

Application of novel analytical ultracentrifuge analysis to solutions of fungal mannans

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Abstract Polysaccharides, the most abundant biopolymers, are required for a host of activities in lower organisms, animals, and plants. Their solution characterization is challenging due to their complex shape, heterogeneity, and size. Here, recently developed data analysis approaches were applied for traditional sedimentation equilibrium and velocity methods in order to investigate the molar mass distribution(s) of a subtype of polysaccharide, namely, mannans from four *Candida* spp. The molecular weight distributions of these mannans were studied using two recently developed equilibrium approaches: SEDFIT-MSTAR and MULTISIG, resulting in corroboratory distribution profiles. Additionally, sedimentation velocity data for all four mannans, analyzed using $ls-g^*(s)$ and Extended Fujita approaches, suggest that two of the fungal mannans (FM-1 and FM-3) have a unimodal distribution of

molecular species whereas two others (FM-2 and FM-4) displayed bi-modal and broad distributions, respectively: this demonstrates considerable molecular heterogeneity in these polysaccharides, consistent with previous observations of mannans and polysaccharides in general. These methods not only have applications for the characterization of mannans but for other biopolymers such as polysaccharides, DNA, and proteins (including intrinsically disordered proteins).

Keywords AUC · Extended Fujita approach · MULTISIG · SEDFIT-MSTAR · Sedimentation

Introduction

Mannans are polysaccharides containing D-mannose, and are found as cell wall components in bacteria, fungi, (moulds and yeast) and plants. Pure mannan is uncommon in plants but it is one of the major components of the yeast cell wall together with glucan, chitin, and protein such as mannoprotein. Mannans have different kinds of structures in various organisms. Figure 1a describes the structure of plant mannan, which has a backbone of linear chains made up of $\beta(1 \rightarrow 4)$ -linked Mannopyrosyl (Man_p) residues (Tombs and Harding 1998). Plant mannans occur in the cell walls as heteropolysaccharides, i.e., glucomannans, galactoglucomannans, and galactomannans (Ebringerova et al. 2005). Generally, *Candida* spp. mannans have an $\alpha(1 \rightarrow 6)$ linked backbone (Fig. 1b) substituted mostly at O-2 by different number of linear or branched side oligomannosyl chains composed of $\alpha(1 \rightarrow 2)$, $\alpha(1 \rightarrow 3)$ and $\alpha(1 \rightarrow 6)$ links with or without terminal $\beta(1 \rightarrow 2)$ linkages (Shibata et al. 1992, 1996, 2003, 2007). It was also reported that there are subtle variations in the linkage or number

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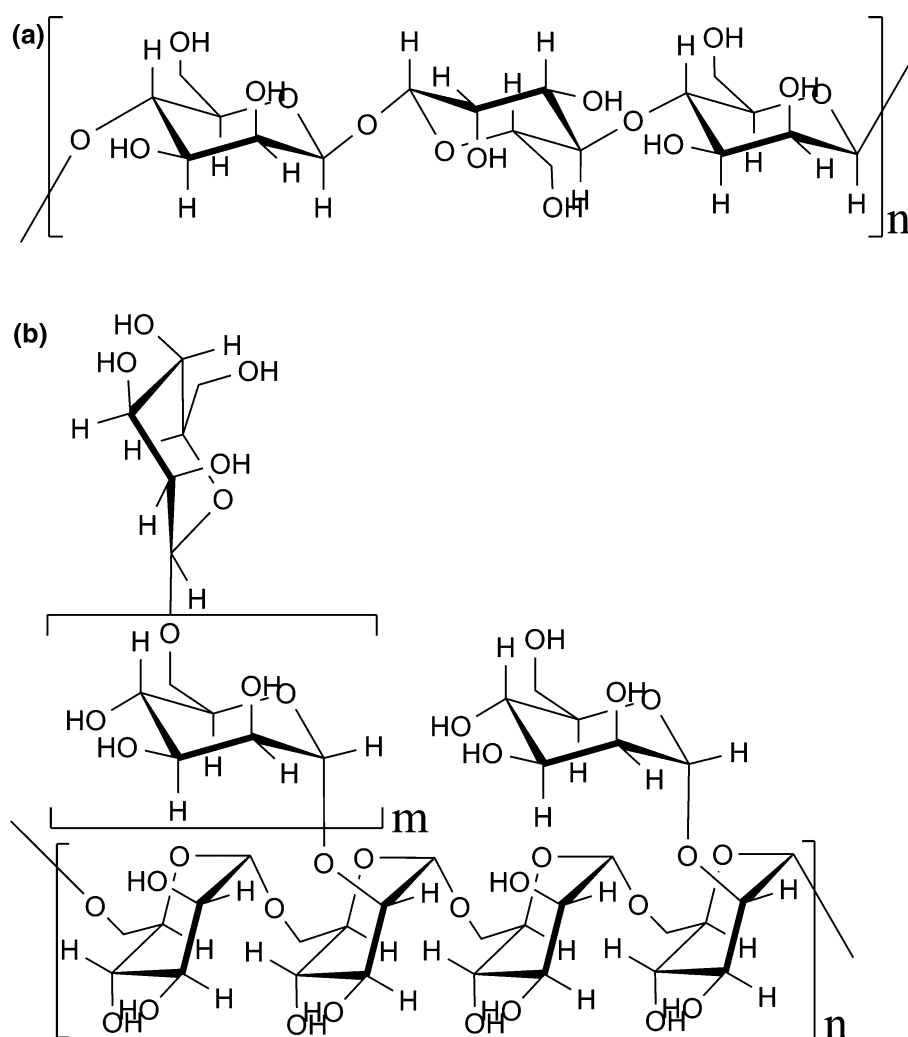
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Fig. 1 Chemical structure of mannan from **a** plant sources and **b** fungal sources



of mannose residues in side chains of *Candida* spp. mannan molecules (Nelson et al. 1991). *C. tropicalis* mannan is composed of an $\alpha(1 \rightarrow 6)$ -linked backbone substituted only with $\beta(1 \rightarrow 2)$ linked and $\alpha(1 \rightarrow 2)$ linked Man_p units without $\alpha(1 \rightarrow 3)$ linked ones observed in *C. albicans* mannan (Kobayashi et al. 1994; Suzuki et al. 1997). *C. dubliniensis* has been isolated from the HIV-positive individuals in the beginning of 1990 [(Pujol et al. 2004) and references cited therein] and later grouped separately from *C. albicans* by Sullivan et al. (1995) Mannan from *C. dubliniensis* (Ližičárová et al. 2005), as well as mannan from *C. parapsilosis* (Shibata et al. 1995), have very similar structures to that of *C. albicans*.

