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Cellulose has been intensively studied in a wide range of organisms (Norkram 1967 Adv. Appl. Microbiol. 9:91), and has recently been investigated in Neurospora from both an industrial (Kuroda 1968 Chem. Abs. 68: 18955d) and a theoretical standpoint (Meyers and Eberhard 1966 Biochem. Biophys. Res. Commun. 24: 782). Although there have been attempts to rationalize the cellulose assay (β -1,4-glucanase - Cx) on an absolute basis (bonds broken per second - Almin and Eriksson 1968 Arch. Biochem. Biophys. 124: 129) there is a lack of standardization between published methods. For example, carboxymethyl cellulose (CMC) has been used in several different states each with various degrees of substitution (DS) and of polymerization (DP), which control the rate of the reaction. Its ionic character limits its use in viscometric assays, as the viscosity is dependent on pH, ionic strength and polyvalent cation content of the assay medium. These limitations are magnified in the more enzymically reactive, lower substituted CMC's. Glycol cellulose (hydroxy ethyl cellulose - HEC) has been proposed to replace CMC in order to circumvent these difficulties (Iwasaki et al. 1964 J. Biochem. (Tokyo) 55:30) but has been rarely used, presumably because these authors noted the laborious procedure of making this non-ionic substituted cellulose. A rigorously controlled range of HEC's is available commercially and this allows the facile development of more standardized redoximetric and viscometric cellulose assays. For this purpose we have used the readily soluble Natrosol 250 M HEC (4,500-6,500 centipoises at 2%, mean substitution 2.5, DS ca. 1.0, DP 565 - Hercules Powder Co., Wilmington, Delaware). Equivalent HEC's are available from Farbwerke Hoechst A. G., Frankfurt, Germany (Tylose H 4000) and Union Carbide Corp., New York (Cellotize WP 4400H). Viscometric assays were carried out using a Cannon-Fenske Viscometer (No. 200: efflux time of solvent 10.2 sec.) at 25°C with a reaction mixture of 7 ml 0.44% HEC (250 M Natrosol), 1 ml 0.5 M sodium acetate buffer pH 4.8 and 2 ml enzyme. The substrate was dissolved in water by shaking overnight at room temperature. Blending to aid the rapid solution of the substrate for as little as five seconds gave a much reduced viscosity level. Redoximetric analyses were performed using equivalent reaction mixture, reducing end groups being estimated by the ferricyanide method modification proposed by Park and Johnson (1949 J. Biol. Chem. 181: 149). The colorimetric Somogyi-Nelson method could not be used as it caused precipitation of the substrate and concomitant absorption of the colored complex. Titrimetric analyses of the oligosaccharides by the Somogyi method or by direct reduction with alkaline iodine proved practical but tedious. Enzyme units are defined: one redoximetric unit is that amount of enzyme which produces 1% degradation in 1 minute, under the above conditions, while a viscometric unit is that amount of enzyme which causes a change of Δf (specific fluidity) of 0.10 in 10 minutes when incubated under the above conditions. Apparent zero order kinetics were maintained up to a change of Δf of 0.10. It is important to note that substituted celluloses are degraded at a changing rate

throughout the reaction and hence it is necessary to define the **reduciometric unit at some standard % degradation (1%)**. One **reduciometric unit approximately equals 21 viscometric units, but varies** with the particular enzyme **under investigation**.

A **comparison has** been made of the levels of Cx cellulose produced by Neurospora crassa STA4 (FGSC"262) and N. crassa ATCC 10355. The organisms **were grown** in Vogel's mineral medium N containing **0.5% glucose plus 1% "cellulose"** (750 ml/2 l flask, 30°C with shaking). **Celluloses** used included: HEC (N atrosol 250 M), CMC (Cellofas BIO I.C. I. Ltd., England) and Avicel (Microcrystalline cellulose, FMC Corp., MARCUS Hook, Pennsylvania). Relatively **small amounts** of enzyme were produced under these conditions (Table 1). For example, **a crude commercial cellulose (Trichoderma viride)** has a specific activity of 20.8 (viscometric). More enzyme **was excreted** into the medium **than retained intracellularly** but intracellular levels **were measurable** with these techniques. CMC proved **a better inducer of cellulase** than the other two substrates. Attempts to release **additional bound cellulase** by further incubation of the harvested **mycelium or mycelial homogenates** at pH 7.5 (P04) 16 hours (3°C) proved successful.

Table 1. The induction of cellulase by "celluloses" in two strains of Neurospora crassa (viscometric units).

Inducer	<u>N. crassa</u> 10355 (4 days)				<u>N. crassa</u> STA4 (4 days)				<u>N. crassa</u> STA4 (8 days)			
	Extracellular		Intracellular		Extracellular		Intracellular		Extracellular		Intracellular	
	μ/ml	SpAct	μ/ml	SpAct	μ/ml	SpAct	μ/ml	SpAct	μ/ml	SpAct	p/ml	SpAct
HEC	0.14	0.72	0.30	0.21	0.28	2.55	0.12	0.06	0.17	1.70	1.16	0.39
CMC	0.68	4.86	0.11	0.06	1.20	14.20	0.64	0.42	1.46	13.2	0.94	0.40
AVICEL	0	■	0	■	0.10	1.25	0.01	0.18	0.42	4.2	0.47	0.66