plumose setae (Fig. 1E). The posterior segments are crowded. A few posterior segments are missing in the present specimen.

In all characters, the present specimen fully agrees with the description of the type specimen.

The occurrence of P. johnsoni was first reported by Hartman¹ from the partly enclosed bays of southern California, from the longshore canyons at a depth of 97-230 m and from the southern United States in the littoral region. This was not reported by Day2. However, Hartman3 reported Poecilochaetus from Madras Harbour but the species was not mentioned.

Occurrence of P. johnsoni in the Vasishta Godavari estuary is reported for the first time from the Indian waters. Work on the rich polychaete fauna inhabi-

ting this estuarine system is in progress.

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Observations on Faecal Contamination of Cherai Beach in Kerala

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Faecal pollution of Cherai during premonsoon, monsoon and postmonsoon seasons of 1976 is studied, and the counts of 'indicator bacteria' like E. coli and Streptococcus faecalis occurring in the beach sand and beach waters are presented. Based on FC/FS ratio, the faecal pollution in Cherai beach is primarily of nonhuman source.

FAECAL pollution of any given environment is primarily due to the sewage discharge and faecal matter both of human and nonhuman sources. Coliform and faecal streptococci constitute broadly the 'indicator' group of bacteria. Although several detailed studies on this group are available data on faecal pollution of beaches and sea water are meagre¹⁻⁶. A survey of the indicator species in a beach environment should throw considerable light on the senitary condition and also serve as a guideline for the development of a tourist resort. Such a study has been undertaken on 2 beaches — Cherai and Fort Cochin of Kerala in 1976. The present paper deals with only Cherai, a more or less straight

beach, north of Cochin city. The data are collected during 9 months (April-December 1976).

Sand samples (approximately 15 g each) from a fixed transect along the Cherai beach were collected in small sterile petri dishes, using a sterile spatula. Surf beaten sea water was collected directly in a sterilized glass bottle. Sixteen samples were collected per month at fort-nightly intervals. The samples were brought to the laboratory in iced condition and processed immediately. Sand samples (5 g) were weighed aseptically and mixed thoroughly with 100 ml autoclaved sea water. Sand samples were shaken well in order to release the adhering bacteria and 1 ml of this was plated on sodium desoxy-cholate agar. Water sample (1 ml) was directly plated. The plates were incubated for 24 hr at 37°C. The red colonies which appeared were taken as coliforms and the number was recorded. These red colonies were then streaked on eosine methylene blue agar (EMB agar) and the plates incubated at 45°C for 24 hr. Colonies with metallic sheen were identified as E. coli. When, however, metallic sheen was not developed within 24 hr a further incubation for 24 hr was carried out and the counts recorded as E. coli. These colonies were inoculated in test tubes containing bacto EC medium with inverted Durham tubes and incubated at 45°C for 24-48 hr. Faecal E. coli

produced gas in the Durham tubes.

Detection of faecal streptococci was carried out as follows: processed sand and water samples were

plated on KF agar medium and incubated at 37°C for 48 hr. KF agar medium had the following

composition (%): proteose peptone (No. III), 1;

bacto yeast extract, 1; sodium chloride, 0.5; sodium

glycerophosphate, 1; maltose C.P., 2; lactose, 0.1;

sodium azide, 0.04; sodium carbonate, 0.06; bromocresol purple, 0.0015; and agar, 2. Agar and pro-

teose peptone were added to half of the distilled water and steamed until agar was completely

dissolved. Remaining ingredients were added to

the rest of the water and steamed for 10 min. The

2 mixtures were combined, filtered through cotton and bromocresol purple was added. After the

addition of the indicator the medium was distri-

buted in 100 ml quantities in 100 cc conical flasks

and sterilized at 15 lb pressure for 30 min. Pink or red colonies of faecal streptococci developed

after 48 hr. Parameters of sea water like dissolved

oxygen, salinity, pH and temperature were also

recorded.

