

Factors influencing fluffy layer suspended matter (FLSM) properties in the Odra River – Pomeranian Bay – Arkona Deep System (Baltic Sea) as derived by principal components analysis (PCA), and cluster analysis (CA)

J. Pempkowiak¹, J. Beldowski¹, K. Pazdro¹, A. Staniszewski¹, A. Zaborska¹, T. Leipe², and K. Emeis³

¹Institute of Oceanology, ul. Powstanców Warszawy 55, 81-712 Sopot, Poland

²Institute of Baltic Sea Research, Seestrasse 15, 18119 Rostock-Warnemuende, Germany

³Institute of Biogeochemistry and Marine Chemistry, Hamburg University, Bundesstrasse 55, 20146 Hamburg, Germany

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Abstract. Factors conditioning formation and properties of suspended matter resting on the sea floor (Fluffy Layer Suspended Matter – FLSM) in the Odra river mouth – Arkona Deep system (southern Baltic Sea) were investigated.

Thirty FLSM samples were collected from four sampling stations, during nine cruises, in the period 1996–1998. Twenty six chemical properties of the fluffy material were measured (organic matter-total, humic substances, a variety of fatty acids fractions, P, N, $\delta^{13}\text{C}$, $\delta^{15}\text{N}$; Li; heavy metals-Co, Cd, Pb, Ni, Zn, Fe, Al, Mn, Cu, Cr). The so obtained data set was subjected to statistical evaluation.

Comparison of mean values of the measured properties led to conclusion that both seasonal and spatial differences of the fluffy material collected at the stations occurred. Application of Principal Component Analysis, and Cluster Analysis, to the data set amended with environmental characteristics (depth, salinity, chlorophyll *a*, distance from the river mouth), led to quantification of factors conditioning the FLSM formation. The five most important factors were: contribution of the lithogenic component (responsible for 25% of the data set variability), time dependent factors (including primary productivity, mass exchange with fine sediment fraction, atmospheric deposition, contribution of material originating from abrasion-altogether 21%), contribution of fresh autochthonous organic matter (9%), influence of microbial activity (8%), seasonality (8%).

1 Introduction

Material transport in the marine environment is important as regards a number of problems, ranging from elemental biogeochemical cycles to the exposure of biota to pollutants. An important aspect in this respect is factors influencing properties of suspended matter in the coastal environment. The dispersal of chemical species in the water column is determined by a variety of processes such as coastal circulation, particle absorption and adsorption, vertical and horizontal scavenging (Duursma and Carroll, 1996). The contribution of the processes to particulate matter formation varies seasonally, since the climate modifies both hydrological factors and biological activity (Alldredge and Gotschalk, 1990; Mayer-Reil, 1994; Gustafsson et al., 1998; Lund-Hansen et al., 1999; Mayer, 1999; Leipe et al., 2000; Löffler et al., 2000; Emeis et al., 2002; Martino et al., 2002; Pempkowiak et al., 2002).

An important feature of the phenomenon is suspended matter formation and transport in the near-bottom water layer. As depicted in Fig. 1, particulate matter there is formed from suspended matter entering the coastal environment with river run-off, produced by abrasion of the shore and shallow sea bottom, primary production, atmospheric deposition, autogenic mineral formation, and direct anthropogenic discharges (Jago and Jones, 1998; Jones et al., 1998; Bianchi et al., 2002; Pempkowiak et al., 2002). En route from the high-energy, shallow, coastal environment to the low-energy, deep, depositional basins the matter is a subject to later re-suspension, migration driven by near-bottom currents, and sedimentation (Emeis et al., 2002). Wind-induced currents (Lund-Hansen et al., 1997) and aggregation of suspended matter particles (Thomsen and Ritzrau, 1996) are thought to

Correspondence to: J. Pempkowiak
(pempa@iopan.gda.pl)

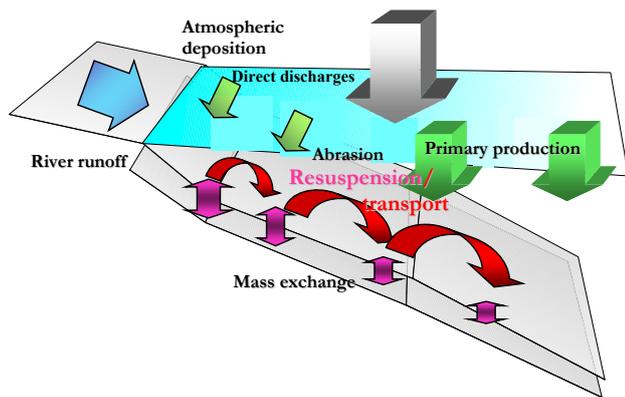


Fig. 1. A schematic indicating sources and fate of FLSM in the near shore environment.

be major factors influencing near bottom transport. However, in periods of calm weather, suspended-matter sedimentation leads to the formation of a high-density fluffy suspended matter (FLSM) layer resting on the sea bottom. Once the FLSM is at rest, both physical and biological processes contribute to the incorporation of the suspended matter into the sediments (Gerino, 1990; Wheatcroft, 1992). The same factors cause fine particles to be released from the sediment and, at least temporarily, incorporated into the suspended matter layer (Pempkowiak et al., 2002). As a consequence, components of FLSM originate from a number of sources, while a number of processes acts to integrate the components into a homogenous material eventually deposited in the low energy depositional areas.

Although individual processes contributing to the FLSM formation and properties are obvious, little is known on their relative importance in an actual coastal environment. This is, at least partly, due to high variability of the coastal environment. Thus a prolonged study, in a course of which a transect from a shallow, highly dynamic near shore environment, to a low energy, depositional basin, was probed repeatedly for FLSM, was carried out, in order to overcome the environmental variability (Emeis et al., 2002; Christiansen et al., 2002).

In this paper we report results of this study carried out in the Odra River – Pomeranian Bay – Arkona Deep System (southern Baltic). Samples of FLSM were collected in the course of a two years long program conducted in the frame of the Baltic Sea System Study (Emeis et al., 2002; Christiansen et al., 2002). Composition and properties of the samples were then measured, and have been a subject of several publications (Emeis et al., 2002; Pempkowiak et al., 2002; Pazdro et al., 2001; Witt et al., 2001; Löffler et al., 2000; Leipe et al., 2000). However, due to the diversified nature of the collected material, no factors responsible for the matter properties were evident and have been named yet. Therefore statistical evaluation, by means of multivariate statistical

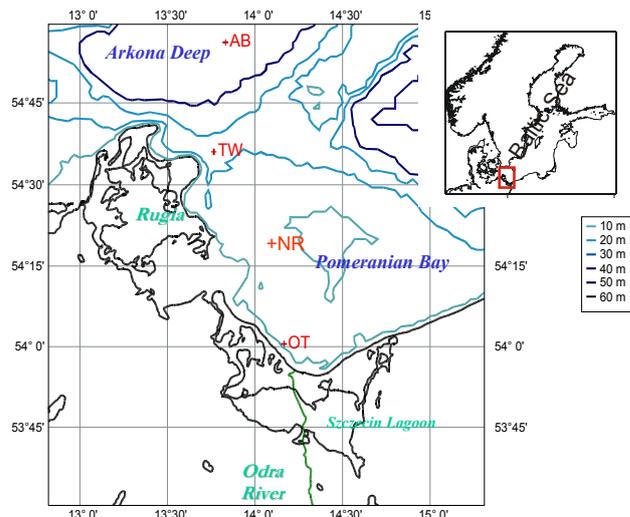


Fig. 2. Distribution of sampling stations in the study area (OT – Odas Tonne, NR – Nordperd Rinne, TW – Tromper Wiek, AB – Arkona Becken).

