0.067 0.040 0.040 0.040 0.040	0.021 0.021 0.042 <0.021 0.021

Table 24. Cont'd.

Station and Location	Depth in Core (cm)	Нg nMoles/gram	As µMo1es	s yMoles/gram
4 South Basin	10-14 14-18 18-22 22-30 30-40 40-50 50-60 60-70	0.30 0.43 0.47 0.40 0.20 0.27	<ul> <li>6.013</li> <li>0.013</li> <li>0.053</li> <li>0.040</li> <li>0.053</li> <li>0.053</li> <li>0.053</li> <li>0.053</li> <li>0.053</li> <li>0.053</li> <li>0.053</li> </ul>	60.021 60.021 60.021 60.021 60.021 60.021
South Basin	0-2 2-4 6-8 6-8 10-14 114-18 18-22 22-30 22-30 50-60 60-70	0.60 1.42 0.88 0.56 0.78 0.54 0.54 0.54 0.54	0.080 0.013 0.013 0.013 0.027 0.040 0.040	6.021 0.042 0.042 0.042 0.021 0.031 0.031 0.031
6 South Basin	0-2 2-4 4-6 6-8 8-10 10-14 14-13 14-22 22-30 30-40 50-50 60-70			0.021 0.042 0.042 0.042 0.042 0.042 0.021 0.021 0.021
7 South Basin Traverse Bay	0-2 2-4 4-6 6-8 8-10 10-14 14-18	0.27 0.40 0.47 0.34 0.43 0.33	0.053 0.067 0.027 0.093 0.053 0.067	0.021 0.021 <0.021 <0.021 0.021 0.031 0.031 <0.021

Table 24. Cont'd.

Station and Location	Depth in	Hg	As	Мо
- COCAL TON	Core (cm)	nMoles/gram	µМо1е	s/gram
8	0-2	0.40	.027	0.021
_	2-4	0.40	.053	0.021
raverse Bay	4-6	0.27	.053	0.021
	6–8	0.27	.053	<0.021
	8-10	0.21	.027	0.042
	10-14	0.17	.027	<0.021
	14-18	0.14	.013	0.031
	18-22	0.14	<.013	0.042
	22-30	0.16	.013	0.042
	30-40	0.14	.013	0.031
	40-50	0.14	.027	0.031
	50-60	0.14	.013	<0.021
9	0-2	0.60	.067	<0.021
	2-4	0.53	.040	<0.021
raverse Bay	4-6	0.34	.053	0.021
	6-8	0.27	.093	<0.021
	8-10	0.27	.067	<0.021
	10-14	0.27	.027	0.021
	14-18	0.24	.040	0.021
	18-22	0.24	.027	0.031
	22-30	0.22	.013	0.021
	30-40	0.21	.013	0.021
	40-50	0.22	.013	<0.021
	50-60	0.21	.013	<0.021
	60–70	0.14	< .013	< 0.021

<sup>†</sup> These data were provided to us by Dr. R. J. Allan, and the total element concentrations were measured by radio frequency plasma emission spectroscopy. Station locations are given in Fig. 24.

Table 25. Overlying lake water and interstitial water chemistry (in pmoles/L) of sediment samples (0-20 cm homogenized) from Lake Winnipeg, collected October 3-12, 1969, squeezed February 1970. Interstitial water samples were immediately filtered through a Whatman GF/C filter. LW = Lake water, IW = Interstitial water, (-) = Not measured.

tation	Ca		Mg		Na		К		Cl		. SO <sub>1</sub>	+	HCO <sub>3</sub>	Si		Fe	Mn
	LW	IW	LW	WI	LW	IW	LW	IW	LW	IW	LW	IW	LW	LW	IW	IW	IW
01	-	1525		1872	_	2109	_	34	_	1211	_	-	2740	186	-	3	35
02	1015	2475	790	2243	900	1630	118	46	379	789	583	-	-	123	-	23	152
03	-	3025	-	3004	-	1243	-	269	412	546	593	13	2860	123	694	$\frac{1}{1}$	64 77
03C	-	1988	-	1564	-	965	-	157	158	721	250 364	13	1420 1880	68 86	925 -	<0.4	77
04	716	1150	502	1111	883	1109	76	174 120	237	507 310	30 <del>4</del>	26	-	-	758	1	66
048	-	1288	-	945	-	648 765	-	113	_	372	_	13	_	134	865	. 1	64
05	-	1301	-	1029 872	-	591	_	552	_	676	_	13	_	103	769	ī	54
06 07	- 324	1163 1668	173	741	- 78	461	18	58	39	828	62	_	_	66	_	36	46
07 08	334	800	202	691	87	365	21	59	39	197	62	_	800	70	-	20	44
10	-	810	-	593	-	478	-	90	175	197	323	<13	1720	133	936	2	69
12	_	770	_	761	_	578	-	101	-	254	_	-	-	129	-	<0.4	58
14	599	740	395	774	287	565	58	113	152	225	239	-	1520	109	-	_	59
15	_	1988	-	1325	-	357	-	59	-	-		13	-	43	-	2	78
17	_	605	-	461	-	626	-	818	327	451	167	350	1350	38	665	2	17 31
18	422	1075	284	765	296	796	38	190	203	535	146	26	1160	40	-	49	12
21	691	715	494	700	565	1000	58	123	429	676	239	-	1940 2220	33		<0.4 <0.4	36
230	723	1475	551	576	639	687	61	104	305	535 479	281 271	-	1980	23 26	-	<0.4	38
23E	736	1350	518	638	639	804 605	61	86 59	496	394	4/1	38	- 1500	-	_	<0.4	57
23H	-	1063	-	716	- 648	696 687	- 61	57	513	451	250	13	2020	29	_	<0.4	63
25	721	1050 660	502	720 802	-	817	- 01	147	712	451	-	-	-	36	_	<0.4	54
26	-	2575	_	1626	-	848	_	86	_	423	_	625	-	36	-	<0.4	96
27 31	-	1850	-	1169	_	8696	_	143	_	569	_	50	_	27	_	22	82
34	_	1175	_	757	_	678	_	59	_	366	-	26	-	27	-	2	40
35	793	2190	539	630	626	774	60	94	429	535	177	-	8240	19	-	<0.4	54
39	753	830	526	588	648	648	53	57	468	423	250	50	2120	18	-	<0.4	30
41	851	3543	588	630	617	904	60	91	389	451	208	-	2300	26	-	<0.4	29
47	-	1200	-	872	<del>-</del>	1009	-	109	-	845	-	100	1040	23	-	<0.4	32 36
48	589	590	428	576	548	870	52	86	440	563	239	- 363	1640 -	36 40	-	57 <0.4	26
50	-	685	-	473	-	328	-	32	21	225	42	202	310	73	_	3	161
51	-	1600	103	845 658	78 -	491 600	13	76 78	34 107	197 197	125	-	1020	48	_	98	26
52	-	745 670	-	617	_	543	-	83	-	254	_	_	-	-	_	8	31
53 54	- 536	760	358	597	252	587	47	87	147	225	156	_	1340	40	_	26	39
55	-	1088	-	823	-	385		101	. –	197	_	375	_	_	587	1	44
57	654	2819	457	733	413	665	74	107	181	338	323	-	1820	140	-	<0.4	80
59	619	3019	440	774	378	709	63	244	158	366	250		-	130	-	<0.4	62
59B	-	1250	-	938	-	991	-	81	_=_	197	<del>-</del>	63	_	-	_	1	131
60	1000	2343	769	774	870	652	116	134	372	366	604	-	2640	118	770	<0.4	73
60B	996	1838	790	1374	861	743	119	141	367	197	65	-	-	194	779	1	101
60C	-	1500	-	1095	-	826	-	137	-	394	_	688	-	70	762 826	1 2	49 73
60D	-	1150		897	-	539	-	102	210	225	489	13	- 2340	100	820	<0.4	76 76
61	868	1791	650	794	709	761	101 52	157 107	310 491	422 845	219	_	1420	42	_	<0.4	16
64	549	1804	403	617	557	1122 922		141		507		_	1420	37	-	<0.4	126
68	-	2725	-	1975	-	922 426	-	95	- -	225	-	963	_	38	402	2	11
69	_	770	-	584	-	426	-	95	-	223	-	303	-	36	706	L	

Table 26. General definition of graphic moment measures applied to sediment particle size distributions (modified after Thomas  $\underline{et}$   $\underline{al}$ ., 1976).

Parameter	Moment	Value for normal distribution	Comments
Mean particle size	First moment about zero	any	Large positive values = fine sediments.
Standard Deviation	Second moment about the mean	>0	Large values = poor sorting of sediment particle sizes.
Skewness	Standardized th moment about th mean		Relative to a normal distribution: positive values = extension of the tail of the distribution towards fine sediment particle sizes.
			Negative values = extension of the tail of the distribution towards coarse particle sizes.
Kurtosis	Standardized fo moment about th mean		Relative to a normal distribution: large positive values = distribution of sediment sizes narrowly peaked at the mean.
			Large negative values = distribution of sediment sizes flat or bimodal.