The anti-tumor activity of polysaccharides was reported by Diller (1947), followed by several other authors who suggested that mannans are potent anticancer agents (Mankowski et al. 1957; Bradner and Clarke 1959; Kamasuka et al. 1968; Oka et al. 1969; Suzuki et al. 1969; Ukai et al. 1983; Peng et al. 2003). It is also critical to characterize these polysaccharides because certain factors,

such as polymer length (proportionally linked to molecular weight), can impact their potency, for example their antigenicity (Kabat and Bezer 1958).

This article is focused on the characterization of fungal mannans using analytical ultracentrifugation (AUC)—a well-established, matrix-free method for the determination of molar mass and sedimentation coefficients. In addition, recently presented analysis techniques have been utilized to determine the molar mass distribution of fungal mannans, along with providing distributions of sedimentation coefficients.

Methods

Four mannan samples, FM-1 (*Candida tropicalis* CCY 29-7-6), FM-2 (*C. dubliniensis* CCY 29-177-1), FM-3 (*C. albicans* CCY 29-3-32), and FM-4 (*C. parapsilosis* CCY 29-20-1) were prepared from the above-mentioned yeast strains obtained from Culture Collection of Yeast (Institute

of Chemistry, Bratislava, Slovakia). The mannans were prepared from fresh yeast biomass as described by Bystrický et al. (2003). Under the used strong alkaline conditions, nearly all covalently linked protein was split from the mannoproteins. FM-3 contained 0.4 % nitrogen whereas FM-1, FM-2, and FM-4 only traces.

Sample preparation

Samples were dissolved in phosphate buffered saline (PBS, pH 7.0, $I = 0.1$ M), made with 0.05 M sodium chloride, and 0.05 M phosphate salts (dibasic sodium and potassium dihydrogen) in screw-capped tubes with constant stirring at low speed. During this period, the temperature was raised to 80.0 °C for 10 min to obtain maximum solubility. Stirring continued overnight at room temperature at low speed. Samples were subsequently centrifuged at 10,000 rpm ($11,600\times g$) for 15 min (Beckman L8-55 M Ultracentrifuge). Concentrations of stock solutions were measured using a differential refractometer (Atago DD-5, Jencons Scientific) and a refractive index increment of 0.171 ml g⁻¹ (Mueller et al. 2000).

Molar mass determination using sedimentation equilibrium

All mannan samples were subjected to sedimentation equilibrium analytical ultracentrifugation (AUC-SE) experiments using the Optima XL-I Analytical Ultracentrifuge (Beckman Instruments, Palo Alto, CA). Double-sector carbon-filled epoxy 12-mm path-length centerpieces, loaded into aluminium housings and sealed with sapphire windows, were used to load solvent (90 μl), and sample (80 μl) at 1.0 mg ml⁻¹. Cells were loaded into eight-hole titanium rotor (An50Ti) and placed in the centrifuge. Samples were centrifuged at 16,000 rpm ($\sim 20,600\times g$) at (20.0 ± 0.1) °C. Scans were taken using Rayleigh interference optics once every hour until equilibrium was achieved.

Data analysis was performed using two independent algorithms. SEDFIT-MSTAR (Schuck et al. 2014), which utilizes a smart-smooth method to fit the raw data curve, and M^* function (Harding et al. 1992), to provide the weight average of the entire distribution of molar masses:

$$M^*(r) = \frac{J(r) - J}{kJ_m(r^2 - r_m^2) + 2k \int_{r_m}^r (J(r) - J)r \cdot dr} \quad (1)$$

$$k = \frac{(1 - \bar{v}\rho)\omega^2}{2RT} \quad (2)$$

In Eqs. (1) and (2), r is the radial distance from the center of rotation, r_m the corresponding value at the meniscus, J is the concentration in fringe displacement units, and

J_m the corresponding value at the meniscus. \bar{v} is the partial specific volume, ρ is the solvent density, ω is the angular velocity of the rotor, R is the gas constant and T is the absolute temperature.

MULTISIG was then used to fit the relative concentration proportions of 17 molar masses, logarithmically spaced to achieve a tenfold range, to yield a molar mass distribution (Gillis et al. 2013). The total (fringe) concentration at a set radial position is given by:

$$J(r) = \sum_{i=1}^{i=17} J_{\text{ref}} \exp\left\{0.5\left(0.5kM_i 1.15^{(i-1)}\left(r^2 - r_m^2\right)\right)\right\} + E \quad (3)$$

where M_i is the species molar mass, J_{ref} is the reference concentration (typically the concentration at the consensus hinge point), and E is the baseline. MULTISIG/RADIUS was used to apply this procedure to radial positions along the column length of solution. Both methods were performed in pro Fit™ (QuantumSoft, Switzerland).

Sedimentation velocity analysis

Sedimentation coefficient distributions of mannans were measured using sedimentation velocity (AUC-SV) in the analytical ultracentrifuge. Solvent (400 μl) and sample (390 μl, 1.0 mg ml⁻¹) were loaded into similarly constructed cells as for AUC-SE. Cells were centrifuged in the Beckman Optima XL-I analytical ultracentrifuge at 40,000 rpm ($\sim 130,000\times g$) at (20.0 ± 0.1) °C. Data were analyzed using least squares apparent distributions of sedimentation coefficients (ls-g*(s) vs. s) from SEDFIT (Schuck and Rossmanith 2000), and curve fitting module MULTIG in pro Fit™ (QuantumSoft, Switzerland). Weight-average sedimentation coefficients (s) for particular components were corrected to standard solvent conditions (density and viscosity of water at 20.0 °C) to yield $s_{20,w}$ (S), using SEDNTERP (Laue et al. 1992), and a \bar{v} of 0.625 ml g⁻¹ (Gray and Ballou 1971). The $s_{20,w}$ was measured at a range of concentrations (0.2–2 mg ml⁻¹) for all samples and extrapolation performed for each to zero concentration to obtain $s_{20,w}^0$ to eliminate the effects of non-ideality.