Table 1. Geographical positions, water depths, and average near-bottom water properties at the sampling stations.

Station	Latitude (N)	Longitude (E)	Water depth (m)	Oxygen ($\text{ml}\times\text{l}^{-1}$)	Salinity (PSU)
OT	54°05′	14°10′	16	5.8	7.5
NR	54°22′	13°52′	20	6.9	7.7
TW	54°36′	13°46′	26	5.9	9.1
AB	54°56′	13°50′	47	2.6	16.4

methods, has been carried out in order to find and quantify the most important factors conditioning the FLSM formation. Both cluster and principal components analyses were used. The results, presented in this publication, point out at, and quantify the contribution of lithogenic material, fine sediment fraction, and freshly produced organic matter; all modified by bacterial activity, as factors actually responsible for FLSM properties.

2 The study area

The Pomeranian Bay – Arkona Deep system extends from the Odra River estuary to the Arkona Basin (Fig. 2). Four sampling stations – Odas Tonne (OT), Nordperd Rinne (NR), Tromper Wiek (TW), and Arkona Becken (AB) were established there, along a transect with the water depth increasing from 16 m at OT to 47 m at AB. The geographical co-ordinates and characteristics of the sampling stations are given in Table 1.

Table 2. Expeditions, station coverage, and the collected samples numbers.

Expedition	Period	Odas Tonne (OT)	Norperd Rinne (NR)	Tromper Wiek (TW)	Arkona Becken (AB)
1	1–11 October 1996	P21,P22	P14,P15	P6,P7	P1
2	14–19 March 1997	P23	P6	P8	P2
3	10–15 June 1997	P24		P9	P3
4	18–22 August 1997	P25			P4
5	12–17 October 1997	P26	P17	P10	
6	1–9 December 1997	P27	P18	P11	
7	16–22 March 1998	P28			
8	22–27 June 1998	P29	P19	P12	
9	8–13 December 1998	P30	P20	P13	P5

The hydrography of this area is influenced by the Odra with its annual water run-off of about 17 km^3 , and by the weather conditions, which influence the water circulation. Sediments in the Pomeranian Bay are temporarily covered by a thin fluffy layer of suspended matter (FLSM). On a weight basis the quantity of FLSM is estimated to range from 2 to 4 mg/cm^3 . The layer is 1 to 3 cm thick. The sediments are composed of a well sorted fine sand (0.1 to 0.25 mm) grading to muddy sand below 20 m water depth and to mud in the depositional basin of the Arkona Deep.

The winds are the main factor governing the currents in the Pomeranian Bay (Löffler et al., 2000). Density-driven currents and tides are of minor importance. Easterly winds dominate between February and May. At this time surface water moves to the north while the near-bottom water layer is transported to the east. Westerly winds are dominant between June and September, while westerly to south-westerly winds generally occur between October and January. During these periods the surface water is transported eastwards along the coast. This surface current is compensated by a near-bottom current, that flows towards the Arkona Deep along the Island of Rugen (Siegel et al., 1999). The sampling stations are situated along a transect following the main transport route of the near-bottom water layer and the suspended matter it carries (Leipe et al., 2000). Although the grain size at the sediment surface was different at the sampling stations (Lund-Hansen et al., 1999), the critical shear stress at all stations was surprisingly uniform, corresponding to flow velocities of about 4 cm s^{-1} . This low value is due to the presence of an organic-rich layer resting on top of the actual sediment surface – FLSM. A flow velocity of 4.5 cm s^{-1} is required to transport the FLSM (Leipe et al., 2000; Christiansen et al., 2002). The Arkona Deep acts as a final deposition area for the material, with an average mass accumulation rate from 1.0 to $1.8 \text{ kg (dry mass) m}^{-2} \text{ year}^{-1}$ (Widrowski and Pempkowiak, 1986; Pempkowiak, 1991; Emeis et al., 2002).

In the period from October 1996 to December 1998 nine expeditions were carried out to the study area. Station coverage is given in Table 2.

3 Sources and composition of FLSM in the study area

Altogether, the Odra discharges about 425 000 t (dry matter)/year of suspended solids. The material is rich in organic matter (15.3% particulate organic carbon) and heavy metals (Pb – $200 \mu\text{g g}^{-1}$, Zn – $1700 \mu\text{g g}^{-1}$, Cu – $120 \mu\text{g g}^{-1}$, Cd – $9 \mu\text{g g}^{-1}$, Hg – $2.5 \mu\text{g g}^{-1}$) (Emeis et al., 2002). In comparison to pelagic suspended matter (PSM), FLSM is rich in hydrated three-layer and mixed-layer silicates such as illite and smectite (Leipe et al., 2000). This is due to their ability to enhance aggregation and sedimentation of suspended matter particles (Leipe et al., 2000). The organic matter content in FLSM is much smaller than in the nepheloid layer suspended matter (NLSM), indicating that organic matter may be undergoing mineralisation in the course of sedimentation. Another possibility is the exchange of particles with surface sediments (Pempkowiak et al., 2002). This conclusion is supported by the fact that the organic matter content at the deepest station (AB – 47 m) is smaller than at the most shallow one (OT – 16 m). Fe-oxihydroxides are present in FLSM: the content of these oxides decreases from about 10% at OT to 5% at TW and 3% at AB. Mn-oxihydroxides are present only at OT (2%) (Leipe et al., 2000). This is most likely due to the anoxic conditions in the uppermost sediments of deeper stations, causing reduction of Fe(III) to Fe(II) and of Mn(IV) to Mn(II). The reduced species migrate to the overlying water, where they are oxidised and precipitated (Pempkowiak et al., 2000).

4 Experimental

The fluffy layer covering the sediment surface was collected by scuba divers, or by a remotely operated vehicle (ROV). The material (FLSM) was siphoned from the sediment surface by an on-board pump and transferred into 100 l containers.