1	1 1																					
ation, icle sizes Our	Kurtosis		1.64	0.973	0.984	1.10 2.1	0.981	1.02	1.25	1.07	1.34	1.21	1.01 1.03 1.02	0.916			0.798 0.778 0.751	0.710	0.987	0.973	770.0	67.0
size, standard deviation, istribution of particle sizes s given in Table 2. Our 1 (1957).	Skewness		0.479	-0.2/1 -0.009	-0.107 -0.286	-0.148	-0.172	0.022	-0.236	-0.144	-0.29/	-0.223 -0.314	-0.168 -0.238 -0.194	-0.384			-0.074 0.187 -0.076	-0.483	-0.246	-0.150	T67'n=	-0.272
(mean particle size, standard tosis) of the distribution of sediment samples given in Tablow Folk and Ward (1957).	Standard Deviation	South Basin	1.91	2.14 1.71	1.83 2.76	1.74	2.02	2.20 1.66	1.58	1.54	1.72	1.55	1,64 1,73 1,63	2 40	•	Narrows	2.45 2.47 2.59	2.33 2.68	1,96	1.74	0/17	2.59
Moment parameters (mean particle skewness, and kurtosis) of the disof Lake Winnipeg sediment samples calculations follow Folk and Ward	Mean 🏚		4.20 8.43	8.23 8.00	8.20 7.43	8.70	8.00	6.40 8.73	9.40	8.97	9.20 9.83	9.30 9.07	8.93 8.93		÷		6.93 6.13 7.03	8.77	8.50	7.40	7.33	7.53
Table 27. Morski	Station		01 02	33. 03.C	03D .	048	09 00	07	92	27	29 298	90 909	600 60D 61	Composite of	פון ארפרותנו		14 15 51	52	54	55 55 55	ň	Composite of all stations

Table 27. Cont'd.

Station	Меал ф	Standard Deviation	Skewness	Kurtosis
;   		North Basin		
17 221 231 232 233 233 333 333 44 44 44 45 50 50 50 50 50 50 50 50 50 50 50 50 50	5.63 8.37 8.37 7.10 7.10 7.14 7.10 8.87 7.97 8.40 8.40 8.40 8.03 9.10 9.10	2.87 1.588 1.77 1.77 1.62 1.62 1.51 1.51 1.63 1.65 1.65 1.65 1.65 1.65 1.65 1.84	0.611 0.163 0.163 0.163 0.026 0.026 0.030 0.365 0.069 0.009 0.108 0.108 0.009 0.108 0.	0.611 0.993 1.0597 1.02 1.02 0.984 1.23 1.02 1.02 0.998 0.998 0.990 0.990 0.990 0.643
Composite of all stations	7.43	2,55	-0.230	0.78

Table 28. Clay mineralogy of the total clay fraction (<4 µm) of selected south basin Lake Winnipeg sediment samples. This data is from Kushnir (1971).

1	Kaolinite and Chlorite†	13	14	13	12	15	18	13	14	14	13	11	14	15	13	11	13	26	17	11	10	
of Clay Fraction	Illite	34	33	39	28	36	34	27	32	34	27	27	30	49	27	28	40	45	40	28	29	1
94	Montmorillonite	53	53	48	90	49	48	60	54	25	99	29	56	36	90	61	47	53	43	61	61	ć
	Station No.	. 00	01	02	03	350	030	04	048	05	90	20	08	10	. 21	23	59	598	09	60B	209	1 4 4

+ Kaolinite was found to be more abundant than chlorite in most samples subjected to heat treatment.

Table 29. Comparison of present-day sedimentation rate data for Lake Winnipeg and the St. Lawrence Great Lakes.

Winnipeg (Red River Basin)* 11: Erie¹ 28			average inickness of sediment per year (mm)	Range
•	1170	0-1970	1.1	0-2
	2880	0-5050	6.1	0~24
Ontario <sup>2</sup> 56	563	0-1156	2.5	9.6-0
Huron³ 26	266	65-495	1.2	0.3-B
Superior <sup>4</sup> 36	380	25-780	1.2	
Michigan <sup>5</sup> 14	147	0-165	0.8	0-3.2

Minimal sedimentation rate estimate, based on annual Red River suspended sediment supply only, see Table 19.

 $<sup>^1\,</sup>$  Data from Kemp et al. 1976; Sly & Thomas 1974; Kemp et al. 1974.

<sup>&</sup>lt;sup>2</sup> Data from Thomas et al. 1972; Sly & Thomas 1974.

<sup>3</sup> Data from Kemp et al. 1974; Kemp et al. 1978.

<sup>4</sup> Data from Bruland et al. 1975; Kemp et al. 1978.

<sup>5</sup> Data from Edgington & Robbins 1976.

Table 30. Comparison of average, deep basin, elemental sediment supply rates.

California Coastal Basins <sup>3</sup>	l	•	t	322	2.7	1000	0.08	0.2	0.09	90.0	0.8	0.3	600.0	0.08	1
Lake Ontario <sup>2</sup> (Range)	(491-842)	(171-630)	(23-52)	1	•	•	1.0	99.0	0.22	1.5	0.57	3.2	0.020	0.48	0.0047
Lake Erie <sup>2</sup> (Range)	(770-8710)	(300-1630)	(32-213)	1	ı	ı	2.4	1,3	0.70	3.7	1.6	6.1	0.044	0.72	900.0
Lake Huron <sup>2</sup>	935	94	8.4		,	ı	0.38	0.27	0.12	0.19	0.37	0.84	0.005	0.18	0,0005
Lake Superior <sup>2</sup>	1060	102	19	1	1	,	0.34	0.93	0.14	0.33	0.70	0.98	0.009	0.20	0,0004
Lake Winnipeg <sup>1</sup> (Red River Basin only)	(4,000)	(420)	22(34)	480	21	2950	0.06	9.0	0.5	2.9	4.6	3.3	0.02	0.06	0.014
Element (mMoles m <sup>-2</sup> yr <sup>-1</sup> )	IJ	z	<b>c</b> .	Fe	Mn	A1	Ni	7	00	ភ	>	Zn	p	Pb	Ħg

Probably a minimal estimate, based on Red River sediment supply only. See Table 19. Data in parentheses are from twice monthly sampling of the Red River at Lockport, Manitoba. In this column, C is total carbon = organic plus inorganic C.

Table 31. Comparison of 1926 and 1969 sediment particle size data for Red River and Lake Winnipeg bottom sediments.+

	% Sand	% Silt	% Clay
Ward's Lake Winnipeg Station, 4 miles north of the mouth of the Red River (1926)	6.82	36	57.13
Our Station 4, 4½ miles off mouth of Red River (1969)	19	36	44
Ward's St. Vital Bridge sample of Red River sediments (1926)	6.8	57.8	35.1
Our Red River sediment sample at the St. Vital Bridge (1969)	.e.	44.9	. 51.5

<sup>+</sup> The 1926 data are from Ward (1926), and the 1969 data are from Table 2. Ward's size limits were probably as follows: Clay = <4 µm, silt = 4-62.5 µm, and sand = >62.5 µm. These limits are slightly different than ours (see Table 2). The St. Vital Bridge is in the City of Winnipeg.

pius norganic t.  $^2$  Data from Kemp et al. (1974, 1976, 1978).  $^{\rm C}$  = organic carbon.

<sup>3</sup> Data averaged form Bruland et al. (1974).

<sup>&</sup>lt;sup>4</sup> Estimated from Derksen (1973) and Armstrong and Hamilton (1973), assuming no release from sediment particles.

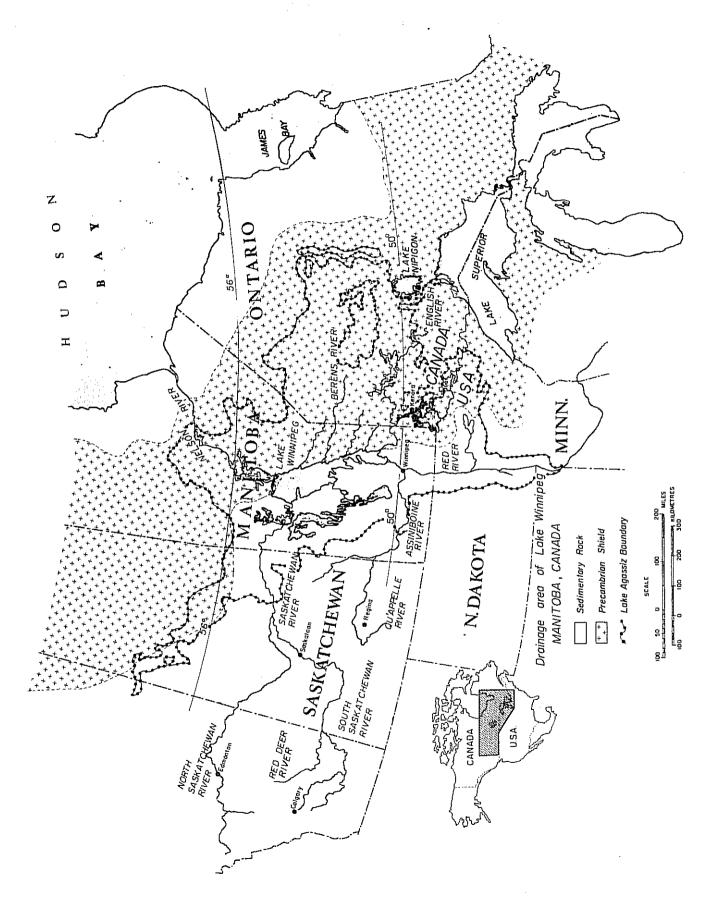


Fig. 1. Map of the watershed of Lake Winnipeg, its major tributaries, and the maximum extent of glacial Lake Agassiz sediments (dashed line, from Elson'1967).