Results and discussion

Molar mass distribution

Mannan samples, both dialyzed and undialyzed, were probed for their molar masses using sedimentation equilibrium. There was little observable difference in terms of their molar masses between dialyzed and undialyzed

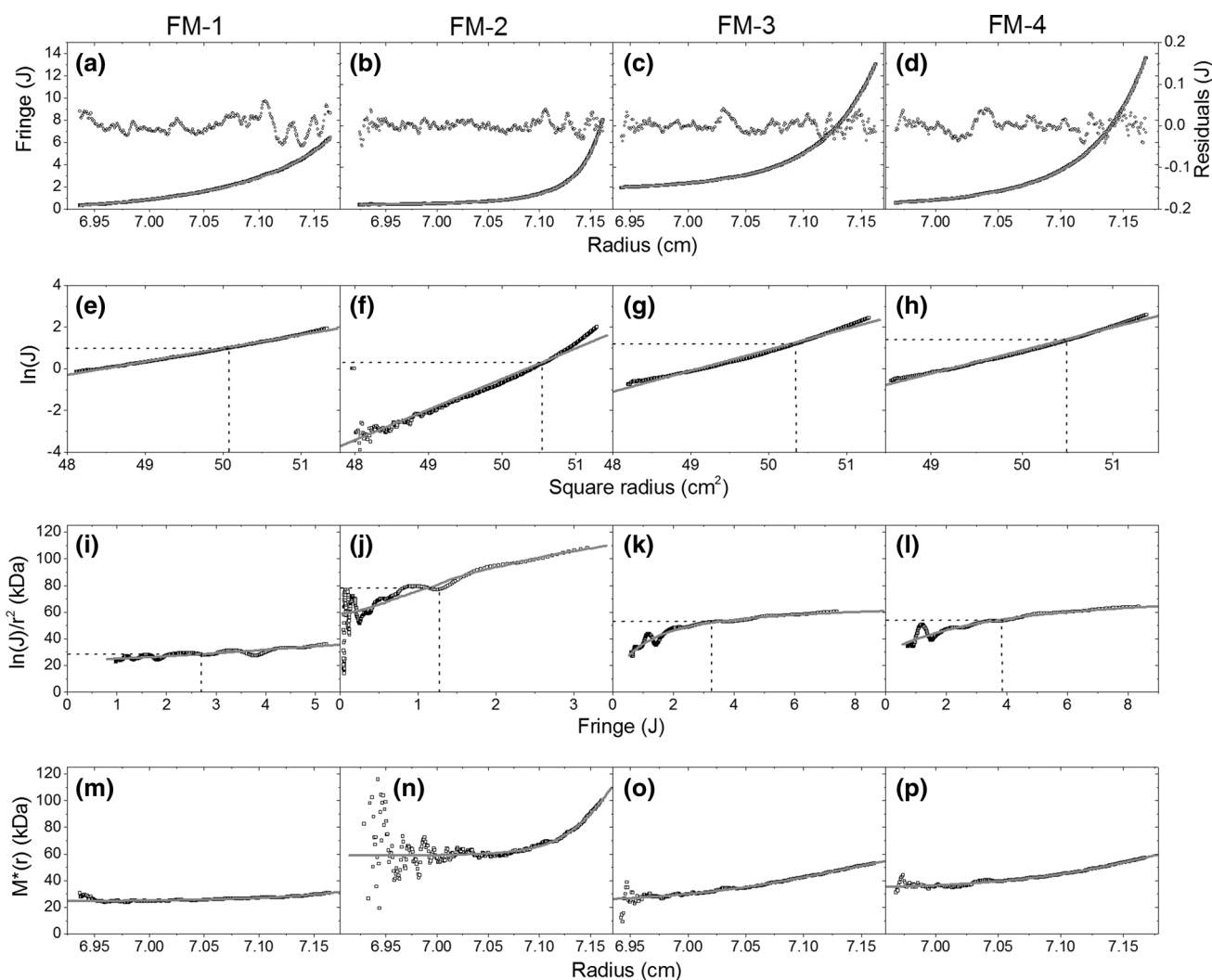


Fig. 2 Output from SEDFIT-MSTAR of FM1-4 (left to right). **a–d** Fringe displacement (J) vs. radius with residual between raw data (hollow square) and fit (grey line); **e–h** natural logarithm of baseline-

corrected fringe displacement J vs. square radius; **i–l** differential of **e–h** yielding apparent molar mass vs. concentration; **m–p** $M^*(r)$ algorithm extrapolating to cell base

samples (data not shown), suggesting the sample was of high purity. Results presented below are from dialyzed samples.

SEDFIT-MSTAR

SEDFIT-MSTAR fitted the raw fringe displacement data using the smart-smooth analysis (Fig. 2a–d) with results shown in Table 1. The grey lines represent the result of the fitted parameters. Residuals show no overall trend and do not deviate beyond 0.1 fringes in all samples. The natural logarithm of (baseline-corrected) fringe displacement (J) versus the square of the radius (Fig. 2e–h) shows a near-straight line but with small positive curvature, particularly in FM-2. This indicates polydispersity. The differential of Fig. 2e–h yields Fig. 2i–l representing point-average,

apparent $M_w(r)$ as a function of the concentration $c(r)$ across the cell. Positive slopes suggest polydispersity, especially in FM-2 (Fig. 2j) where values for the $M_w(r)$ appear to fall significantly as zero concentration is approached. This result is consistent with results from ls-g*(s) analysis suggesting a bimodal distribution; thus the lower radial positions in the cell would contain a high proportion of the low-molar-mass species.

The MSTAR algorithm yielded weight-average molar masses (M_w) ranging from 3.1 to $11.2 \times 10^4 \text{ g mol}^{-1}$ through the extrapolation of the $M^*(r)$ to the base of the cell (Fig. 2m–p) and z -average molar masses (M_z) ranging from 4.1 to $14 \times 10^4 \text{ g mol}^{-1}$. The ratio of M_z/M_w provides the polydispersity index (PDI) ranging from 1.2 to 1.3, typical for a polydisperse system. Consensus hinge points (CHP—the point at which, during the approach to equilibrium, the

Table 1 Weight average and *z*-average molar mass estimates from AUC-SE of fungal mannans using SEDFIT-MSTAR

Sample	$\times 10^{-3}$ <i>w</i> -average molar mass ^a (g mol ⁻¹)	$\times 10^{-3}$ <i>w</i> -average molar mass ^b (g mol ⁻¹)	$\times 10^{-3}$ <i>z</i> -average molar mass ^b (g mol ⁻¹)	PDI ^b (<i>z</i> / <i>w</i>)
FM-1	30.2	31.5	41.6	1.32
FM-2	83.3	112	140	1.25
FM-3	56.1	54.8	64.7	1.18
FM-4	58.5	59.6	71.6	1.20

