

ABSTRACT

The anti microbial activities were determined by agar diffusion modified method by using Nutrient agar medium and *Staphylococcus aureus* ATCC 25923, *Pseudomonas aeruginosa* ATCC 27853, *Escherichia coli* ATCC 25922, *Bacillus subtilis*, *Salmonella typhimurium* as test microorganisms. Six species were showed anti microbial activity expressed as clear zone around the colonies agar (diameter 0,8 cm, height 3 mm). Diameters of the clear zone were observed each 24 hours for 10 days in order to establish diversity of the anti microbial activities.

The profile of antimicrobial power of four *Streptomyces* sp. isolates reveal different activities. *Streptomyces* sp. 2 has the highest activity in inhibiting *Staphylococcus aureus* ATCC 25923 at day 2 with inhibitory zone diameter as much as 27.6 mm, *Escherichia coli* ATCC 25922 at day 5 with inhibitory zone diameter as much as 29.2 mm, and *Salmonella typhimurium* at day 7 with inhibitory zone diameter as much as 27.5 mm, *Streptomyces* sp.-3 has the highest activity in inhibiting *Pseudomonas aeruginosa* ATCC 27853 at day 6 with inhibitory zone diameter of 42.4 mm. *Streptomyces* sp. 4 has the highest activity in inhibiting *Bacillus subtilis* at day 5 with inhibitory zone diameter of 20.7 mm.

Phylogenetic analyses based on PCR showed that they formed distinct phyletic band. Three isolates (sp.-1, sp.-3, and sp.-4) were determined based on their 16S rRNA sequences. These isolates were assigned to genus *Streptomyces*. It is evident that by an partial 16S rRNA gene sequence of the strain that it formed a distinct phyletic line within the range of variation encompassed by the genus *Streptomyces* and the sharp separation of the organism from representatives of the genus *Streptomyces* was strengthened by the fact that is sequencing 16S rRNA differed from those of 1489 recognized *Streptomyces* species.

Key words : *Streptomyces*, mangrove, 16S rRNA gene, antibiotic

UCAPAN TERIMA KASIH

Alhamdulillah Robbil 'Alamin, segala puji dan syukur penulis panjatkan ke hadirat Allah subhaanahu wa ta'ala atas rohman dan rohim-Nya, sehingga diberi kemudahan, kekuatan, dan kemampuan dalam menyelesaikan penelitian ini.

Penelitian ini dapat terselesaikan berkat bantuan berbagai pihak, karena itu saya mengucapkan terima kasih yang setinggi-tingginya kepada Pemerintah Republik Indonesia cq Direktorat Jenderal Pendidikan Tinggi, Departemen Pendidikan Nasional yang telah membiayai penelitian ini.

Ucapan terima kasih saya sampaikan kepada Rektor Universitas Airlangga Prof. dr. H. Fasich, Apt atas kesempatan dan fasilitas yang diberikan kepada saya untuk melaksanakan penelitian Hibah Bersaing di Universitas Airlangga Surabaya.

Ucapan terima kasih yang sebesar-besarnya saya sampaikan kepada Ketua Lembaga Penelitian dan Pengabdian Kepada Masyarakat Universitas Airlangga Prof. Dr. Sarmanu, MS, Dekan Fakultas Kedokteran Universitas Airlangga Prof. Dr. H. Muhammad Amin, dr., SpP., Ketua Laboratorium Mikrobiologi Fakultas Kedokteran Universitas Airlangga, dr. Eddy Mudihardi, MS., SpMK(K), saya ucapkan terima kasih atas izin dan bantuan moril maupun materiil. Ucapan terima kasih juga saya sampaikan kepada Ketua *Tropical Disease Center* (TDC) Prof. Dr. Yoes Prijatna Dachlan, dr, M.Sc., yang telah memberikan bantuan fasilitas peralatan dan konsultasi untuk mendukung analisis penelitian ini.

Akhirnya kepada semua pihak yang tidak dapat saya sebutkan satu persatu yang telah membantu saya selama penelitian, saya sampaikan terima kasih yang sebesar-besarnya. Semoga Allah subhanaahu wa ta'ala senantiasa melimpahkan rahmat dan hidayahNya kepada kita semua. Amin.