The particulate matter was separated from the water by filtration through pre-weighed glass-fibre filters and freeze dried before further analyses. A set of 26 properties of

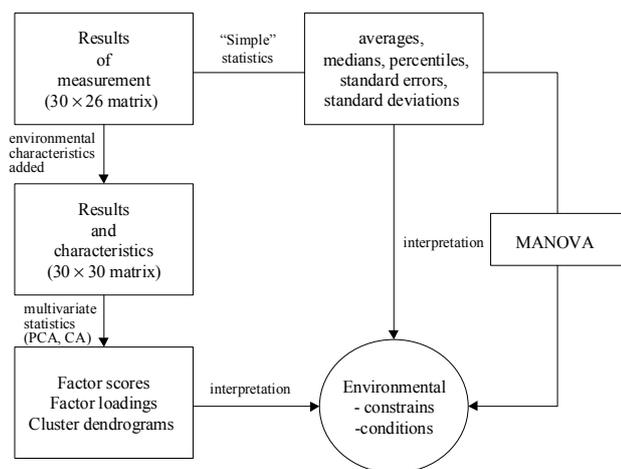


Fig. 3. A schematic of the approach to the statistical evaluation of the data set.

the so obtained material were analysed (total organic nitrogen – N_TOT, total phosphorous – P_PPM, total carbon – TC, loss on ignition – LOI, total organic carbon – C_ORG, total inorganic carbon – CaCO₃, C¹³/C¹² isotopes ratio – δ¹³C, N¹⁴/N¹⁵ isotopes ratio – δ¹⁵N, total free fatty acids – TFAF, the sum of saturated acids – SATURATE, the sum of branched fatty acids – BRANCHED, the sum of monounsaturated fatty acids MONOUNSA, the sum of polyunsaturated fatty acids - POLYUNSA, humic substances – HS, humic acids – HA, fulvic acids – FA, total metals: Zn, Ni, Cu, Pb, Al, Mn, Fe, Co, Li, Cd, Cr). The following procedures were used. Total organic carbon (POC, δ¹³C) and total organic nitrogen (N_TOT, δ¹⁵N) were measured in a Fission Instruments Hd8 CHN Analyser, connected to an isotope ratio mass-spectrometer (Finnigan MAT Delta S). Pure N₂ and CO₂ were used as reference gases. Air nitrogen and NBS-18 reference carbonate were used for standardisation.

Heavy metals (Cd, Pb, Zn, Cu, Ni, Co, Cr, Fe, Mn, Al) were analysed in a model Video 11E atomic absorption spectrometer after 500 mg (dry weight) subsample was wet digested (HF:HNO₃:HClO₄=2:2:1), excess acids were evaporated, and the dry residue dissolved in 10 cm³ 0.1 mol/dm³ HNO₃. Electrothermal (Co, Cd) and flame (other metals) atomizations were applied. Quality assurance of the analyses was assessed by incorporating into the analysis an ICES/SCOR certified, Baltic Sea bottom sediment sample, and ICES marine sediment sample. Accuracy as characterized by recovery was in the range from 83% (Fe) to 108% (Cd), while precision, given by relative standard deviation, was in the range from 5% (Fe) to 13% (Pb).

The samples were analysed for total carbon (TC) by dry combustion and for total inorganic carbon (CaCO₃) by 50% phosphoric acid treatment. An IR detector (Eltra Metotyl CS 1000S C/S analyser) was used to determine evolving CO₂.

Concentration of Li and P were analysed by means of automated electron probe x-ray micro analysis (EPXHA; Leipe et al., 2000).

For fatty acids analyses, a 1–2 g subsample of the freeze-dried material was extracted three times with 2:1 chloroform methanol mixture. The fatty acids contained in the combined extracts are referred to as free fatty acids (TFAF). The solid residue was then saponified with 0.5 mol/dm³ KOH in a 95:5 v/v methanol:water solution and filtered. The acids determined in the resulting filtrate are referred to as bound fatty acids. After separation from other lipid components the fatty acids were derivatized into coumaryl esters and analysed quantitatively using high performance liquid chromatography in a reversed phase column, a gradient methanol:water mobile phase and a fluorescence detector were used. Details of the procedure were described earlier (Pazdro and Falkowski, 1994). Individual fatty acids were quantified using calibration curves based on actual standards (Sigma), and the above described derivatization procedure. The precision of the results, expressed as a relative standard deviation, varied from 1.4 to 20% for individual compounds and from 0.4 to 16% for the total fatty acid content. Concentrations of fatty acids within the following groups were then added and are referred to as total saturated fatty acids – SATURATE (straight chain saturated fatty acids with carbon atoms in the range from 8 to 24 (C8:0 to C24:0), total monounsaturated fatty acids (C14:1 to C22:1) – MONOUNSA, total polyunsaturated fatty acids (C18:2 to C22:6) – POLYUNSA, and branched fatty acids (C_{ai,i}15:0 and C_{ai,i}17:0) – BRANCHED.

Humic substances were extracted repeatedly (five fold) from 2 g of the freeze-dried material by means of a standard alkaline (0.1 mol/dm³ NaOH) extraction. The combined extract was acidified to pH 2.0 with 3 mol/dm³ HCl. Precipitated humic acids (HA) were separated (centrifugation), purified and freeze-dried, while dissolved fulvic acids (FA) were sorbed on Amberlite XAD-2 resin, extracted with 0.1 mol/dm³ NH₄OH, freeze dried and weighted. Details of the procedure are described elsewhere (Pempkowiak, 1989).

5 Statistics

Two approaches have been taken in order to evaluate statistically and characterize the data set originating from the measurements. The strategy employed is depicted in Fig. 3. Results of the measurements (30 samples × 26 properties measured) were a subject to simple statistics. The obtained values were then used to assess spatial and temporal changes of FLSM properties. The average values were also subjected to the MANOVA in order to assess the significance of differences.

The multivariate statistics were applied as another approach. The principal aim of multivariate analysis is to explain variation in a multivariate data set by as few “factors” as

possible, and to detect “hidden” multivariate data structures. This is carried out by developing a set of new “dimensions” – factors, which explain variations in the data matrix (principal components – PCs, in the case of Principal Components Analysis- PCA).

Principal component analysis (PCA), also known as empirical orthogonal function analysis, is an often adopted approach in geochemistry related studies since the pioneering work in this field by Wiwant et al. (1975). PCA objectively extracts the optimal linear lower-dimensional structure from a multivariate dataset. Feature extraction, or pattern recognition, is probably the most appealing of the many purposes for which PCA can be used. The attempt is then made to associate PCA modes with actual physical modes in the dataset. Whereas such an attempt is usually successful in case of few first modes (that is, these that explain most variance in the total dataset), the spatial orthogonality of PCA modes often causes higher-numbered modes to bear predictable geometric relationships to the first mode rather than to represent independent physical modes (Meglen, 1992; Ruessink et al., 2004).