Loading concentrations were approximately 1 mg ml⁻¹. Polydispersity index (PDI) measured as a ratio of *z*-average and *w*-average from SEDFIT-MSTAR *c*(*M*) fit

^a From the Consensus Hinge Point (CHP) method

^b From extrapolation of *M*^{*}(*r*) to the cell base (Eq. 1, Fig. 2) and SEDFIT-MSTAR fit

concentration does not change significantly over time; as well as an indication of the loading concentration of the sample) were measured as an internal check of the rigor of the MSTAR analysis. Hinge points are indicated in Fig. 2 by dashed lines and results are presented, with the other results from MSTAR and *c*(*M*) analysis, in Table 1. CHP results do not deviate greatly from the MSTAR and *c*(*M*) results, suggesting that non-ideality had no significant impact on the overall analysis (Schuck et al. 2014).

MULTISIG

MULTISIG approximates the real solute distribution present by a series of ‘concentration’ coefficients attached to terms in reduced molar mass value, which are logarithmically spaced, thus yielding a distribution (*g*(*M*) vs. *M*) (Gillis et al. 2013). These are presented in Fig. 3. FM-2 and FM-4 both show two peaks, which is consistent with *ls-g*^{*}(*s*) distributions. Number, weight, and *z*-average molar masses, along with polydispersity indices, are shown in Table 2. Estimates are also made in Table 2 for peak molar masses and relative concentrations.

MULTISIG/RADIUS

Whilst MULTISIG fits 17 discrete species at a selected radial position (in this investigation, the CHP, i.e., the point in the curve where the concentration does not change over the approach to equilibrium), MULTISIG/RADIUS provides this estimate at 20 points along the range of the cell. The baseline was fixed based on ten iterations from the previous MULTISIG analysis. Results yielded number, weight, and *z*-average reduced molar masses plotted against the concentration range of the cell and a 3D contour plot of *g*(*M*) vs. *M* vs. concentration range (Fig. 4). Number, weight, and *z*-average molar masses produced similar trends to those found from SEDFIT-MSTAR (Fig. 2i–l), which is unsurprising since these plots should provide equivalent information, despite being independently calculated. The contour plot indicates that the twin-peak

distributions found in FM-2 and FM-4 are present throughout the cell context with little change in relative strength—with the exception of FM-2, which has a low proportion of the high-*M_w* species at the top of the cell. This can be explained by the high *g* force depleting the high-*M_w* component at the meniscus. PDI (*z*/*w*) is also consistent to within a reasonable margin of error compared to SEDFIT-MSTAR values, although a direct comparison is difficult due to the peak-identifying nature of MULTISIG and the whole-solution evaluation from SEDFIT-MSTAR.

Weight averages present themselves directly in-between the two peaks of FM-2 and FM-4, whereas they lie closer to the main peaks present in FM-1 and FM-3; however, they are slightly skewed towards small amounts of smaller (FM-3) or larger (FM-1) material. It is unclear whether these smaller or larger species are indicative of impurities or algorithmic anomalies, however the averages are similar to those found from the independent SEDFIT-MSTAR analysis approach.

Sedimentation coefficient distribution

The *ls-g*^{*}(*s*) profiles for FM-1, FM-2, FM-3 and FM-4 are shown in Fig. 5, at approximately 2 mg ml⁻¹, as normalized distributions. The profiles indicate that FM-1 and FM-3 yielded a single distribution at ~4 S with a small degree of larger material in the high-sedimentation range (up to 20 S). FM-4 peaks at ~4 S and has a larger proportion of faster-sedimenting material, particularly at 11 and 14 S. FM-2 yielded a two-peak distribution at ~4 and ~9 S.

The concentration for each component was re-calculated based on the initial loading concentration, and amount of mannan fraction present, and used to calculate the *s*_{20,w}⁰ (S) from *s*_{20,w}. Extrapolations were performed using the reciprocal sedimentation coefficients as shown in Fig. 5 as before (Patel et al. 2006, 2008). The Galen coefficient, (*k_sc*) was calculated according to Eq. (4) (Rowe 1992) and presented in Table 3.