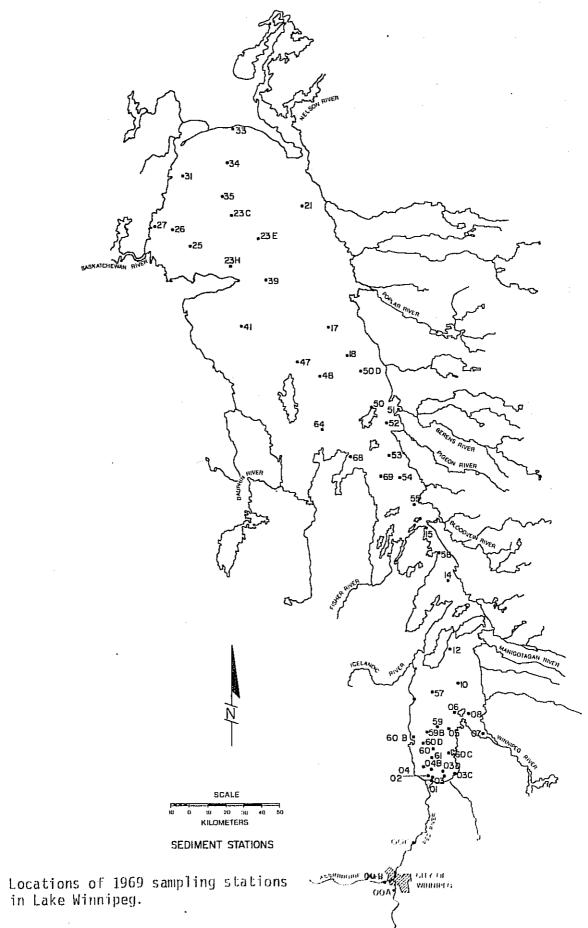


Fig. 2.

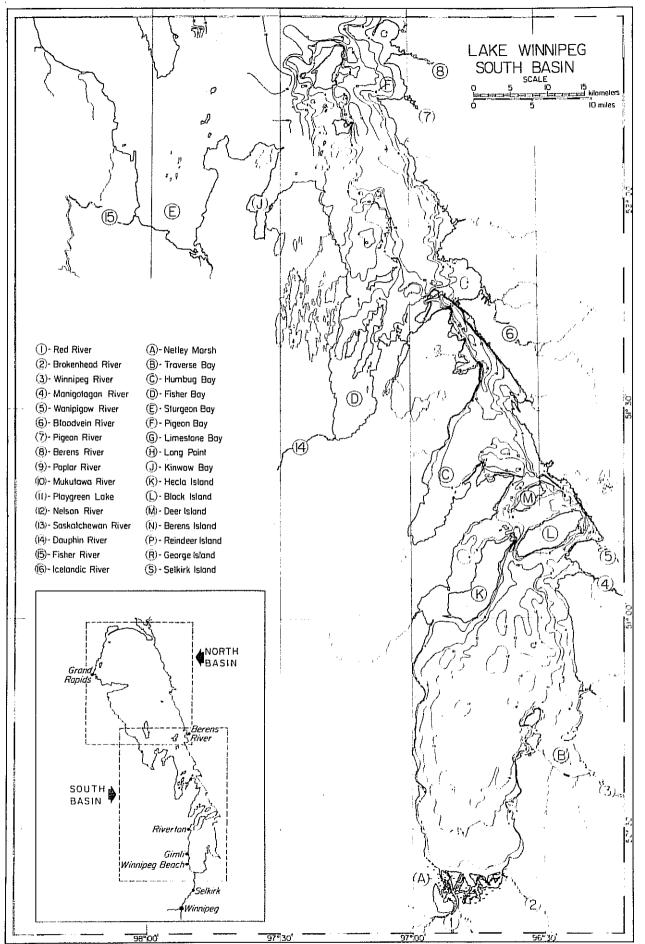


Fig. 3. Bathymetry of the south basin and narrows of Lake Winnipeg, from Brunskill, Elliott, and Campbell (1979). Depth contours are in meters.

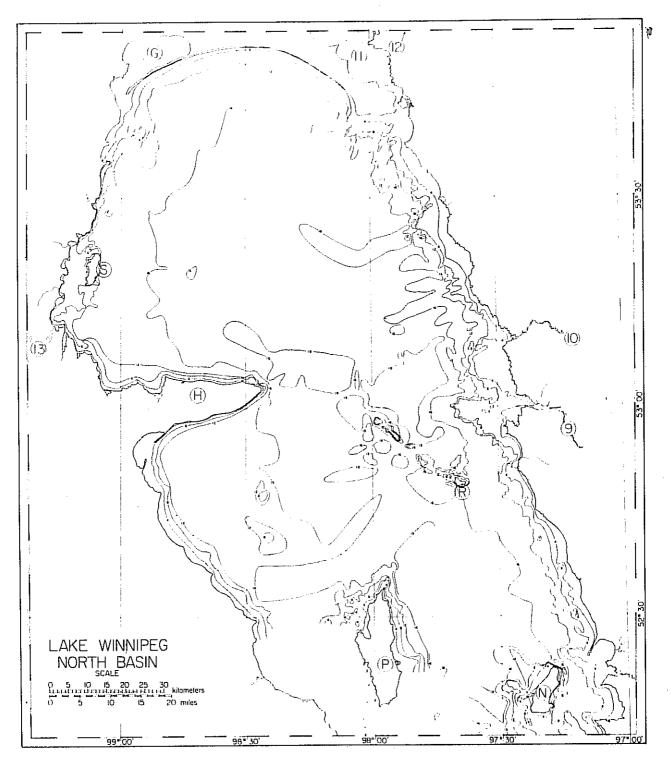


Fig. 4. Bathymetry of the north basin of Lake Winnipeg, from Brunskill, Elliott, and Campbell (1979). Depth contours are in meters.

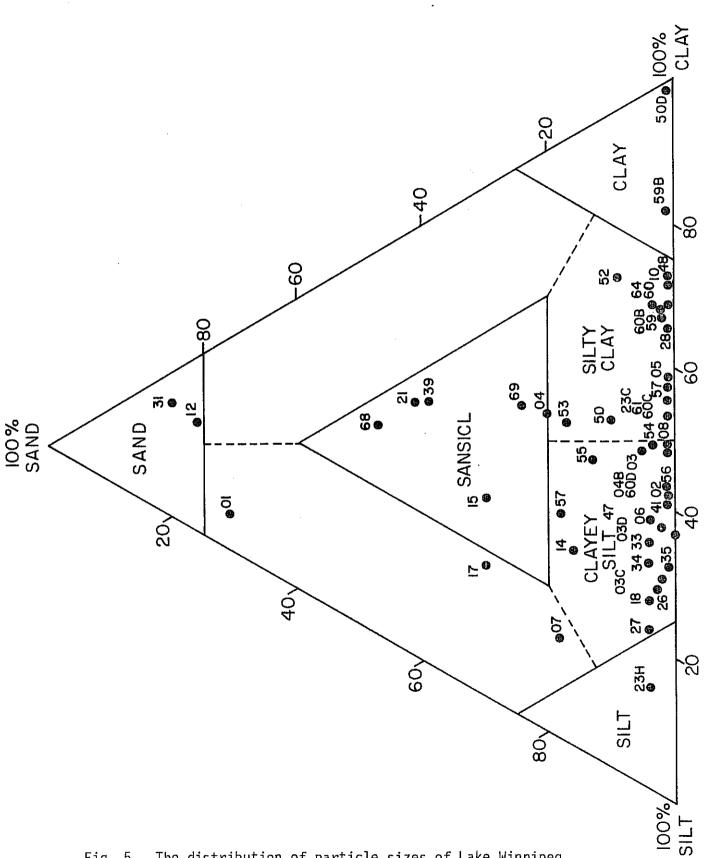


Fig. 5. The distribution of particle sizes of Lake Winnipeg sediments, according to Shepard (1954). Station numbers are from Fig. 2, and Tables 1 and 2 give data for this diagram.

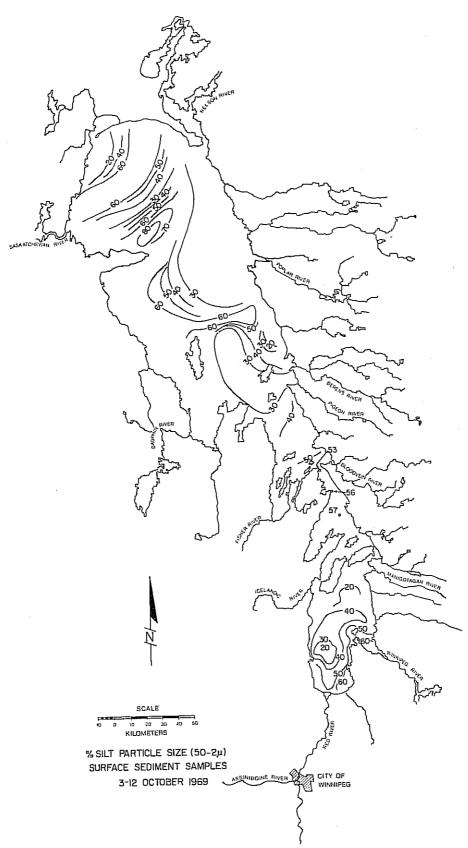
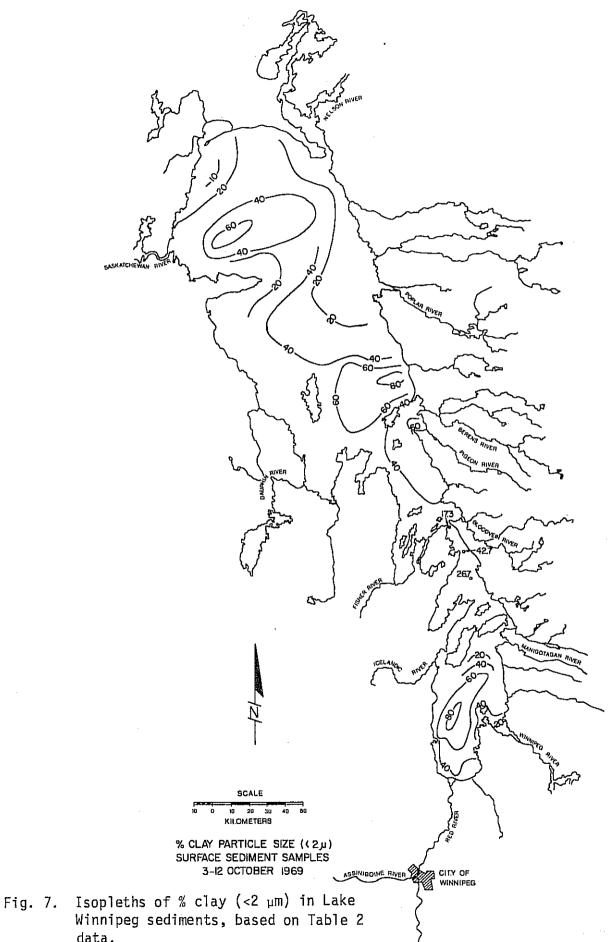


Fig. 6. Isopleths of % silt (50-2  $\mu m)$  in Lake Winnipeg sediments, based on Table 2 data.



data.