PCA is commonly applied to geochemical data sets which consist of columns – representing sample properties and rows representing individual samples. As result of PCA analysis a set of PCs is calculated equal in number to the number of properties (columns). The analysis is successful if just few PCs explain large proportion of the data set variation. Successful PCA implies also that significance (meaning) of the factors is found. This might, and often do, cause problems. Typically correlation between factors and properties (PCs loadings), and numerical values of factors for samples (PCs scores) are used to explain actual meaning of PCs. After analysis, rotation is applied in order to optimize factor loadings (De Lazzari et al., 2004; Liu et al., 2003; Reiman et al., 2002; Smoleński et al., 2002; Persson and Wedborg, 2001; Podani, 2000; Brown, 1998; Shine et al., 1995; Szefer et al., 1995; Meglen, 1992, to name just a few).

Before applying PCA, the data set (26 properties by 30 samples) was amended with environmental characteristics of the sampling stations and seasons (distance from the Odra river mouth, water depth, water temperature, water chl *a* concentration at the sampling stations). The so obtained data table comprised 30 rows (cases or samples) and 30 columns (26 properties measured +4 “properties” characterizing hydrography of sampling stations and seasonality of expeditions). Then the 30×30 data matrix was subjected to analyses of both clusters (CA) and principal components (PCA).

To probe agglomeration of data, cluster analysis was used. City block distances and squared ordering were used throughout the CA of the data matrix. The dimensionality of the data matrix was reduced and meaning assessed using principal components scores (PCS), and principal components loadings (PCL). Only PCL with eigenvalues exceeding 1 were analysed. Varimax rotation was used in PCA in this study.

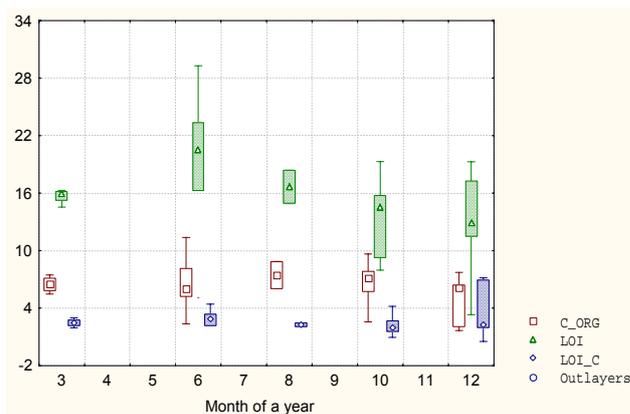


Fig. 4. Seasonal changes of organic carbon (C_ORG), loss on ignition (LOI) and ratio of LOI to C_ORG (LOI_C); C_ORG and LOI in %, LOI_C – dimensionless.

Before applying PCA the data matrix was standardized using the standard deviation approach. This is based on replacing actual values (x_i) with standardized ones (x_{is}). Standardization is meant to eliminate the influence of both different units, and different range of the investigated properties.

$$x_{is} = \frac{x_{ij} - \bar{x}_i}{s_i}$$

where $s_i = \sqrt{\frac{\sum (x_{ij} - \bar{x}_i)^2}{n-1}}$; n – number of samples analysed.

Standardization by maximum values was carried out throughout CA.

$$x_{is'} = x_{ij} / \max_j \{x_{ij}\}$$

In this approach the value of 1 is assigned to the maximum value of a given property while the range of the standardized values is $<0,1>$.

The statistical software package Statistica 6.0 for Windows was utilized.

6 Results and discussion

6.1 Spatial and temporal distribution pattern

To characterize concentrations and properties of organic matter in the samples the following properties of FLSM were used: loss on ignition (LOI), organic carbon (C_ORG), concentrations of humic acids (HA), fulvic acids (FA), and humic substances (HS=HA+FA), total nitrogen (N), total phosphorus (P), a number of fatty acids fractions (saturated, monounsaturated, polyunsaturated, branched). Besides actual concentrations, values standardized to LOI or/and C_ORG were used when appropriate.

In Fig. 4 seasonal changes of C_ORG, LOI and LOI/C are presented. Organic carbon concentration in FLSM decreases

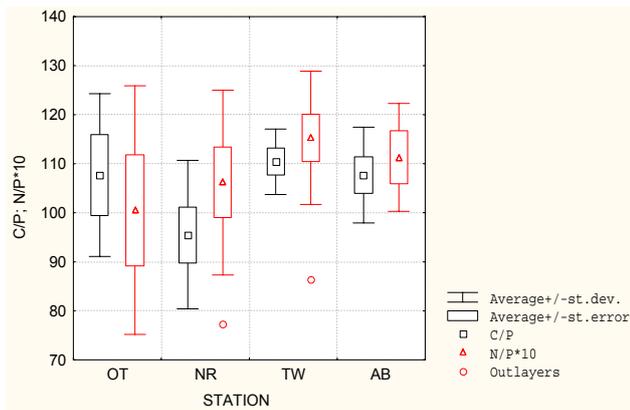


Fig. 5. Spatial distributions of C/P and C/N molar ratios in FLSM.

from 8% in summer to 6% in winter. However, organic carbon contribution to organic matter decreases from some 60% in summer, to 35% in winter. This unexpected feature of fluffy material might be attributed to rapid mineralization of labile organic matter produced in the course of the spring primary production in elevated summer temperatures, while organic matter produced in autumn, after sedimentation, is preserved in the near bottom layer. However, high organic matter concentrations in winter may also suggest participation of other processes, for example contributions of land derived organic matter (15% organic carbon in the Odra river suspended matter was reported), and sedimentary organic matter (Bale and Morris, 1998).

Molar ratios of biogenic elements (C, N, P) in FLSM are smaller than the Redfield ratios (Laima et al., 1999). Smaller ratios of both C/P and N/P in winter, indicate preferential mineralization of organic nitrogen and phosphorus over organic carbon. Combined with the temporal dynamic of the C/LOI ratios, the feature suggests relatively high biochemical stability of organic compounds abundant with carbon, while nitrogen and phosphorous abundant organic compounds are easily biodegraded. The spatial distributions of C/P and N/P, presented in Fig. 5, indicate smaller ratios in the samples collected at the nearshore station (OT and NR). This could be, most likely, attributed to the contribution of land derived organic matter. Bordovsky (1965) indicated that marine autochthonous organic matter is characterized by C/N ratios smaller than 10, while allochthonous organic matter—with values larger than 10. The conclusion is supported by distribution of $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ in the samples, and results of earlier studies on stable isotopes ratios in the area, reported by Voss and Stock (1997). Distributions of $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ clearly indicate contribution of land derived organic matter exceeding 90% at the OT station, in comparison with just above 5% contribution in the AB station, although distribution of lignin oxidation products suggests less steep gradient (Miltner and Emeis, 2001). Organic matter discharged to the sea with river run-off is usually depleted with carbon and nitrogen isotopes

