

Improved techniques for study of caratenoid intermediates

R. E. Subden

G. Turian

Follow this and additional works at: <https://newprairiepress.org/fgr>



This work is licensed under a [Creative Commons Attribution-Share Alike 4.0 License](https://creativecommons.org/licenses/by-sa/4.0/).

Recommended Citation

Subden, R. E., and G. Turian (1969) "Improved techniques for study of caratenoid intermediates," *Fungal Genetics Reports*: Vol. 15, Article 8. <https://doi.org/10.4148/1941-4765.1911>

This Technical Note is brought to you for free and open access by New Prairie Press. It has been accepted for inclusion in Fungal Genetics Reports by an authorized administrator of New Prairie Press. For more information, please contact cads@k-state.edu.

Improved techniques for study of caratenoid intermediates

Abstract

Improved techniques for study of caratenoid intermediates

Subden R.E. and G. Turian. Improved techniques
for study of carotenoid intermediates in *Neurospora*.

Previous studies of *Neurospora* carotenoids have been hampered by low total carotenoid yields (0.08 = 1% of dry weight) or by a distribution of intermediate pool sizes which favored the end product; e.g., neurosporaxanthin accounts for up to 90% of the total carotenoid fraction.

Enhanced intermediate pool sizes have been obtained by using neurosporaxanthin-less or yellow "albino" strains; e.g., ylo-1 or ylo-b, ALS-4, ALS-23. These strains have 55-75% of the total carotenoid fraction yields of the wild type strains, mostly in the form of the early precursor pools (phytofluene, β -carotene, neurosporene, etc.).

Huang (1964 Genetics 49:453) and Harding (1968 Neurospora Newsl. 13: 8) reported yield improvements by culturing in the dark in liquid medium for 5 days and then draining off the medium and exposing the spread-out mycelial mat to intense fluorescent light for 1 to 24 hours. Cold treatments (6 hrs. at 7°C) also seem to improve yield?, but as yet no quantitative data are available.

Using the above techniques, it has been possible to obtain a yield of 1.8% (total carotenoid fraction/dry weight of mycelium) and isolate short-lived intermediates. β -zeacarotene has already been identified as a component of the *Neurospora* carotenoid fraction using this technique, which was developed in conjunction with a genetic study attempting to define the specific biosynthetic lesions caused by the "albino" gene cluster alleles. - - - Laboratory of General Microbiology, University of Geneva, Geneva, Switzerland.