$$\frac{1}{s_{20,w}} = \frac{1}{s_{20,w}^0} (1 + k_s c) \quad (4)$$

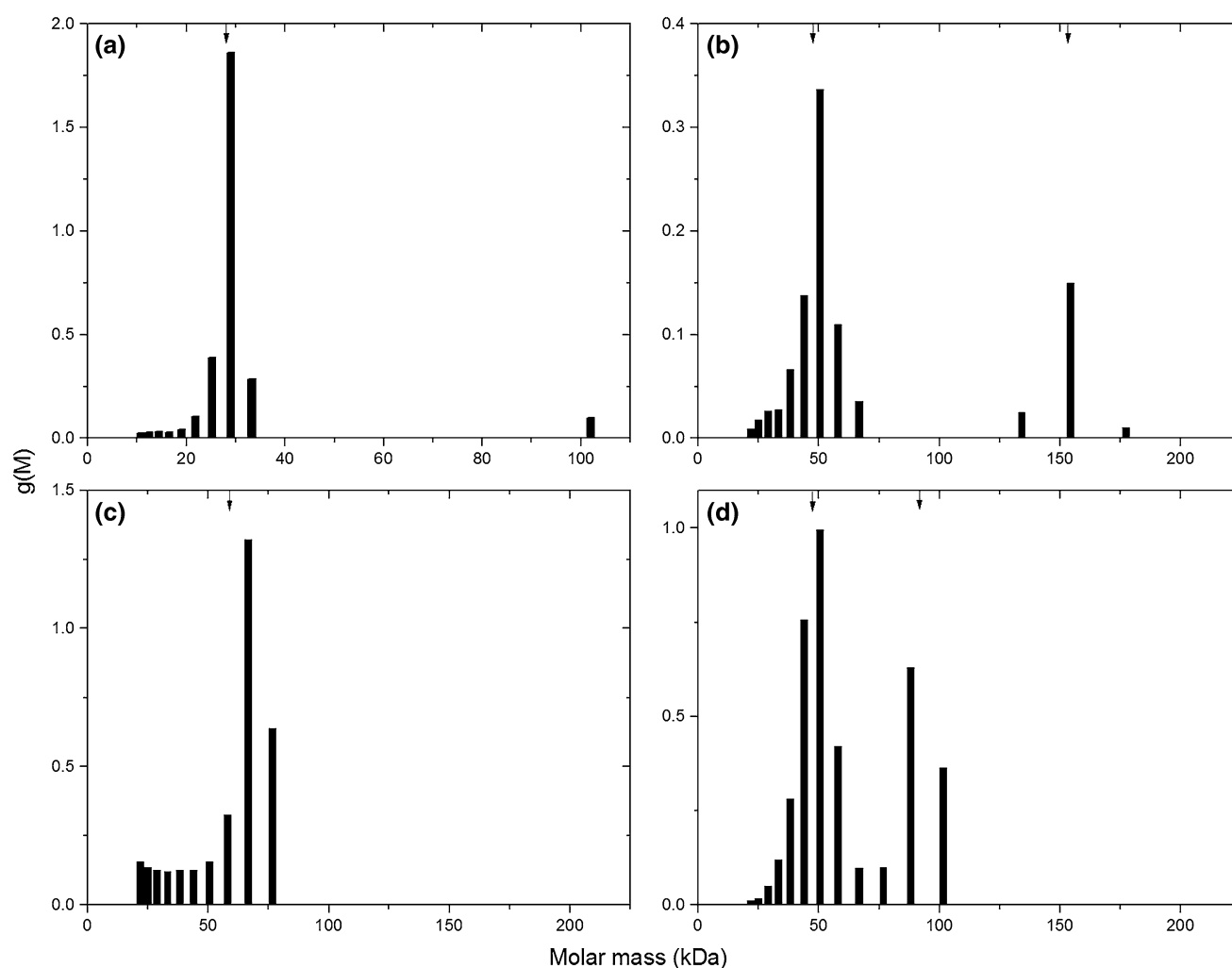


Fig. 3 MULTISIG output of **a** FM-1; **b** FM-2; **c** FM-3; and **d** FM-4, including 17 discrete molar mass values. *Arrows* represent weight-average of peak(s)

Table 2 Output from MULTISIG of fungal mannans measured at a loading concentration of 1 mg ml⁻¹

Sample	Peak	Fraction (%)	$\times 10^{-3}$ <i>n</i> -average molar mass (g mol ⁻¹)	$\times 10^{-3}$ <i>w</i> -average molar mass (g mol ⁻¹)	$\times 10^{-3}$ <i>z</i> -average molar mass (g mol ⁻¹)	PDI (<i>z/w</i>)
FM-1	1	97	28.8	29.7	30.3	1.02
	Total	100	29.5 (± 0.2)	32.3 (± 0.1)	39.2 (± 0.0)	1.22
FM-2	1	81	48.5	50.8	52.8	1.04
	2	19	162	163	163	1.00
	Total	100	56.3 (± 1.0)	67.2 (± 0.2)	101 (± 0.0)	1.39
FM-3	1	80	69.0	70.4	71.5	1.02
	Total	100	54.2 (± 0.6)	62.3 (± 0.1)	67.6 (± 0.0)	1.09
FM-4	1	82	49.1	50.8	52.3	1.03
	2	28	96.9	97.6	98.3	1.01
	Total	100	57.1 (± 0.2)	64.2 (± 0.0)	72.3 (± 0.0)	1.13

Averages (*n*, *w*, *z*) and polydispersity index (PDI, *z/w*) produced from both the overall distribution ('Total') and from individual peaks. Standard error of the mean represented by *parentheses*

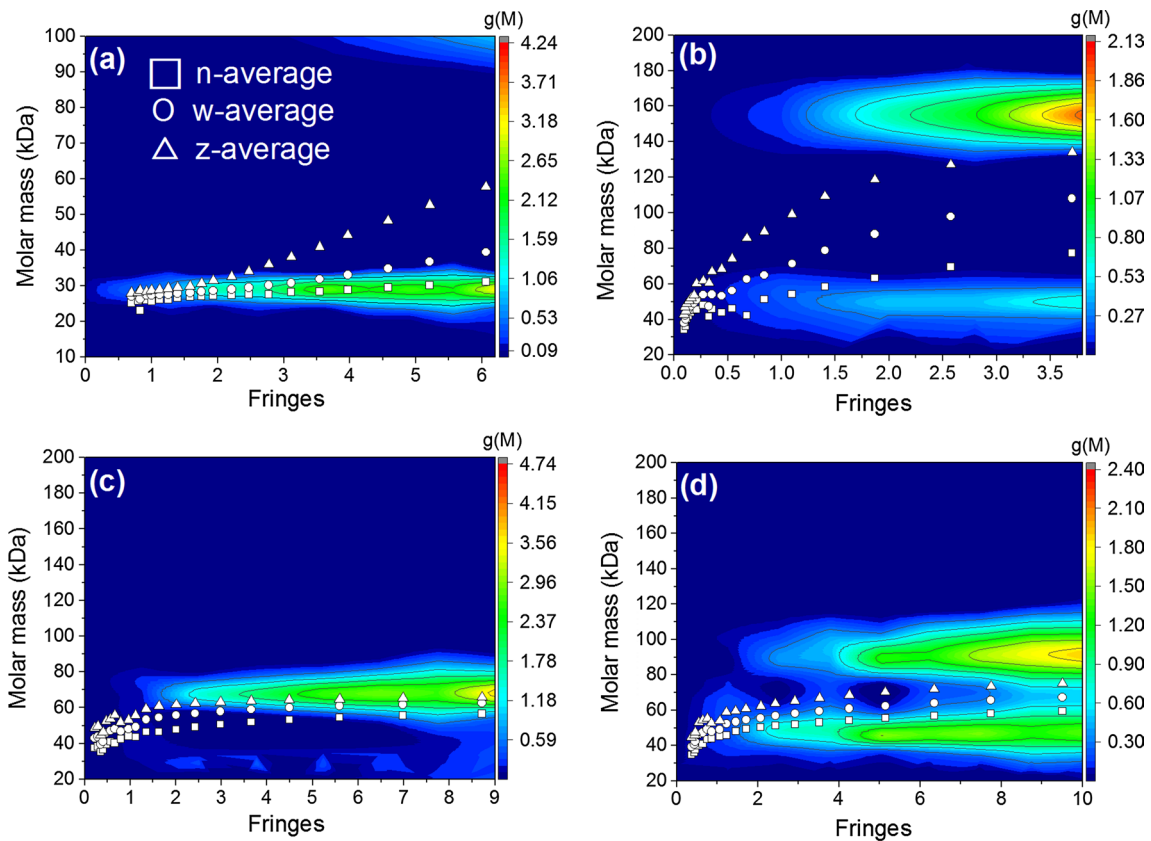


Fig. 4 MULTISIG/RADIUS output of **a** FM-1; **b** FM-2; **c** FM-3; and **d** FM-4. 17 discrete molar masses along 20 points of the cell. Number (square), weight (circle), and z-average (triangle) molar masses overlaid