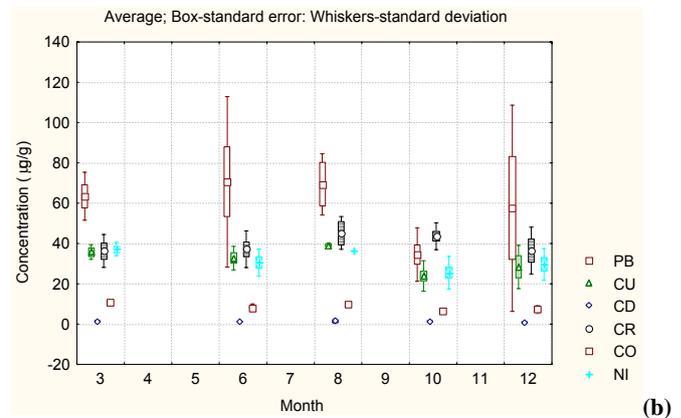
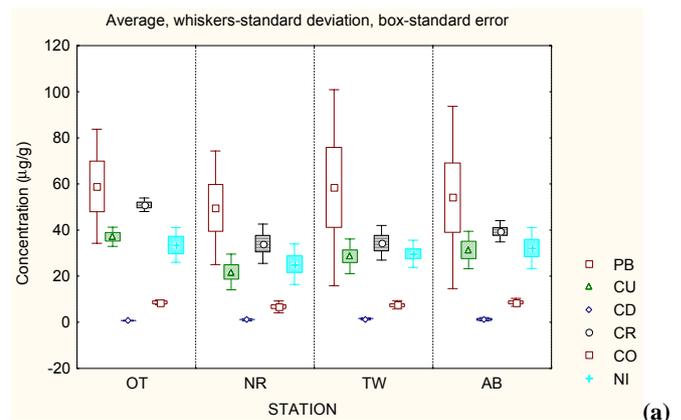


Fig. 6. Variations of heavy metals concentrations (mg/g dry matter) in FLSM (a) spatial, and (b) seasonal.

of larger molecular weight (Voss and Stock, 1997), and with nitrogen (Bordovsky, 1965). Moreover, results of this study show larger proportions of humic substances (humic acids + fulvic acids) in the fluffy material from the nearshore stations. This may be interpreted as resulting from contribution of relatively old, biochemically resistant organic matter derived from land.

On the other hand larger concentrations of labile fractions of organic matter (monounsaturated and polyunsaturated fatty acids) in the summer months, indicate contribution of relatively fresh organic matter (Wakeham et al., 1997; Pazdro et al., 2001). The larger concentrations of saturated fatty acids in winter months result from relative stability of saturated as compared to the unsaturated acids (Sun et al., 1997; Laima et al., 1999). This feature is consistent with discussed reasons for dynamic behaviour of the organic FLSM component, and its elemental composition.

Diagrams showing heavy metals distribution are presented in Fig. 6. Substantial spatial (Fig. 6a) and temporal (Fig. 6b) variations of heavy metals were found in the collected samples of FLSM. This, most likely, reflects diversified origin of fluffy material (river run-off, abrasion of shores and sea

bottom, biogenic input, and red-ox phenomena at the sediment/water interface). After normalizing concentrations to aluminium a pattern governed by both, seasonality and distance from the Odra river mouth, emerges. FLSM is enriched with biogenic metals (Co, Cd, Cu) in summer, while concentrations of these metals in winter are much smaller (Fig. 7a). This was already reported for cadmium in FLSM of the area by Staniszewski et al. (2001). The differences evaluated using the MANOVA proved statistically significant. The explanation that algae (primary production) contribute biogenic heavy metals to FLSM is also substantiated by depletion of dissolved cadmium in pelagic suspended matter, during summer (Pempkowiak et al., 2002), and enrichment of suspended cadmium in the Pomeranian Bay during algal blooms (Pohl et al., 1998). Some metals do not show clear seasonal pattern, although they are regarded as biogenic. This concerns elements characterized by large concentrations in FLSM, hardly affected by incorporation of heavy metals to FLSM in the course of primary productivity. Unusual behaviour of Mn might be attributed to the red-ox phenomena (Heiser et al., 2001; Dehairs et al., 1989). Leipe et al. (2000) indicated that FLSM in the Pomeranian Bay is enriched with hydrated three-layer and mixed-layer silicates such as illite, illite/mixed layers, and smectite. These minerals possess a strong affinity to organic matter, and are readily colonized by microorganisms (Mayer, 1999). This supports conclusion on strong influence of temperature dependent biological processes on concentrations of selected heavy metals in FLSM. No easily recognizable pattern has been found for lead and chromium. Both the metals, attributed to the “scavenged” group, show high concentrations dynamics, reflecting general patchiness, caused by periodical supply of suspended material from the Szczecin Lagoon in this highly dynamic, near shore area of the Pomeranian Bay (Emeis et al., 2002), and both natural and anthropogenic sources of lead (Hinrichs et al., 2002).

A common feature of all metals distributions is a general tendency for decreasing concentrations in the offshore direction (Fig. 7b). This feature is caused by high concentrations of heavy metals in suspended matter supplied to the Pomeranian Bay with the Odra run-off. For example concentration of Cd in the Szczecin Lagoon suspended matter is as high as $5 \mu\text{g/g}$ (Leipe et al., 2000). On entering the marine environment the heavy metals abundant, land derived, material is “diluted” with material originating from coastal erosion. Similar distributions are observed in other estuaries (Belzunce et al., 2001; Borg and Jonsson, 1996). Relatively large concentrations of heavy metals are observed in material collected from the Arkona Deep acting as a depositional ground for suspended matter in the region. The increased concentration of heavy metals there indicates that the deep acts as depositional basin for material originating not only from the Pomeranian Bay but also from other sources, in the large catchment area of the basin (Löffler et al., 2000). However, most organic matter comprised in FLSM is au-

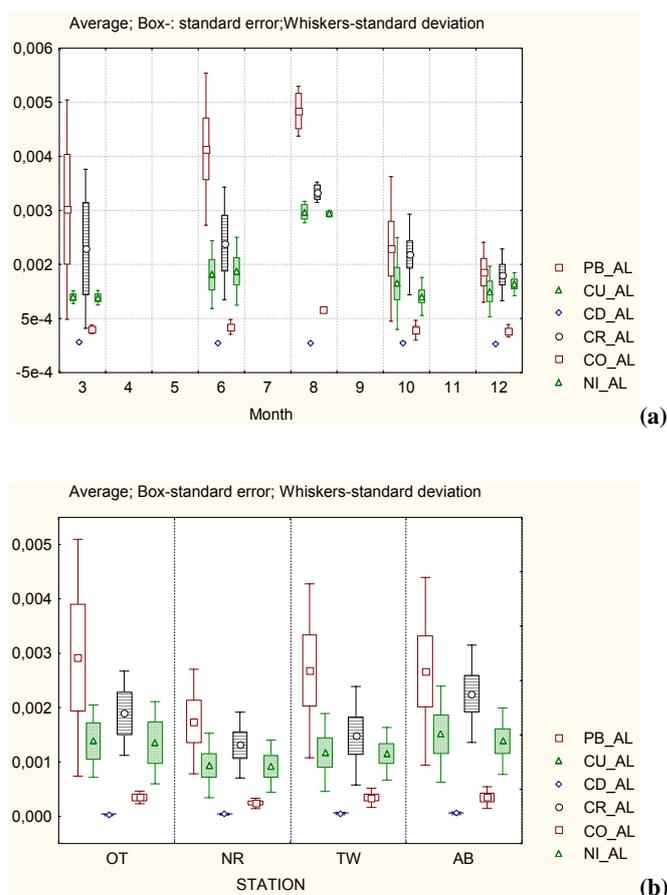


Fig. 7. Changes in heavy metals concentrations in FLSM normalized to Al (a) seasonal, and (b) spatial.

tochthonous, as indicated by stable carbon ($\delta^{13}\text{C}$) and nitrogen ($\delta^{15}\text{N}$) isotopes ratios. This feature was thoroughly discussed and explained by Voss and Stock (1997).