Fig. 8. Isopleths of % sand (2000-50  $\mu$ m) in Lake Winnipeg sediments, based on Table 2 data

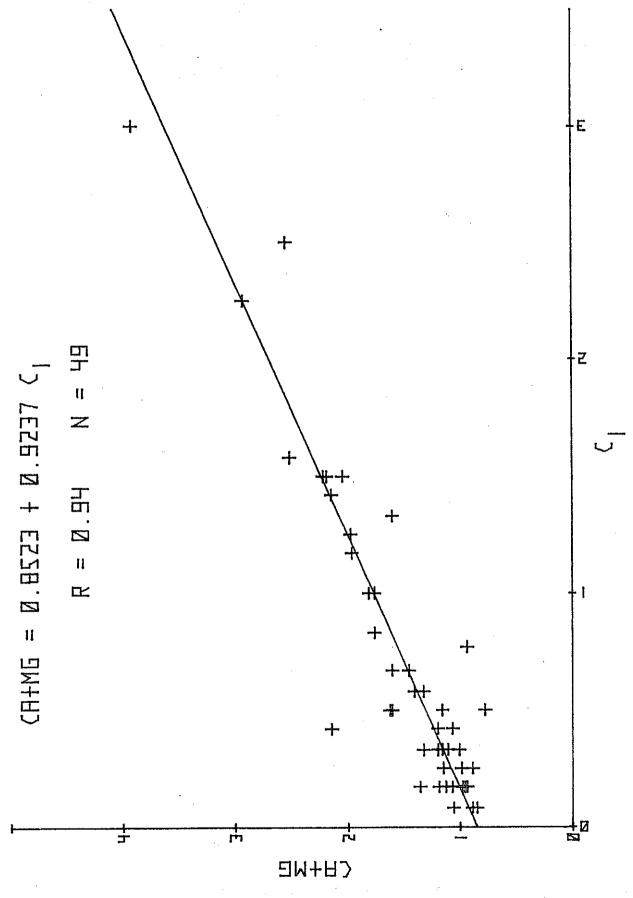


Fig. 9. The relationship between the sum of Ca and Mg (in mMoles/g) dry weight), and inorganic carbon ( $C_i$ ) for sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

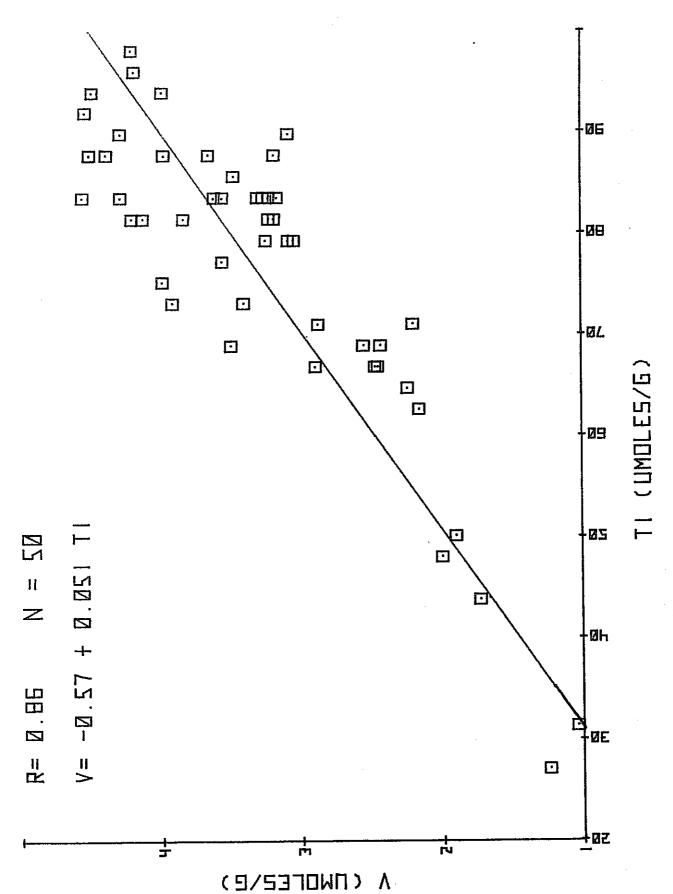


Fig. 10. The relationship between vanadium (V) and titanium (Ti), in  $$\mu$moles/g$  dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

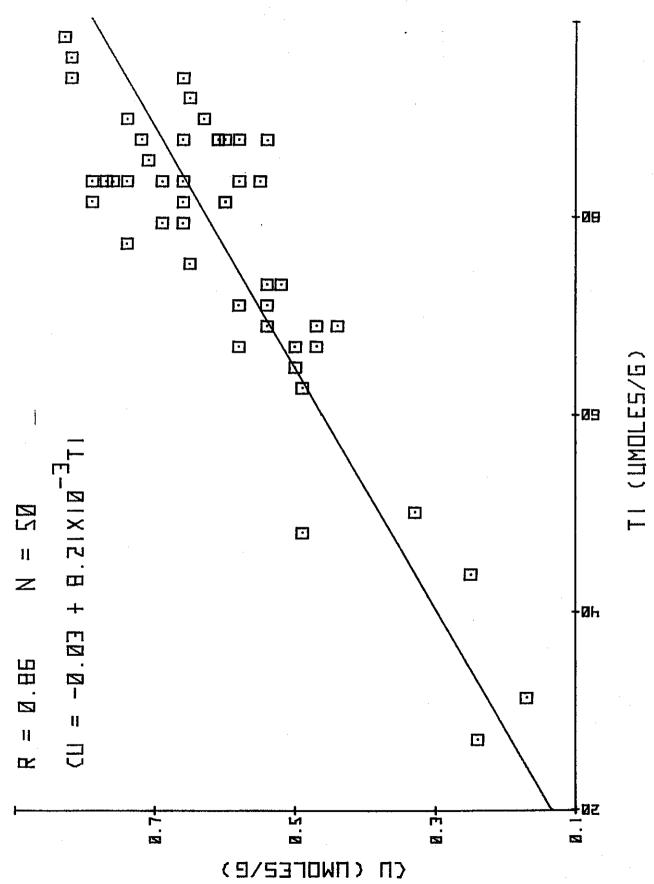


Fig. 11. The relationship between copper (Cu) and titanium (Ti), in  $$\mu$moles/g$  dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

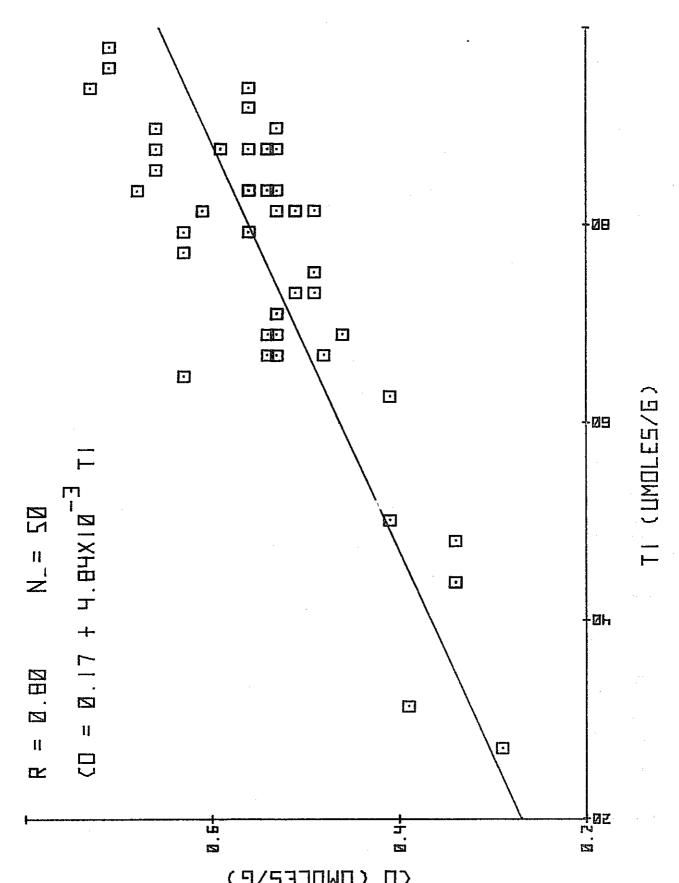


Fig. 12. The relationship between cobalt (Co) and titanium (Ti), in μmoles/g dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

