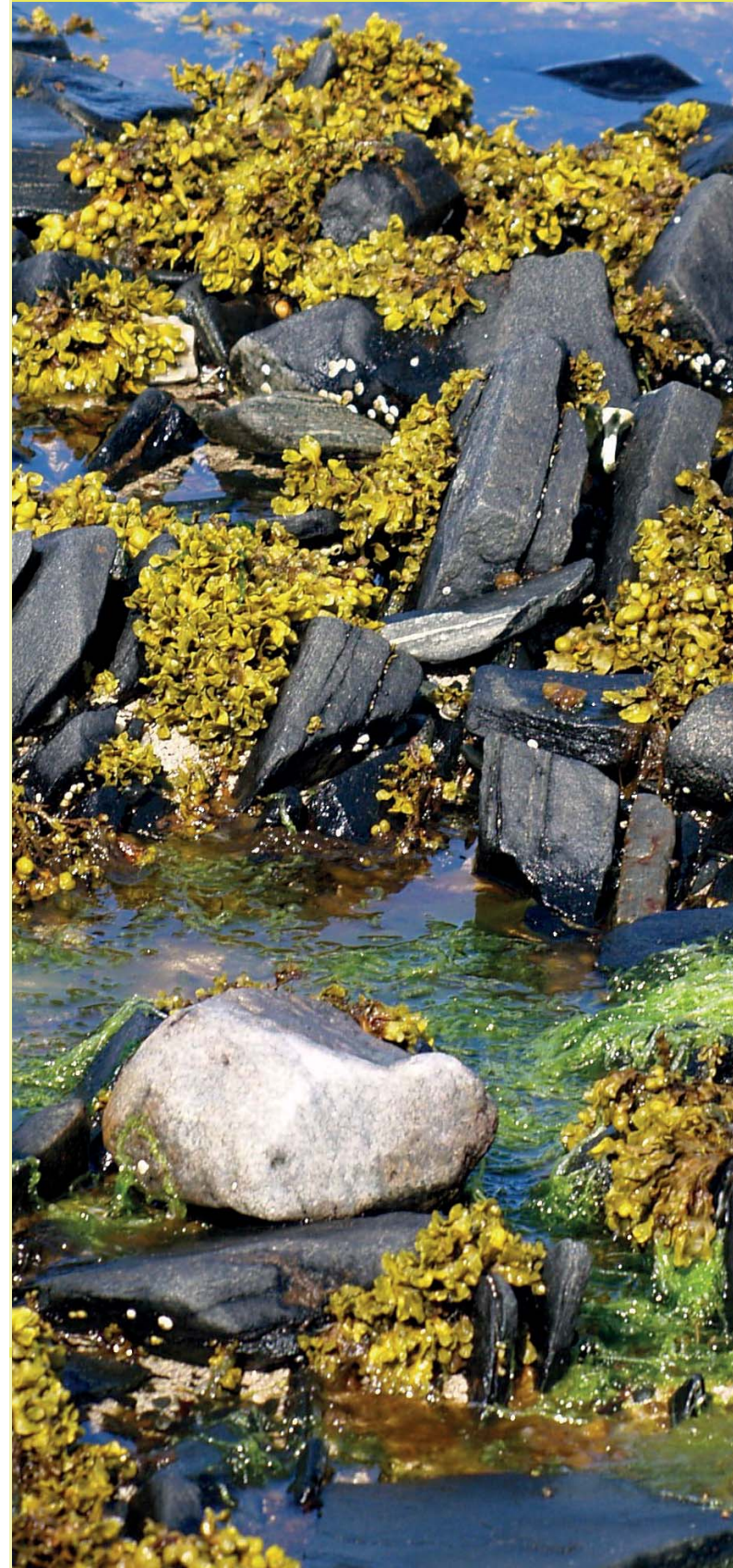




The 12<sup>th</sup> Symposium on Aquatic Microbial Ecology



# SAME12 2011

from August 28<sup>th</sup>  
to September 2<sup>nd</sup>

12<sup>th</sup> Symposium  
on Aquatic Microbial Ecology

Germany  
Rostock–Warnemünde

**MORPHOLOGICAL- VERSUS MOLECULAR-BASED DIVERSITY SURVEYS OF PLANKTONIC MICROBIAL EUKARYOTES: THE CASE OF TINTINNID CILIATES**

Bachy, C.<sup>1</sup>, Lopez-Garcia, P.<sup>1</sup>, Dolan, J.<sup>2</sup>, and Moreira, D.<sup>1</sup>

<sup>1</sup>*Unité d'Ecologie, Systématique et Evolution, Université Paris-Sud, France ,*

<sup>2</sup>*Microbial Ecology, Laboratoire d'Océanographie de Villefranche-sur-Mer, Station Zoologique, France*

During the last decade, the use of culture-independent molecular approaches to describe the communities of microbial eukaryotes present in natural environments has led to the discovery of a huge diversity of these organisms. However, these analyses have rarely taken into account the temporal pattern of variation of diversity and most often lacked of a comparison of morphological- and molecular-based estimates. In this context, we have studied the temporal pattern of genetic diversity of tintinnids (Ciliophora) over a two-year survey in a Mediterranean Sea location (Villefranche-sur-Mer, France). The species-rich order Tintinnida contains freshwater and marine ciliates easily distinguishable based on morphological characters, in particular their conspicuous organic or inorganic tests. This allowed us to couple morphological observations with a double molecular approach (using *single cells and environmental DNA*) to analyse the *SSU-rRNA* and the *ITS coding regions*. Using a fingerprinting technique (DGGE), we detected a strong relationship between the structure of the tintinnid communities and the sampling depth. Despite an extensive work of single-cell isolation, identification, and subsequent *SSU-rRNA* and *ITS* sequencing, the analysis of tintinnid communities by direct PCR amplification and sequencing of rRNA genes from plankton samples revealed a number of phylotypes without any closely related known species. Conversely, several sequences from single-cell analyses were never found in the environmental sequence libraries. Using this well-characterized protist group, we discuss the limitations of morphological- and molecular-based studies to assess the diversity and temporal dynamics of microbial eukaryotic communities.