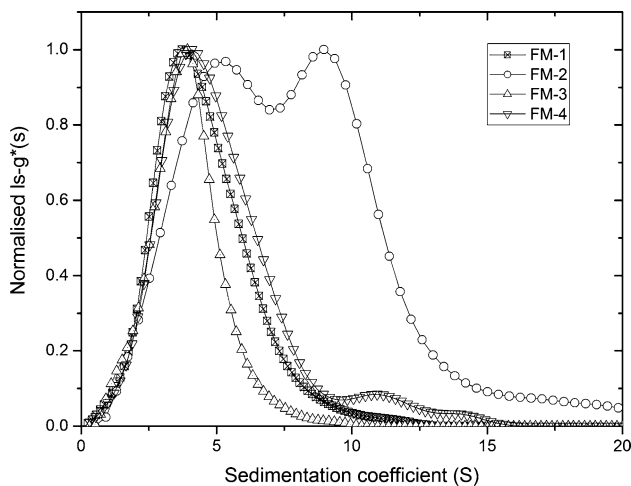


Fig. 5 Normalized $ls-g^*(s)$ vs. sedimentation coefficient of fungal mannans at highest concentration ($\sim 2 \text{ mg ml}^{-1}$)

Figure 6 shows that FM-1 (a) and FM-3 (c) have one component whereas FM-2 (b) and FM-4 (d) had two different components, based on distributions in Fig. 5. It can be observed that these mannan samples show little

Table 3 Summary of sedimentation velocity analysis of fungal mannan samples, including sedimentation coefficients corrected for solvent conditions and extrapolated to infinite dilution

Sample	Peak	Fraction (%)	$s_{20,w}^0$ (S)	k_s (ml g^{-1})
FM-1	1	100	$5.6 (\pm 0.4)$	$470 (\pm 140)$
FM-2	1	53	$5.3 (\pm 0.2)$	$10 (\pm 200)$
	2	47	$9.1 (\pm 0.1)$	$29 (\pm 70)$
FM-3	1	100	$3.3 (\pm 0.1)$	$137 (\pm 65)$
FM-4	1	92	$4.3 (\pm 0.1)$	$60 (\pm 23)$
	2	8	$12.8 (\pm 0.5)$	N/D ^a

Values in parentheses represent standard error of the mean (S) or regression (k_s)

^a Peak 2 of FM-4 was averaged (mean), no linear regression was estimated

dependence of concentration on sedimentation coefficients over the small concentration range studied. The second peak from FM-4 showed no definite trend, thus the values were averaged (gradient = 0) to yield $s_{20,w}^0$ and no k_s estimated because of the very low concentration range.

There is little difference between extrapolated values of native and reciprocal sedimentation coefficients and similar

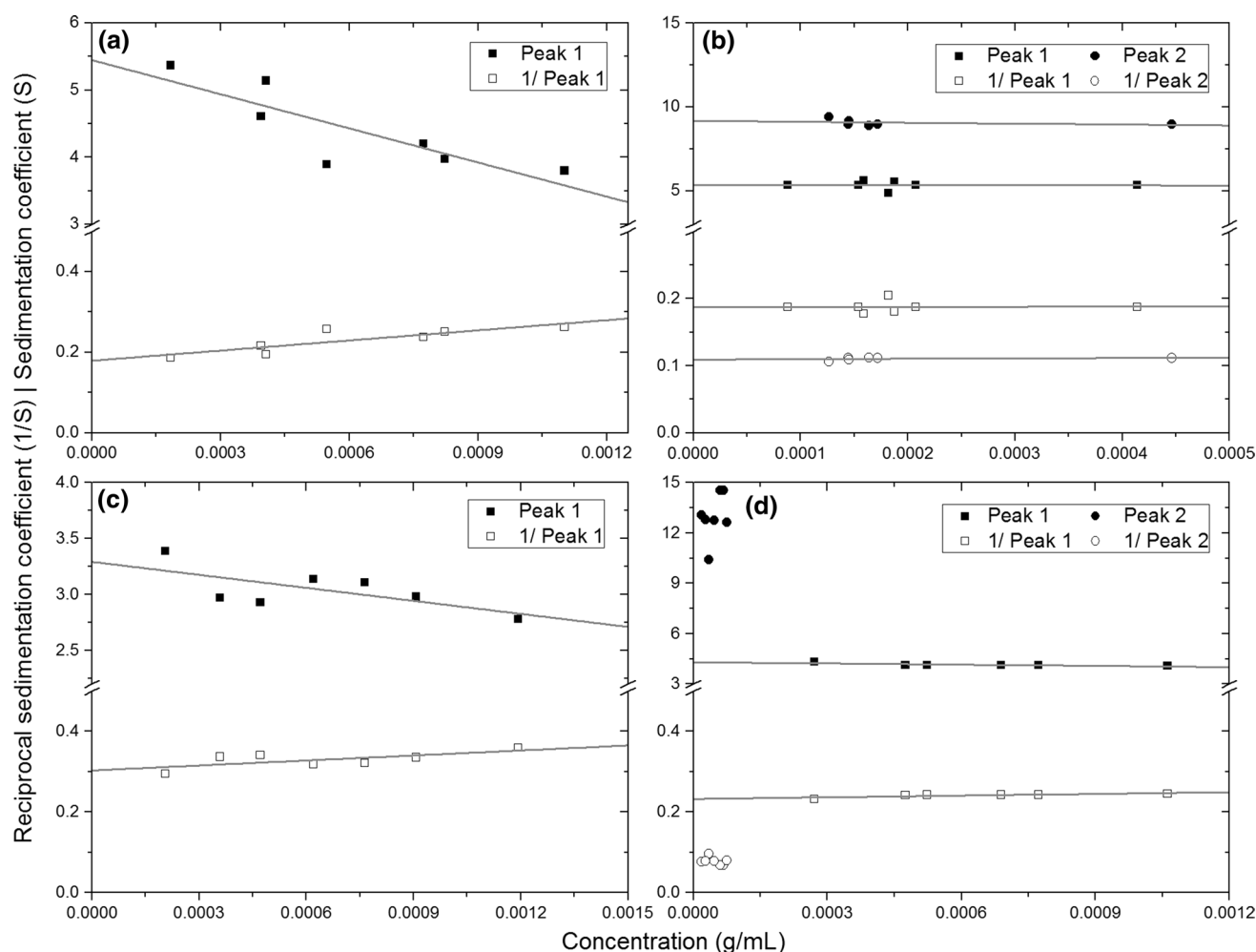


Fig. 6 Extrapolations of integrated peaks from sedimentation velocity measured using linear regression (grey line) of native and reciprocal sedimentation coefficients against concentration of **a** FM-1; **b** FM-2; **c** FM-3; and **d** FM-4