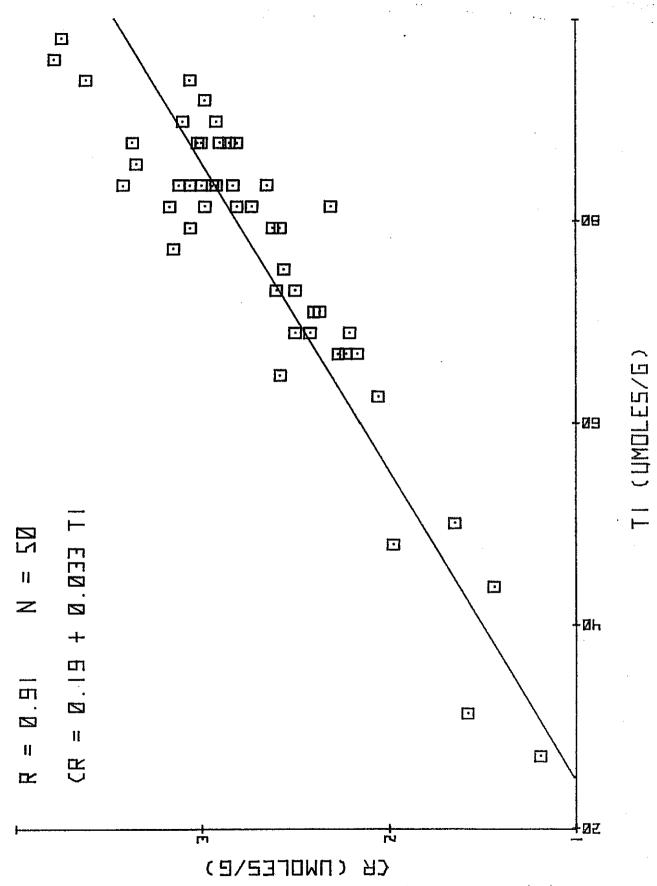


Fig. 13. The relationship between chromium (Cr) and titanium (Ti), in μmoles/g dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

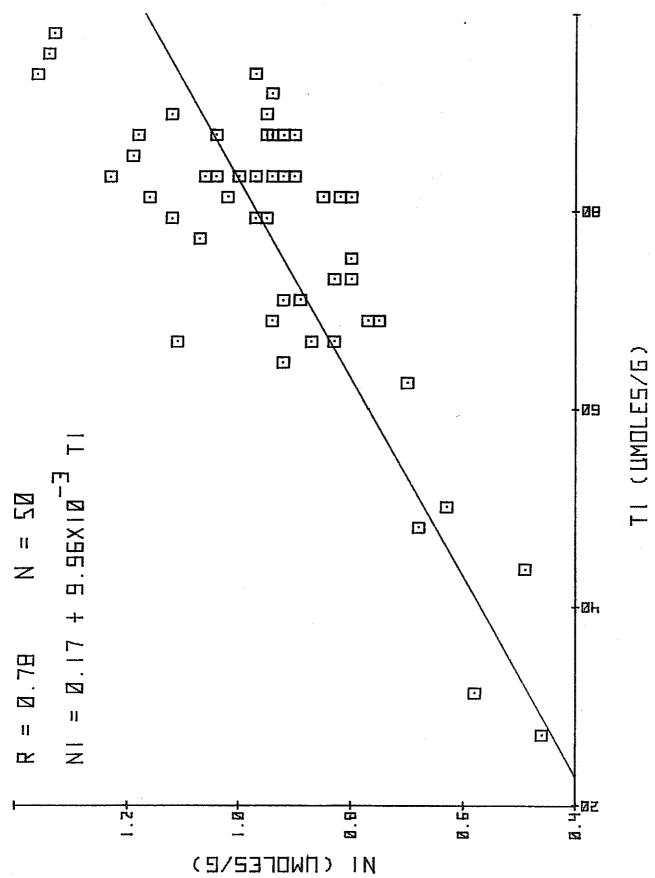


Fig. 14. The relationship between nickel (Ni) and titanium (Ti), in  $_{\mu}$  moles/g dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

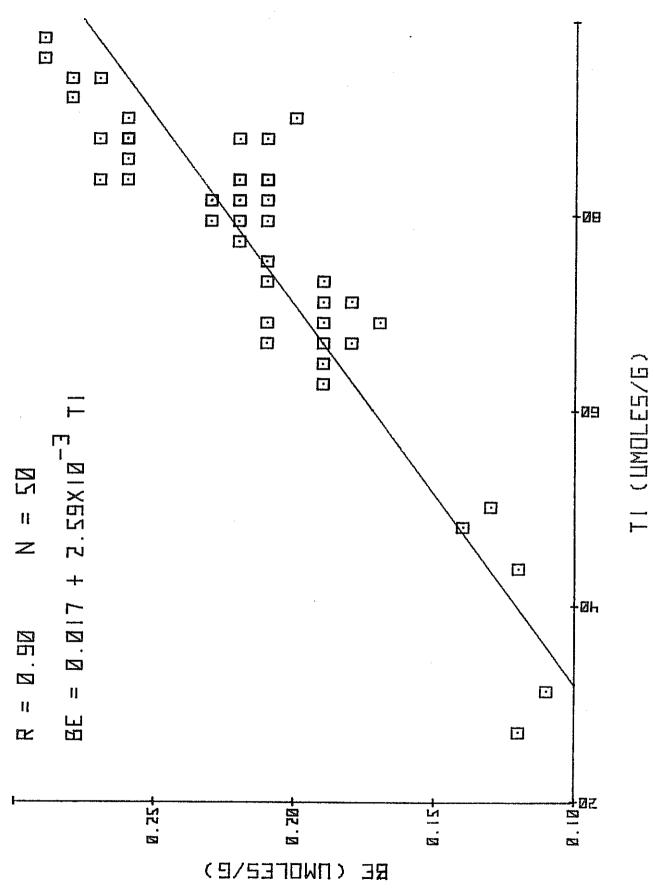


Fig. 15. The relationship between beryllium (Be) and titanium (Ti), in  $_{\mu}moles/g$  dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

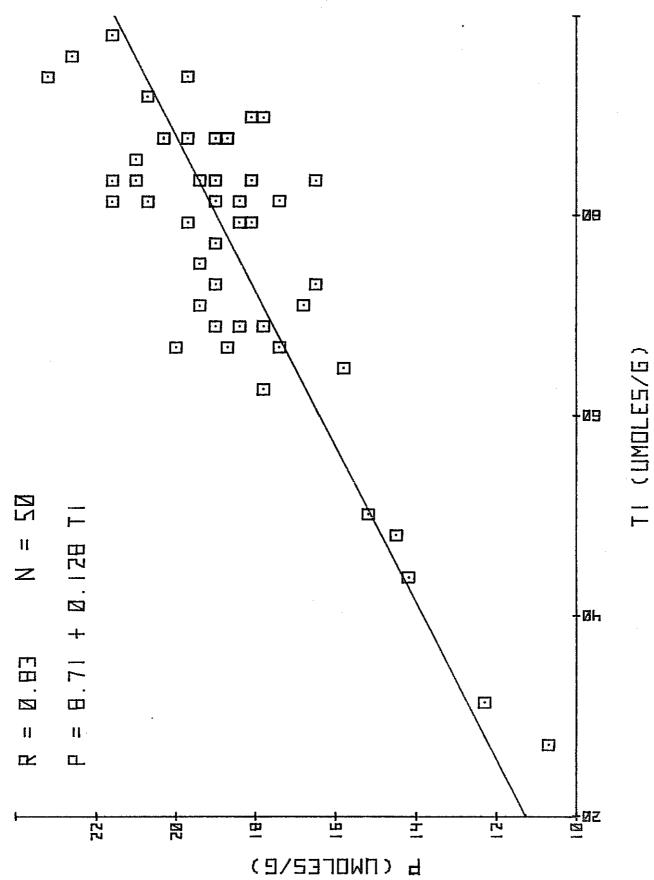


Fig. 16. The relationship between phosphorus (P) and titanium (Ti), in  $_{\mu}$ moles/g dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

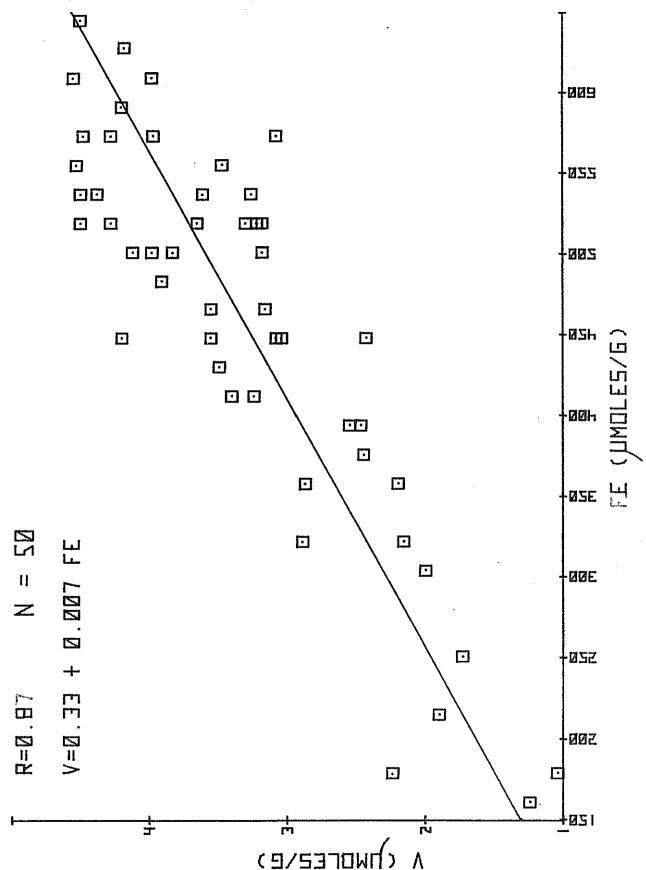
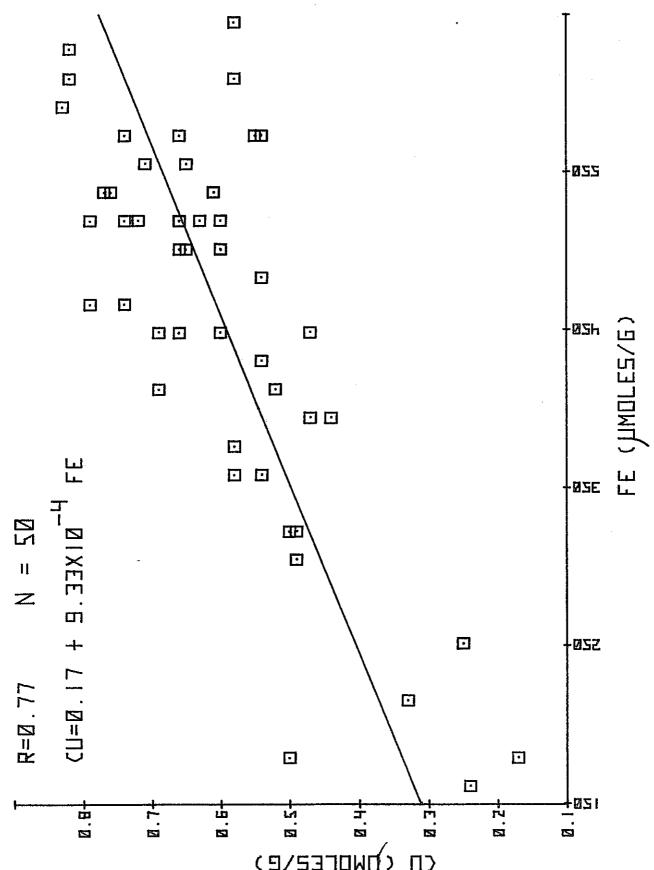


Fig. 17. The relationship between vanadium (V) and iron (Fe), in  $$\mu$moles/g$  dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.