Apart from $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values which clearly differentiate FLSM collected at different stations, chemical properties of FLSM are scattered. Although average values discussed above indicate spatial and temporal pattern, large standard errors underline intensive scatter of results. Properties of FLSM collected at subsequent stations within one expedition, frequently show no pattern at all, or pattern different from that emerging from profiles of average properties. This may originate, as indicated earlier, from diversified sources of suspended matter, supplying the system with suspended matter with varying intensity. Moreover, the shallow, near-shore environment is highly dynamic since wind induced currents cause displacement of FLSM with frequency depending on the water depth. It is estimated that at the OT station events of FLSM resuspension and transport occur twenty to forty times per year, while at the TW station such events occur few times per year at the most. It may also happen that, at least temporarily, direction of FLSM offshore transport is reversed.

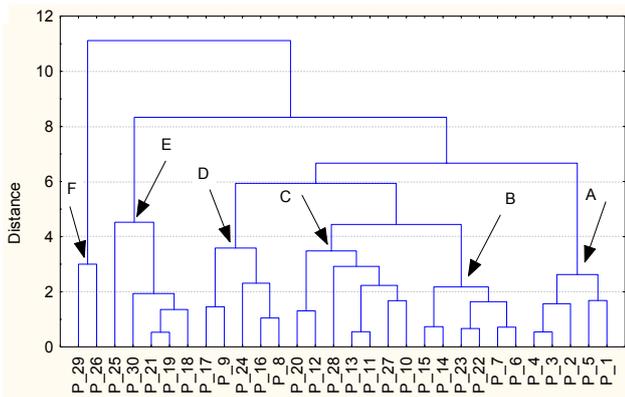


Fig. 8. Dendrogram of the analysed cases (P1–P5: samples collected at AB, P6–P13: samples collected at NR; P14–P20: samples collected at TW; P21–P30 samples collected at OT).

A suitable tool for investigating hidden patterns in data sets of the described sort is multivariate statistical analysis. From among many statistical techniques, principal component analyses (PCA), and cluster analyses (CA) are frequently applied to geochemical and chemical data sets (Manly, 1990; Meglen, 1992; Szefer et al., 1995), and are used in this study to assess factors conditioning FLSM. Meglen (1992) suggests that multivariate analysis begins with the CA, and then is followed and confirmed by a multi-stage PCA.

6.2 Cluster analysis (CA)

A dendrogram showing distribution and association of the analysed samples is presented in Fig. 8. Six major clusters, marked with letters, can be recognized in the dendrogram. Cluster A represents samples collected at the AB station. The samples are clearly separated from other samples collected in the system. This underlines substantial difference between properties of the AB samples and those collected from other stations, closer to the Odra mouth. Both anoxic conditions and prolonged time of the matter residence on the sea floor may be responsible for the phenomenon (Siegel et al., 1999). Moreover, Pempkowiak et al. (2002) indicated substantial mass exchange between FLSM and fine fraction of surface sediments. Substantial contribution of suspended matter from other areas than Pomeranian Bay/Odra River system to FLSM was reported (Emeis et al., 2002), and may contribute to the specific properties leading to the separation of the AB station samples, from others collected in the system. Cluster F, opposite to cluster A is composed of two samples collected at the OT station, the one closest to the Odra River mouth. Both samples were collected following the period of heavy flood in the Odra catchment area and, quite likely, represent allochthonous suspended matter. Cluster B comprises samples collected at stations OT (22, 23), NR (6, 7), and TW (14, 15) in October 1996. Close associ-

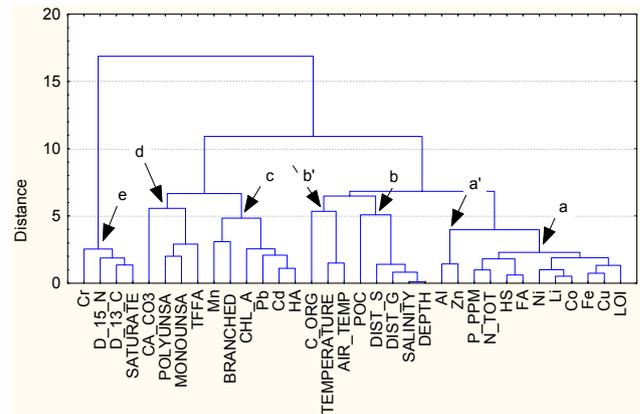


Fig. 9. Dendrogram of the analysed FLSM properties.

ation of samples indicates relative homogeneity of FLSM in the Pomeranian Bay at that sampling time. However, sample 5 collected at the AB station at the same expedition in October 1996 is incorporated in the AB cluster (A), indicating and confirming specific composition of FLSM there.

The other three clusters (C, D, E) comprise samples collected at the OT, NR, and TW stations. It is worth noticing that individual clusters comprise samples differing not only with locations where they were collected, but season as well. It is characteristic that samples from the NR station were collected several months after those from the OT station, while samples from the TW station – still later than those from the NR station (see Table 2 for comparison). This indicates that FLSM retains its properties (a memory effect) when transported from the Odra River mouth (OT station) seaward (NR station, followed by TW station) to a degree enabling time independent clustering. The distribution can be used for assessing time and velocity of migration. In 4 out of 5 instances when the “movement” of samples characterized by similar properties was noticed, the time required for FLSM to move from OT to TW stations is in the range from 4–9 months. The thesis that pattern of FLSM properties can be recognized after several months and after covering some 200 km was also investigated using method developed by Krumpal (1993). The method is based on measuring ratios of heavy metals concentrations (Table 3) and using the values for fingerprinting the samples. Specific values of the ratios can be detected along the transect, corresponding to patterns found in the clusters C, D, and E (Fig. 8).

In Fig. 9, a dendrogram showing distribution and association of the analysed properties is presented. Cluster *a* comprises concentrations of lithogenic elements (Li, Co, Ni, Fe) and properties closely associated with concentration of organic matter (LOI, FA, HS, N, P). Copper characterized by high anthropogenic enrichment factor in the Baltic sediments (Pempkowiak, 1991; Szefer et al., 1995) is also included in the cluster. This might be caused by affinity of the element to organic matter in general, and sedimentary humic

Table 3. Fingerprinting FLSM.