errors. Values ranged between 3 and 6 S, for the smaller peak. Gralen coefficients differed more between the two methods and generally yielded higher error in the reciprocal regression. The single-peak samples were in the 100 range—118–310 ml g⁻¹, whereas the two-peak samples were in the tens range—14–56 ml g⁻¹. This indicates that the single-peak samples have a higher non-ideality than the two-peak systems, however the Gralen coefficients for FM-2 and FM-4 may be affected by the Johnston-Ogston effect (1946).

Molar mass distributions from the extended Fujita approach

Combining information from AUC-SV (sedimentation coefficients) and AUC-SE (molecular weights) allow for the calculation of power-law scaling factors. The slope of a double logarithmic plot of sedimentation coefficients against molar mass yields the Mark–Houwink–Kuhn–Sakurada (MHKS) sedimentation shape factor b (Eq. 5).

$$s_{20,w}^0 = \kappa_s M^b \quad \text{or} \quad M = \left(\frac{s_{20,w}^0}{\kappa_s} \right)^{1/b} \quad (5)$$

where κ_s is an intercept constant (not to be confused with the Gralen coefficient, k_s). This factor ranges between ~0.15 for a ‘rod’ to ~0.67 for a ‘sphere’ and 0.5 is a ‘random coil’. A previously published study showed that the MHKS shape factor was 0.43—this equates to a random coil, but on the ‘stiffer’ end of the scale (Pavlov et al. 1992). The inset of Fig. 7 shows results from this investigation plotted on a double-logarithmic scale. These data points include individual peaks from FM-2 and FM-4, with peak average molar masses obtained from MULTISIG. The gradient was 0.446 (± 0.154), with a κ_s of $4.11 (\pm 3.38) \times 10^{-2}$, which is well within experimental error of the literature value of 0.43, corresponding κ_s of $5.09 (\pm 0.01) \times 10^{-2}$. The agreement of the b from Pavlov et al. (1992) and the present study justifies the assumption we have made that $M_{w,\text{app}}$ measured at 1.0 mg ml⁻¹ is $\sim M_w$.

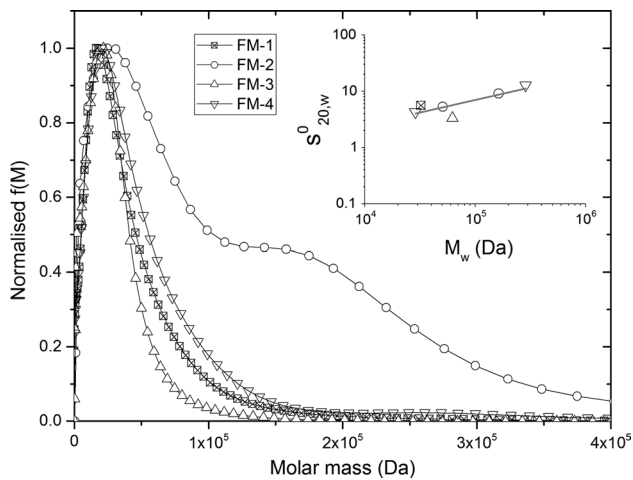


Fig. 7 Extended Fujita approach (normalized $f(M)$ vs. M) for FM-1, FM-2, FM-3, and FM-4. Inset is Mark–Houwink–Kuhn–Sakurada plot of six peaks from AUC-SV and AUC-SE (MULTISIG)

The extended Fujita approach is a method for yielding molar mass distributions from $g(s)$ distributions from AUC-SV (Harding et al. 2011). The original method assumed that the macromolecule was a random coil, which meant that the sedimentation coefficient was directly proportional to the square root of the molar mass:

$$s_{20,w}^0 = \kappa_s M^{0.5} \text{ or } M = \left(\frac{s_{20,w}^0}{\kappa_s} \right)^2 \quad (6)$$

Harding et al. (2011) extended this approach to all conformational types using the general power relation (Eq. 6). To transform a distribution of sedimentation coefficients $g(s)$ vs. s to a distribution of molar masses $f(M)$ vs. M the following transformation equations are used:

$$f(M) = g(s) \frac{ds}{dM} \quad (7)$$

where

$$\frac{ds}{dM} = b\kappa_s^{1/b} s^{(b-1)/b} \quad (8)$$

In this instance, $\ln-g^*(s)$ analysis performed by SEDFIT can be used to substitute $g(s)$ in these equations, with b and κ_s calculated from the double logarithmic plot of sedimentation coefficient and M_w . Using Eq. (5) to modify the abscissa and Eqs. (7) and (8) to modify to ordinate, the (normalized) $f(M)$ vs. M plots are shown in Fig. 7. The distributions show a reduction in peak height for FM-2 peak 2 compared to $\ln-g^*(s)$ from AUC-SV, but correlates more with the $g(M)$ vs. M distribution from MULTISIG AUC-SE. Peak 2 for FM-4 is also reduced, which is less consistent with MULTISIG AUC-SE, which

is likely an over-simplification of species present in solution. FM-1, FM-2, and FM-4 all showed overlapping peaks around $2\text{--}3 \times 10^4 \text{ g mol}^{-1}$ with broad distributions leading to $\sim 1.5 \times 10^5 \text{ g mol}^{-1}$, consistent with information from AUC-SE.