(5/5370WΓ) Π)

Fig. 18. The relationship between copper (Cu) and iron (Fe), in μmoles/g dry weight, for surface sediment samples from Lake Winnipeg.
r = correlation coefficient, n = number of samples.

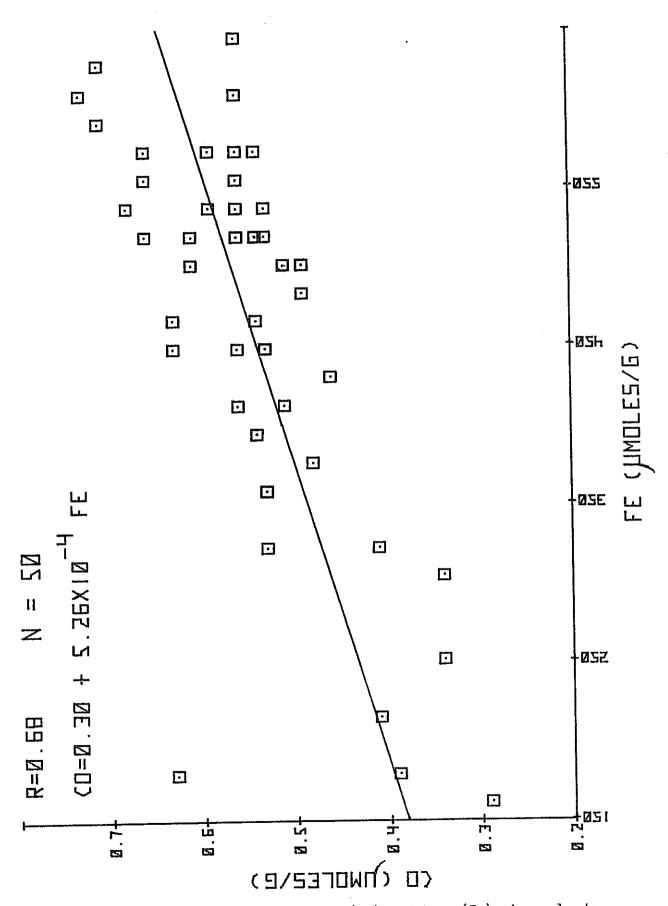


Fig. 19. The relationship between cobalt (Co) and iron (Fe), in pmoles/g dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

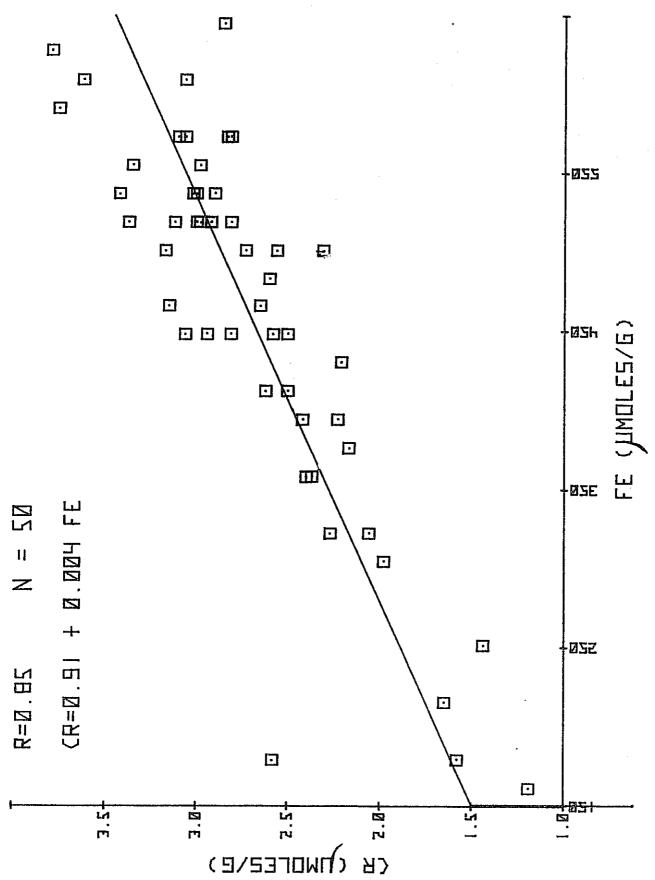


Fig. 20. The relationship between chromium (Cr) and iron (Fe), in  $\mu$ moles/g dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

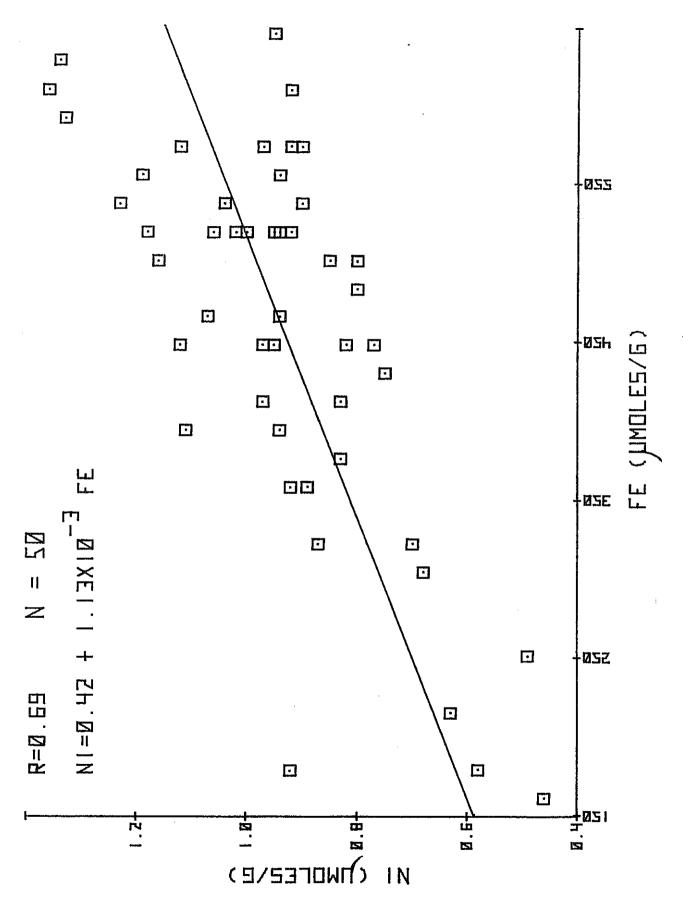


Fig. 21. The relationship between nickel (Ni) and iron (Fe), in  $\mu$ moles/g dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

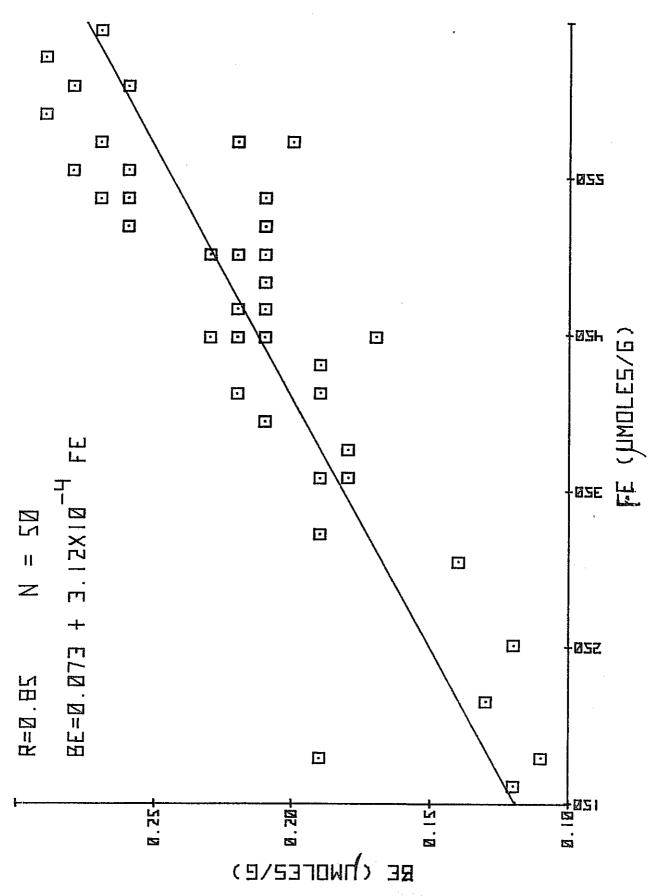


Fig. 22. The relationship between beryllium (Be) and iron (Fe), in  $\mu moles/g$  dry weight, for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