Cruise date	FLSM characteristics							
	OT		NR		TW		AB	
	Cu/Co	Fe/Co	Cu/Co	Fe/Co	Cu/Co	Fe/Co	Cu/Co	Fe/Co
October 1996	2.7*	0.20*	3.8	0.31	3.9	0.30	3.8	0.35
March 1997	4.0	0.25	3.3	0.29	2.6	0.20	2.6*	0.20*
August 1997	3.2	0.44	3.9	0.41	3.7	0.31	3.4	0.36
December 1997	5.2	0.38	3.5**	0.54**	4.1	0.34	3.9	0.27
June 1998	4.4	0.39	4.2	0.36	3.3**	0.51**	3.6	0.40
December 1998	4.0	0.30	4.4	0.45	3.6	0.30	3.3**	0.53**

*, ** Specific ratios of metals in samples collected along the Odra mouth – Arkona Deep transect.

substances in particular (Pempkowiak and Szponar, 1995). The lithogenic elements – organic matter cluster (*a*) is associated with cluster *a'* comprising Al – another lithogenic element, and Zn – most often included in the group of “anthropogenic” metals (Szefer et al., 1995). Both the *a* and *a'* clusters are closely associated with a cluster comprising salinity (SALINITY), depth of water at the sampling sites (DEPTH), and their distance from the Odra mouth (DIST). These associations point at the Odra River as the source of material, and the major factor influencing distribution of lithogenic metals in the coastal environment. Cluster *c* comprises two anthropogenic elements Cd, and Pb as well as humic acids – HA (possibly organic complexes with heavy metals), and Chl *a* (indicative of primary production intensity). These are closely associated with concentrations of Mn (indicator of red-ox potential shifts) and branched fatty acids – BRANCHED (indicator of microbial activity). In cluster *d* fatty acids, excluding the saturated ones, are grouped. These are associated with calcium carbonate concentration, indicating contribution of mussels abundant in the Pomeranian Bay: *Mytilus edulis*, *Mya arenaria*, and *Macoma balthica* (Warzocha, 1994) to the pool of fresh organic matter (unsaturated fatty acids are a labile fraction of organic matter – Pazdro et al., 2001). It is quite likely that the mussels contribute to the enrichment of FLSM with Cd and Pb, since *Mytilus edulis* is known to accumulate Cd, while *Mya arenaria* and *Macoma balthica* preferentially accumulate Pb (Szefer, 1987).

Cluster *e* indicates close relation between distribution of stable isotopes in organic matter, and concentration of chromium. Distribution of stable isotopes is closely related to the provenience of organic matter. It is also the property characterized by the smallest standard deviations of average concentrations at subsequent stations, and exhibiting the well developed trend (95% of land derived organic matter at the OT station; 5% of land derived organic matter at the AB station). It may be assumed, therefore, that chromium in FLSM is supplied to the system with the Odra River run-off, and distributed similarly to land derived organic matter (steep gradient of land derived components off-shore). It is interesting

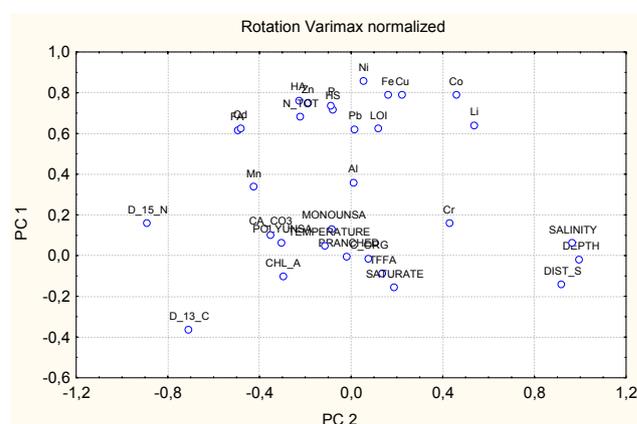


Fig. 10. Scatterplot of the object samples properties loadings in space spanned by axis PC1 and PC2 of the analysed data set.

to see that saturated fatty acids, are also included in the *e* cluster. The feature strongly supports conclusions of Pazdro et al. (2001) who pointed at long chain saturated fatty acids, a stable fraction of organic matter, as closely related to the land derived organic matter.

6.3 Principal components analysis (PCA)

In Table 4 results of PCA are presented. The first five principal components (PC) are responsible for some 74% of the data scatter. The PC6 eigenvalue indicates contribution differing little from contributions of each of the thirty variables included in the data set. Basing on PCs eigenvalues the number of evaluated PCs was limited to the first five. Loadings of the PCs, subjected to further analysis, are presented in Table 5. In Fig. 10 distribution of PC1 and PC2 loadings is presented in the scatterplot of individual variables in space spanned by axis PC1 and PC2.

A number of FLSM properties exhibit PC1 loadings exceeding 0.6. These include Cu, Co, Ni, Fe, LOI, Li, P, N (components of the *a* cluster – Fig. 9), Zn, and Pb. The

Table 4. Results of PCA.

Principal Component	Characteristics			
	Eigenvalue	Explained variability(%)	Cumulative eigenvalue	Cum.explained variability (%)
1	8.53	25.11	8.53	25.11
2	7.22	21.23	15.76	46.35
3	3.67	10.85	19.45	57.21
4	2.88	8.49	22.33	65.70
5	2.73	8.05	25.07	73.75
6	1.67	4.92	26.75	78.68
7	1.37	4.05	28.13	82.73

Table 5. The first five PCs loadings.

Properties	PC loadings (Varimax rotation)				
	PC_1	PC_2	PC_3	PC_4	PC_5
LOI	0.615827	-0.131215	0.135703	-0.116643	-0.158778
HA_%	0.764951	0.244689	0.031128	0.016921	0.063215
FA_%	0.622782	0.521088	0.096235	-0.122844	0.048274
HS_%	0.717535	0.091280	-0.017029	0.200468	-0.341112
TFAF	-0.103623	-0.148583	0.804567	0.298030	0.152843
SATURATE	-0.151261	-0.197577	-0.894538	0.149026	-0.069372
MONOUNSA	0.134285	0.103319	0.802867	-0.324078	0.060624
POLYUNSA	0.056482	0.312121	0.825113	0.093179	0.133130
BRANCHED	0.023038	0.065904	0.227170	-0.770905	0.094131
PB	0.622738	-0.018872	-0.103822	-0.339346	0.230029
CU	0.806217	-0.195065	-0.110383	-0.374885	0.184319
CD	0.610996	0.475611	0.341783	0.007825	0.148894
FE	0.795886	-0.147878	0.073624	-0.285510	-0.050702
CR	0.169184	-0.408616	0.061850	0.203638	0.626316
MN	0.358417	0.465929	-0.277963	-0.379490	-0.297228
CO	0.798616	-0.444187	-0.174920	-0.054096	-0.024167
NI	0.854760	-0.051373	0.072531	0.194649	0.105215
ZN	0.744949	0.187217	0.080519	0.118211	-0.202666
AL	0.331586	-0.051579	0.516195	0.403984	-0.258158
N_TOT %	0.674489	0.228624	0.342183	0.298404	0.214571
CA_CO3 %	0.081751	0.333692	0.252231	0.718889	0.159130
$\delta_{15}\text{N}$	0.151160	0.893782	0.164908	0.043205	0.215380
$\delta_{13}\text{C}$	-0.372765	0.705305	0.265600	0.145884	-0.069663
LI_PPM	0.642370	-0.535120	-0.078994	0.360288	0.150337
P_PPM	0.735087	0.099368	0.251127	0.273612	0.131260
C_ORG	0.689340	0.229371	0.042478	-0.224912	0.138206
Depth	-0.005356	-0.989345	-0.055603	-0.043507	-0.006959
Distance	-0.134794	-0.924796	0.005893	-0.034832	-0.136303
Salinity	0.076838	-0.953152	-0.090243	-0.043946	0.071806
CHL_a	-0.113655	0.286276	0.260015	0.128807	0.736589
Eigenvalue	7.774360	6.036083	3.816995	2.534828	2.349314
Expl.variance	0.2511	0.2123	0.1085	0.0849	0.0805

