Maintenance of Cultures

Cultures sent to customers remain alive for at least 14 days provided that the following precautions are observed. Cultures should be unpacked immediately after receipt and stored at 15-18°C under low light intensity (north window, no direct sun light, or weak white fluorescent light). Screw caps or vessels should be loosened but not removed. Further maintenance or multiplication of cultures requires transfer into new culture media. This presupposes experience in simple microbial techniques.

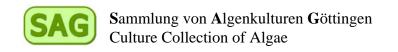
Many species are cultivated and dispatched on agar media for safety reasons but develop their morphological characteristics only in liquid media, e.g. flagellates, colony-forming Volvocales and Chlorococcales. For teaching purposes these species should be transferred into liquid media 2-3 weeks before demonstration, e.g. into Soil Water Media, Basal Medium, or Desmidiacean Medium.

Culture Media

The following media have proved suitable for the maintenance of cultures in test tubes at the SAG for many years. The recipes originate from E. G. Pringsheim and W. Koch, unless stated otherwise. It must be emphasized that the maintenance medium indicated is not always the best medium for the cultivation of a species. There are other media which are just as suitable, e.g. those given in the catalogues of other culture collections of algae (Thompson et al., 1988, Watanabe and Nozaki, 1994, Andersen et al., 1991, Rippka and Herdman, 1992, Starr and Zeikus, 1993). Mass algal culture often requires more concentrated media (for recipes and methods consult Kuhl and Lorenzen, 1964; Starr, 1971; Stein, 1973; Guillard, 1975; Werner, 1982; Castenholz, 1988; Richmond, 2004; Andersen, 2005).

All solutions should be made up with de-ionized water. Media are usually prepared from stock solutions of macronutrients, trace metals, and vitamins which are added to a large proportion of the final volume of water in order to avoid precipitation.

Media may be used as liquid or solidified by 1.0-1.5% agar. Before sterilization the agar has to be dissolved in the medium in a steamer. After this test tubes should be filled with 10 ml of the hot medium, closed with cotton plugs, sterilized (usually by autoclaving at 121°C for 15 min.) and may be stored for several weeks, after cooling, in a refrigerator. Solid media for Cyanobacteria are prepared by mixing, after cooling to 50°C, equal volumes of separately autoclaved double strength solutions of the mineral salts medium and either agar to give a final agar concentration of 0.6-1.0 %.



19. Z-Medium for Cyanobacteria (= Z)

NaNO ₃	467.0 mg
$Ca(NO_3)_2$. $4H_2O$	59.0 mg
K_2HPO_4	31.0 mg
$MgSO_4$. $7H_20$	25.0 mg
Na_2CO_3	21.0 mg
Fe-EDTA complex *	10.0 ml
micronutrient solution **	0.08 ml
de-ionized or distilled water	1000.0 ml

^{*} Preparation of the Fe-EDTA complex: 5 ml of a 0.1M solution of FeCl $_2$. 6 H $_2$ O (= 27g/l) in 0.1N HCl and 5 ml of a 0.1M solution of the dinatrium-salt of EDTA (=37.2g/l Titriplex III, Merck) in de-ionized or distilled water were combined and made up to 500 ml.

** Composition of the micronutrient solution (after H. Gaffron)

To 100 ml of de-ionized or distilled water add:

H_3BO_3	310.0 mg
$MnSO_4$. $4H_2O$	223.0 mg
Na_2WO_4 . $2H_2O$	3.3 mg
$(NH_4)_6Mo_7O_{24}$. $4H_2O$	8.8 mg
KBr	11.9 mg
KJ	8.3 mg
$ZnSO_4$. $7H_2O$	28.7 mg
$Cd(NO_3)_2$. $4H_2O$	15.4 mg
$Co(NO_3)_2$. $6H_2O$	14.6 mg
CuSO ₄ . 5H ₂ O	12.5 mg
$NiSO_4(NH_4)_2SO_4$. $6H_2O$	19.8 mg
$Cr(NO_3)_3$. $7H_2O$	3.7 mg
$VOSO_4$. $2H_2O$	2.0 mg
$Al_2(SO_4)_3K_2SO_4$. 24 H_2O	47.4 mg

a) Z 45/4 Medium for Cyanobacteria (without nitrogen) (= Z 45/4)

$\mathrm{KH_{2}PO_{4}}$	41.0 mg
K_2HPO_4	17.0 mg
$CaCl_2$. $2H_2O$	37.0 mg
$MgSO_4$. $7H_2O$	25.0 mg
Fe-EDTA complex	10.0 ml
micronutrient solution	0.06 ml
de-ionized or distilled water	1000.0 ml

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Further recommended literature about culturing algae:

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- Belcher & Swale (1982) Culturing Algae a guide for schools and colleges. ISBN 1-871105-04-8 (ask for at ccap@sams.ac.uk). (Currently unavailable).
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