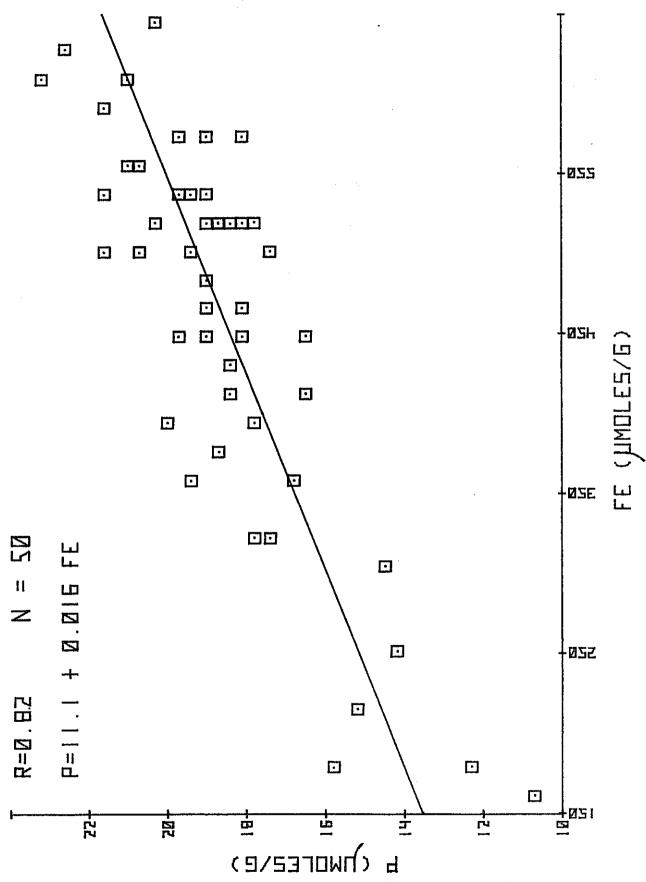


Fig. 23. The relationship between phosphorus (P) and iron (Fe), in  $\mu$ moles/g dry weight; for surface sediment samples from Lake Winnipeg. r = correlation coefficient, n = number of samples.

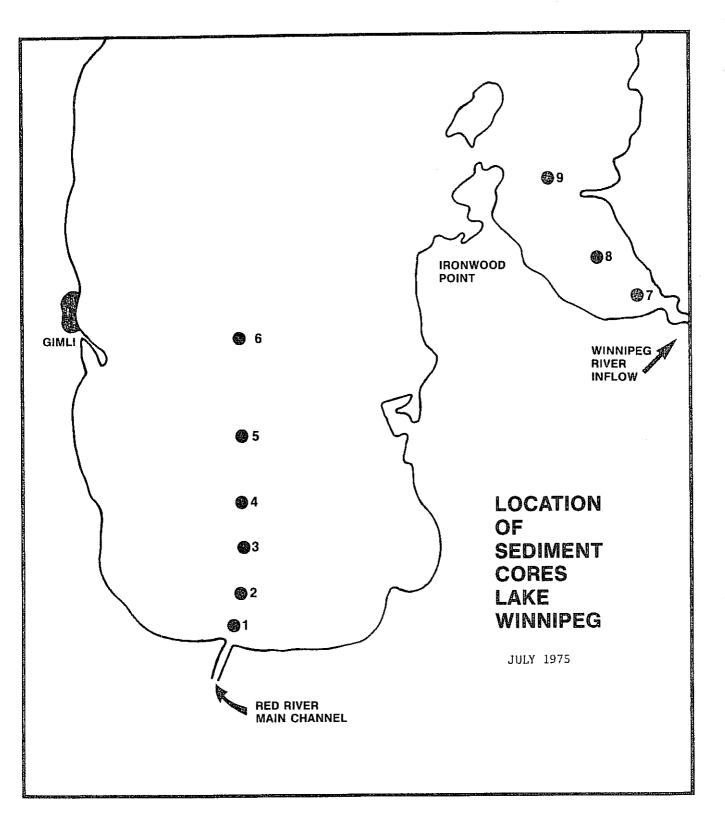


Fig. 24. Location of Lake Winnipeg sediment cores taken by Dr. R. J. Allan in 1975. These cores were analyzed for Hg, As, and Mo, and are reported in Table 24.

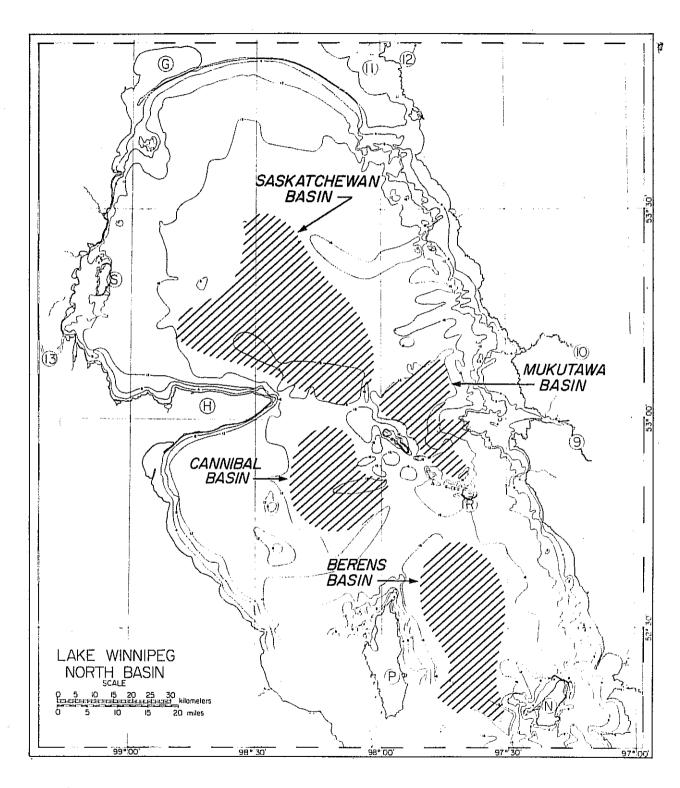


Fig. 25. Major regions of sediment accumulation in the north basin of Lake Winnipeg. The stippled areas are the depositional basins. See Fig. 26 for key to geographic features. Depth contour intervals are 4 m.

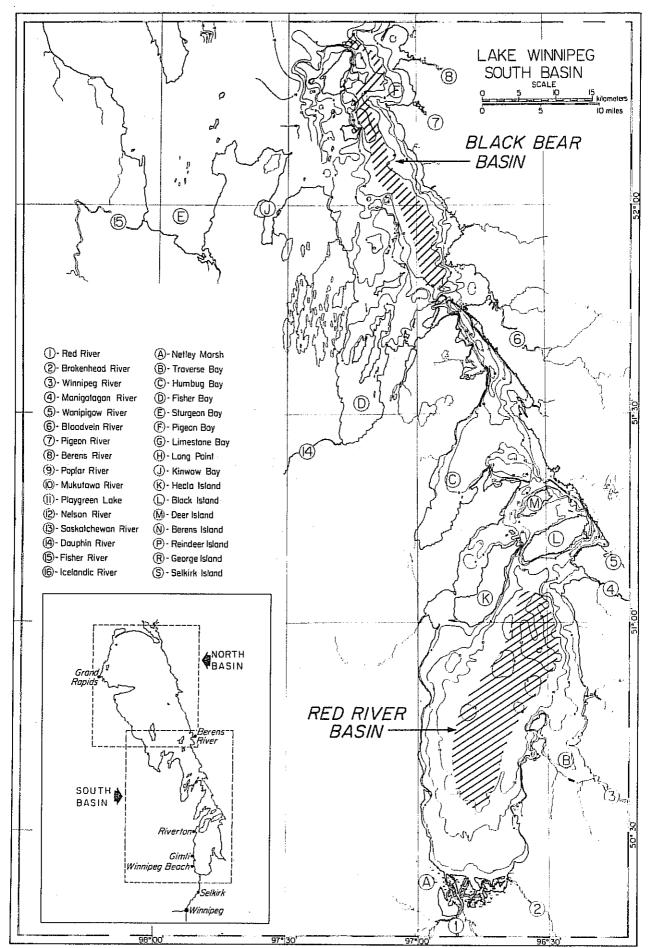


Fig. 26. Major regions of sediment accumulation in the narrows and south basin of Lake Winnipeg. The stippled areas are the depositional basins.

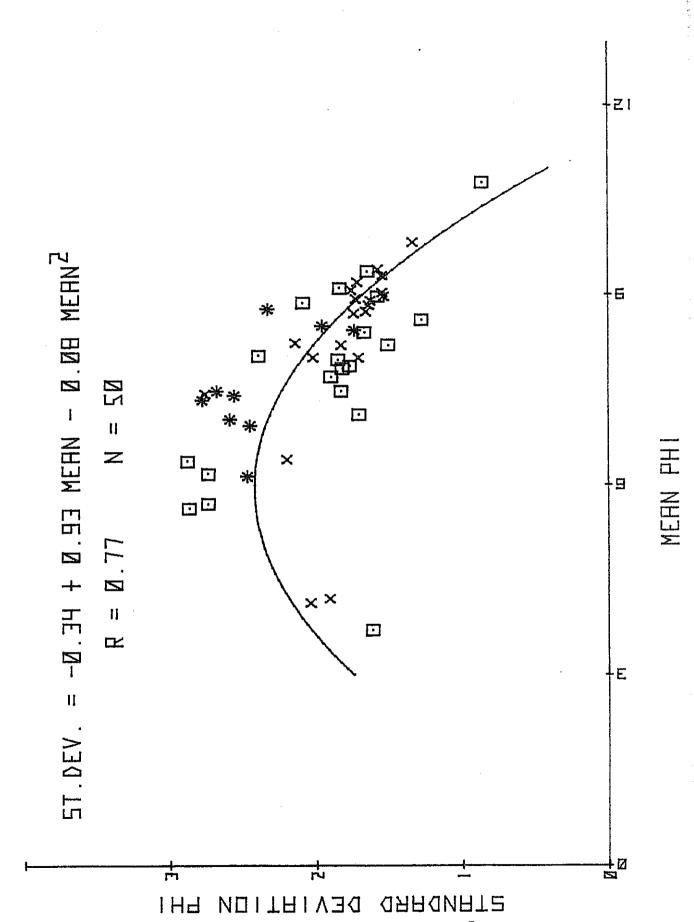


Fig. 27. The relationship between mean particle size  $(\bar{X}_{\varphi})$  and graphic standard deviation of  $X_{\varphi}$  for Lake Winnipeg off-shore sediment samples. X = south basin; \* = narrows;  $\square$  = north basin.