Coliforms, E. coli and faecal streptococci occurred in most of the months both in sea water and in sand. From Table 1 it can be seen that coliforms, E. coli, and faecal streptococci were present in most of the months both in sea water and in sand samples, and the counts were higher during monsoon and immediately after monsoon (July-October). The counts were higher in beach sand than in water samples. High counts during monsoon and immediate postmonsoon months may be due to the low salinity and also due to lesser exposure to sunlight and temperature, since these organisms cannot tolerate high salinities even for 24-48 hr. Higher counts in beach sand compared to sea water may be explained by the fact that beach sand offers a

Table 1 — Occurrence of Different Bacteria in Cherai Beach During April-December 1976

(Values are average of 3 observations)

		Sand	counts/g		Water count/ml			
•	Coliforms	E. coli	Faecal streptococci	FC/FS	Coliforms	E. coli	Faecal streptococci	FC/FS
Apr.	7	2	5	0.4*	3	1	2	0.5*
May	11	3	8	0-3*	7	nil	2	
June	17	4	9	0-4*	15	2	3	0.6*
July	17	6	9	0.6*	6	2	1	1*
Aug	19	8	4	2†	11 .	nil	1	
Sept.	19	4	3	1.33†	11	niI	1	_
Oct.	18	6	3	2†	18	3	nil	3†
Nov.	13	. 2	1	1*	10	nil	1	_
Dec	7	1	2	0.5*	16	nil	nil	

^{*}Nonhuman faecal pollution. †Human faecal pollution.

TABLE 2 - AVERAGE HYDROGRAPHICAL PARAMETERS

Month	Temp	. (°C)	Dissolved	Salinity	рΗ
	Water	Sand	oxygen (ml/litre)	%.	
Apr.	30.3	36	5·16	32-43	7-4
May	30.6	37.8	6.61	33	7.3
Tune	30.2	36	5.88	31.8	>7
July	27.5	38.2	6	24.1	>7
Aug.	28.3	37.2	6.15	25•4	7.5
Sept	30-1	40	7.03	30.81	7-5
Oct.	29.5	32-3	6.6	31.09	7.4
Nov.	29.7	31.7	6.73	32.03	7.5
Dec.	30.5	37.6	6-34	28.6	7.4

better substratum for the organism. As the salinity rises making the environment more and more marine the organisms get destroyed resulting in low counts. Gore and Singbal⁴ and Dwivedi et al.⁵ have reported E. coli counts from 2 sandy beaches of Goa; a maximum of 50/g of sand in Siridaon beach which is more estuarine and 10-15/g in Colva, a plain sandy purely marine beach. In the present study, the beach is marine and the count ranges between 7 and 10/g of sand. In this respect it resembles the Colva beach (Goa) revealing a more or less similar bacteriological status. There appears to be no significant effect of chemical parameters except salinity, on counts of the above organisms (Table 2). Since there is no significant variation in the pH values there appears to be no direct relation with bacterial counts.

Examination of faeces from a large number of animal species reveals that with one exception the faecal streptococci (FS) group outnumbers the faecal coliform group (FC). But the FC of human intestines are more abundant than FS. This fact may be utilized to distinguish between human and animal faecal pollution in waters. The available evidence suggests that FC/FS ratio in excess of 2, probably represents primarily human faecal pollution, while ratio of 1 or less may be due to nonhuman sources. Intermediate value cannot be interpreted reliably. In accordance with this FC/FS ratio, the faecal pollution in Cherai beach may be principally of nonhuman sources. FC/FS ratio for beach sand and sea water in various months is incorporated in Table 1. The results indicate that Cherai beach is not free from faecal pollution. Considering the acceptable counts for coliform and S. faecalis given by Brisou⁷, in the beach under study the counts of coliforms ranging between 7 and 19 of sand and 6 and 18/ml in sea water, indicate that they are beyond the limit of acceptance; the faecal streptococci counts, however, do not fall within the acceptable range. In general, the counts of coliforms and S. faecalis both in sea water and in sand suggest that the beach is contaminated to a considerable degree during the monsoon and immediate postmonsoon months. However, the water did not show any macroscopic faecal pollution during the period of investigation.

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