pattern indicates that PC1 meaning could be attributed to contribution of lithogenic (land) derived material in FLSM. The eigenvalue of 8.53 corresponds to 25.1% of total vari-

ability in the data set. The PC2 (eigenvalue 7.22, variance 21.2%) is characterized by large loadings of stable isotopes ($\delta^{15}\text{N}$, $\delta^{13}\text{C}$), salinity, water depth, and distance from the

Table 6. Summary of PCA.

PC	Explained		Assigned meaning
	Variability %	Cum. Variab. %	
1.	25.1	25.1	Distribution of heavy metals and humic substances. <i>Contribution of lithogenic and land derived material.</i>
2.	21.2	46.3	Distance from the shore, contribution of autochthonous material. <i>Influence of time dependent processes (abrasion, fine sediment fraction, autochthonous organic matter, atmospheric deposition).</i>
3.	10.8	57.2	Distribution of straight chain fatty acids. <i>Contribution of fresh organic matter.</i>
4.	8.5	65.7	Distribution of branched fatty acids. <i>Microbial activity.</i>
5.	8.1	73.8	Chl. <i>a</i> , Temperature. <i>Seasonality of properties.</i>

shore (the Odra river mouth). Salinity and water depth increase with increasing distance from the river mouth, while contribution of land derived organic matter decreases sharply from 95% at the nearshore OT station, to 5% at the offshore AB station (Emeis et al., 2002). Therefore the distance from the Odra River mouth can be assigned as actual meaning of PC2. It is worth stressing that distance from the shore, in this instance, describes collectively a number of time dependent factors decisive for the FLSM properties. These will include contribution of autochthonous organic matter (Voss and Stock, 1997; Emeis et al., 2002), time factor important for the load of fine sediment fraction contributing to FLSM (Pempkowiak et al., 2002), atmospheric input (Löffler et al., 2000; Witt et al., 2001), and both autochthonous organic matter production and biochemical degradation (Pazdro et al., 2001).

This assignment is supported by the PC2 loadings exceeding 0.9 for a set of variables closely attributed to the distance from the shore. It is also strongly supported by distribution of sample scores presented in Fig. 11, in space spanned by axis PC1 and PC2. Samples collected at subsequent stations are separated into clusters according to PC2 scores. It is interesting to notice the strong separation of the Arkona Becken samples, and much weaker separation of samples collected at the other stations. This is in agreement with cluster analyses presented in Fig. 8. No seasonal pattern can be assigned to either PC1 or PC2. Some attention is to be paid to the fact that PC1 scores of samples collected at different stations do not separate them. This originates from, and supports, the meaning assigned to PC1. Material of land origin is, after all, a major component of FLSM collected at all stations (Löffler et al., 2000; Leipe et al., 2000). One exception is organic matter as indicated by stable isotopes ($\delta^{15}\text{N}$, $\delta^{13}\text{C}$), and chromium, most likely, due to the steep gradient, and lack of land associated contents at the offshore stations.

Distribution of straight chain fatty acids sets PC3 (Table 5). Since saturated fatty acids (large positive PC3 load-

**Fig. 11.** Scatterplot of the object samples scores in space spanned by axis PC1 and PC2 of the analysed data set.

ing) are relatively resistant to biodegradation, while mono- and poly-unsaturated ones (large negative PC3 loadings) are considered to be labile (Pazdro et al., 2001; Canuel and Martens, 1996), the PC3 differentiate between fresh and old organic matter. Distribution of branched fatty acids sets PC4. This fraction of fatty acids originate from bacterial activity (Brown et al., 1996; Pazdro et al., 2001). Therefore, it can be safely assumed that PC4 represents the influence of bacterial activity on FLSM properties. Both Al and CaCO_3 concentrations are characterized by substantial PC4 loadings values. Both are opposite to branched fatty acids loading. Both water temperature and Chl *a* show large PC5 loading values. Water temperature is closely related to season as is Chl *a*. Therefore, conclusions drawn from the time related pattern of properties discussed earlier and presented in Fig. 8 and Table 3, combined with the meaning of PC5 indicate seasonality of the properties. In Table 6 summary of PC eigenvalues and actual meaning assigned to subsequent principal components is given.

7 Conclusions

FLSM samples were collected from four stations situated along a main route of currents leading from the mouth of the Odra River to the depositional area at the Arkona Deep in the period December 1996–December 1998.

Twenty six properties of the samples were analysed. These included concentrations of heavy metals (Ni, Co, Fe, Mn, Zn, Cd, Cu, Pb, Cr, Al), organic matter (Loss on ign., C_{org} , N_{tot} , P_{tot} , $\delta^{13}C$, $\delta^{15}N$), organic matter fractions differing with resistance to degradation (humic substances, fatty acids), and shell debris ($CaCO_3$). The so obtained data set, amended with environmental characteristics of the sampling sites, was then subjected to statistical analyses. The FLSM spatial changes caused by the land derived material dilution with the autochthonous one were documented. Seasonal changes, although less visible, were also noticed. Applying multivariate statistical analysis (both PCA and CA) led to dimensionality reduction in the data set. Contribution of lithogenic material (PC1 – 25%), influence of time dependent processes including contributions of autochthonous organic matter, atmospheric input, fine fractions of sediments, biochemical degradation (PC2 – 21%), contribution of fresh organic matter (PC3 – 13%), bacterial activity (PC4 – 8%), and seasonality (PC5 – 8%) were recognized as main factors controlling properties of FLSM in the investigated system. Samples clustering indicated both Arkona Deep and the Odra River mouth as end members of the FLSM in the system. The FLSM properties changed substantially on passage from the high energy shallow environment to the depositional area. On several instances the passage time was evaluated to be in the range from 4 to 9 month.

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