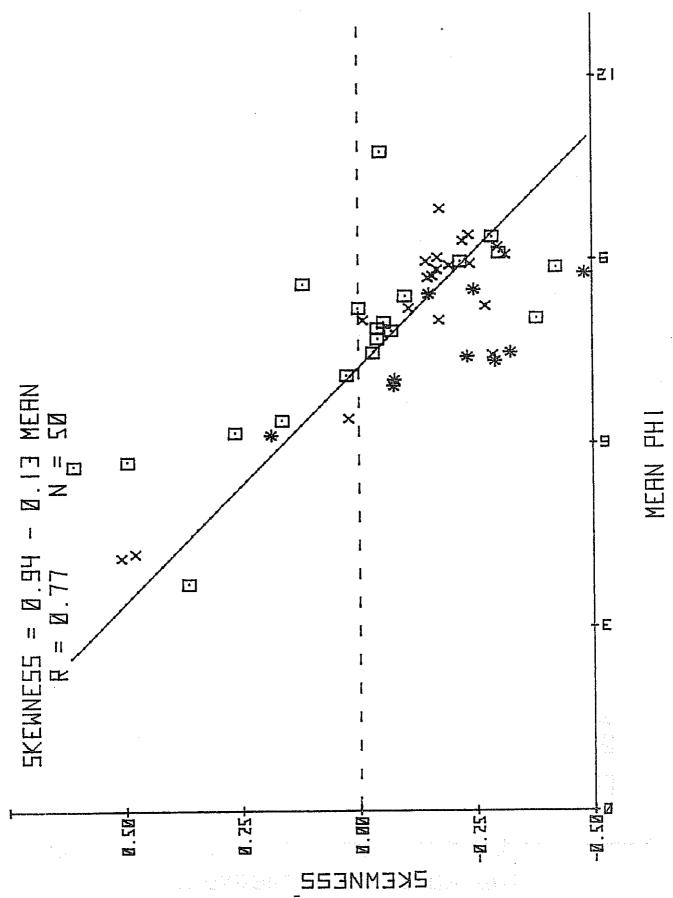


Fig. 28. The relationship between  $\bar{X}_{\varphi}$  and graphic skewness of the particle size distributions of Lake Winnipeg off-shore sediments. X = south basin; \* = narrows;  $\square$  = north basin sediments.

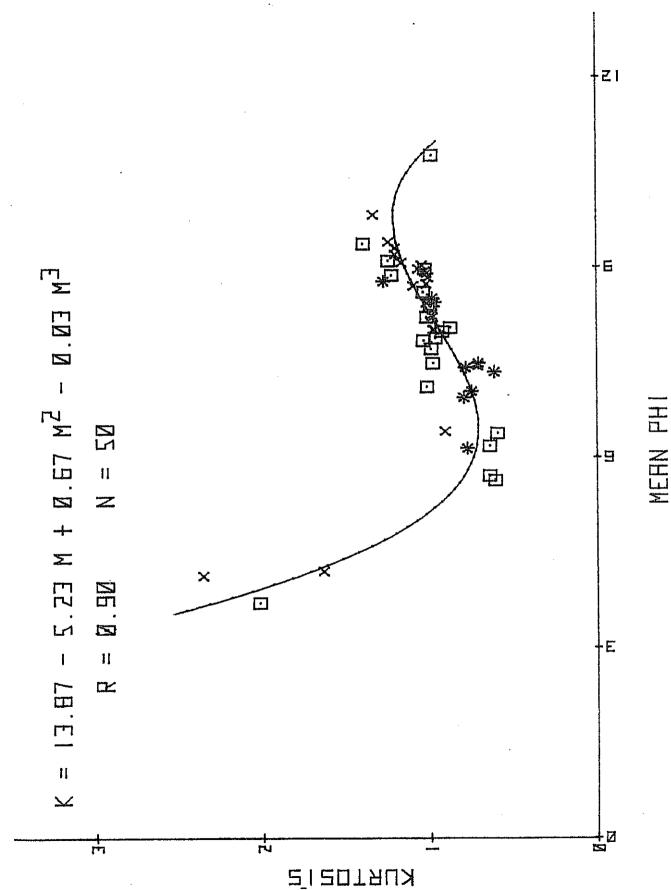


Fig. 29. The relationship between  $X_{\phi}$  and graphic kurtosis of the particle size distributions of Lake Winnipeg off-shore sediments.  $X = \text{south basin; } * = \text{narrows; } \square = \text{north basin.}$ 

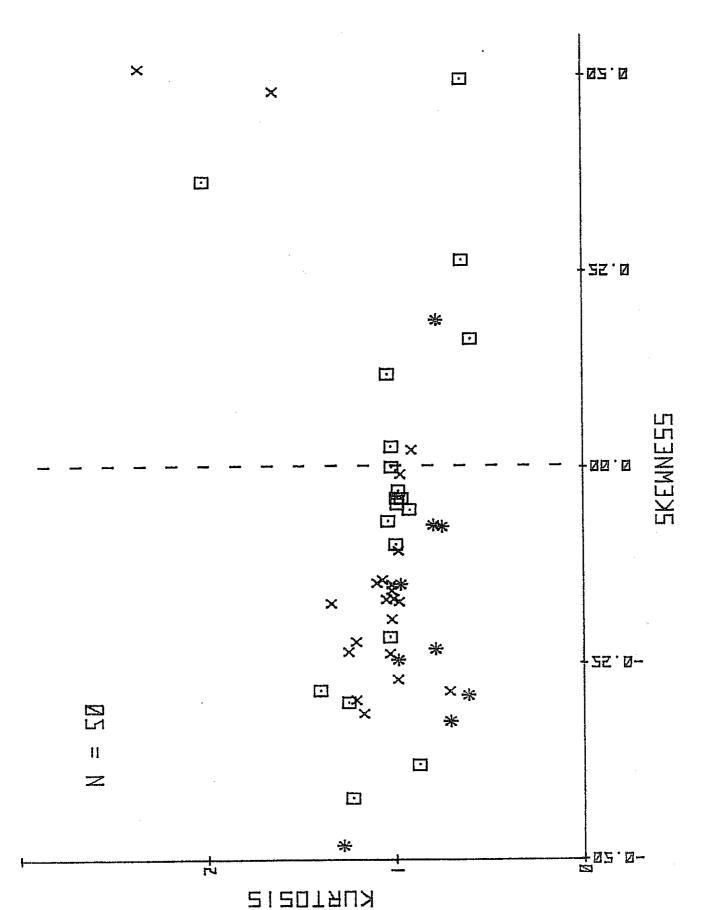


Fig. 30. The relationship of graphic skewness to kurtosis for the particle size distributions of Lake Winnipeg off-shore sediments. X =south basin; \* = narrows;  $\square$  = north basin.

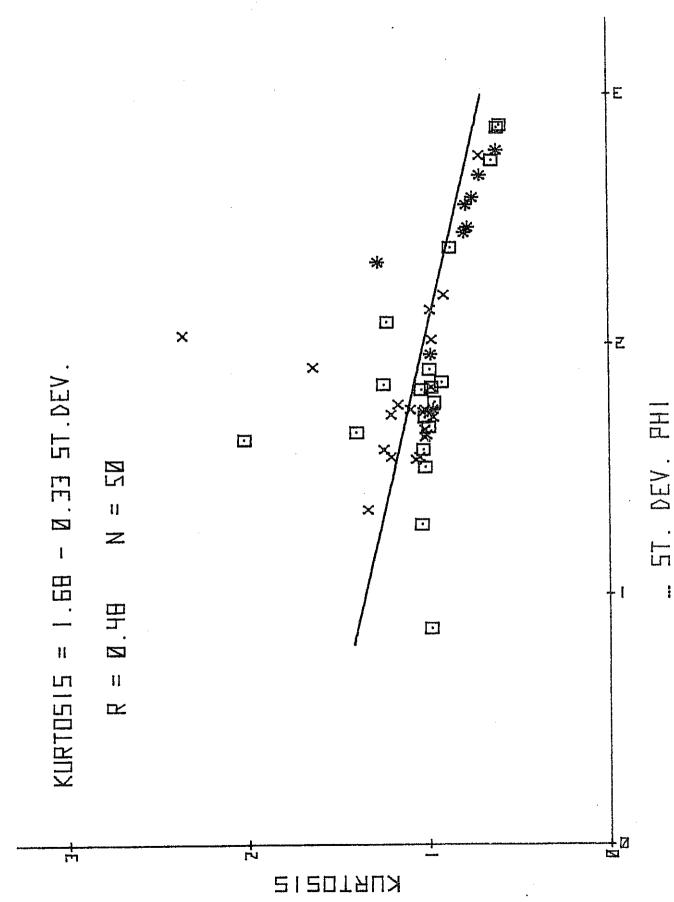


Fig. 31. The relationship of graphic kurtosis to standard deviation of  $\bar{X}_{\varphi}$  for the particle size distributions of Lake Winnipeg off-shore sediments. X = south basin; \* = narrows;  $\square$  = north basin.