Polydispersity indices calculated from these distributions were significantly higher than those for MULTISIG and SEDFIT-MSTAR, with approximately 50–60 % higher estimations for z/w . The extended Fujita approach is based on κ_s , which was calculated with a high standard error ($4.11 (\pm 3.38) \times 10^{-2}$, ± 82 %) from Fig. 7 inset. From Eq. (8), it is shown that κ_s has a large influence on the calculation of ds/dM and thus on the spread of the distribution. The high apparent polydispersity from $f(M)$ can therefore be attributed to this high error in the κ_s value obtained from Fig. 7 inset. Although, in this instance, the extended Fujita approach provided a poor estimation of polydispersity, it did yield accurate estimates for molar mass and heterogeneity.

Non-ideality

On the subject of non-ideality, concentration dependence was directly measured using AUC-SV to yield k_s values, which showed very low non-ideality for all four mannans. AUC-SE did not directly measure concentration dependence, however SEDFIT-MSTAR does provide indications where non-ideality significantly impacts the result. For example, discrepancies between consensus hinge-point and extrapolated $M^*(r)$ values, poor $c(M)$ fits and negatively sloping point-average plots would all indicate the presence of significant levels of non-ideality—however they were not observed in the samples analyzed in this investigation. There would also be a poor correlation between the MHKS value obtained from the literature and these mannans. We can therefore say with confidence that there is a negligible effect of non-ideality present, but the values we report are likely to be slight under-estimates from the true values, although not significantly so.

Conclusions

Four fungal mannan samples were probed for their molar mass and sedimentation coefficients using well-established techniques in the field of polysaccharide characterization.

The two independent analysis techniques used for AUC-SE (SEDFIT-MSTAR and MULTISIG) showed very good agreement. The obvious advantage for MULTISIG was the ability to yield molar mass distributions, particularly insightful for FM-2 and FM-4, as well as reliable values for M_n and M_z , but at a cost of processing time (typically, a set of 20 fits can take between 20 and 30 min). Compared to

this, SEDFIT-MSTAR is a much faster analysis method (a fit taking no more than 10 s) providing reliable and accurate weight and z -average molecular weights but with limited information of molar mass distribution. Combining information from AUC-SE and AUC-SV provided conformation information consistent with previously established results.

These methods (MSTAR, MULTISIG, extended Fujita) have shown a rapid assay for determining the molecular weight distribution of mannans, although they can also be adapted for use with other biopolymer solutions, including polysaccharides, DNA, and proteins—particularly relevant for the characterization of intrinsically disordered proteins. The significance of this assay is the characterization of a polydisperse, heterogeneous biopolymer with significance in various healthcare applications.

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Compliance with ethical standards

Conflict of interest The authors declare no competing financial interests.

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References

- Bradner WT, Clarke DA (1959) Stimulation of host defense against experimental cancer II. Temporal and reversal studies of the Zymosan effect. *Cancer Res* 19:673–678
- Bystrický S, Paulovičová E, Machová E (2003) *Candida albicans* mannan–protein conjugate as vaccine candidate. *Immunol Lett* 85:251–255
- Diller IC (1947) Degenerative changes induced in tumor cells by *Serratia marcescens* polysaccharide. *Cancer Res* 7:605–626
- Ebringerova A, Hromadkova Z, Heinze T (2005) Hemicellulose. *Adv Polym Sci* 186:1–67. doi:10.1007/b136816
- Gillis RB, Adams GG, Heinze T et al (2013) MultiSig: a new high-precision approach to the analysis of complex biomolecular systems. *Eur Biophys J* 42:777–786
- Gray GR, Ballou CE (1971) Isolation and characterization of a polysaccharide containing 3-*O*-methyl-D-mannose from *Mycobacterium phlei*. *J Biol Chem* 246:6835–6842
- Harding SE, Horton JC, Morgan PJ (1992) MSTAR: a FORTRAN program for the model independent molecular weight analysis of macromolecules using low speed or high speed sedimentation equilibrium. In: Harding SE, Rowe AJ, Horton JC (eds) Analytical ultracentrifugation in biochemistry and polymer science. Royal Society of Chemistry, Cambridge, pp 275–294
- Harding SE, Schuck P, Abdelhameed AS et al (2011) Extended Fujita approach to the molecular weight distribution of polysaccharides and other polymeric systems. *Methods* 54:136–144
- Johnston JP, Ogston AG (1946) A boundary anomaly found in the ultracentrifugal sedimentation of mixtures. *Trans Faraday Soc* 42:789–799
- Kabat EA, Bezer AE (1958) The effect of variation in molecular weight on the antigenicity of dextran in man. *Arch Biochem Biophys* 78:306–318
- Kamasuka T, Momoki Y, Sakai S (1968) Antitumor activity of polysaccharide fractions prepared from some strains of Basidiomycetes. *Gan* 59:443
- Kobayashi H, Matsuda K, Ikeda T et al (1994) Structures of cell wall mannans of pathogenic *Candida tropicalis* IFO 0199 and IFO 1647 yeast strains. *Infect Immun* 62:615–622
- Laue T, Shah B, Ridgeway T, Pelletier S (1992) Computer-aided interpretation of sedimentation data for proteins. In: Harding SE, Rowe AJ, Horton JC (eds) Analytical ultracentrifugation in biochemistry and polymer science. Royal Society of Chemistry, Cambridge, pp 90–125
- Ližičárová I, Matulová M, Machová E, Capek P (2005) Mannan from the yeast *Candida dubliniensis*: isolation and structural characterization. In: 13th Eur. Carbohydr. Symp
- Mankowski ZT, Yamashita M, Diller IC (1957) Effect of *Candida guilliermondii* polysaccharide on transplantable mouse sarcoma 37. *Exp Biol Med* 96:79–80
- Mueller A, Raptis J, Rice PJ et al (2000) The influence of glucan polymer structure and solution conformation on binding to (1 → 3)- β -D-glucan receptors in a human monocyte-like cell line. *Glycobiology* 10:339–346
- Nelson RD, Shibata N, Podzorski RP, Herron MJ (1991) *Candida* mannan: chemistry, suppression of cell-mediated immunity, and possible mechanisms of action. *Clin Microbiol Rev* 4:1–19
- Oka S, Kumano N, Sato K et al (1969) Antitumor activity of some plant polysaccharides. II. Chemical constituents and antitumor activity of yeast polysaccharide. *Gann = Gan* 60:287
- Patel TR, Picout DR, Ross-Murphy SB, Harding SE (2006) Pressure cell assisted solution characterization of galactomannans. 3. Application of analytical ultracentrifugation techniques. *Bio-macromolecules* 7:3513–3520
- Patel TR, Morris GA, Garcia de la Torre J et al (2008) Molecular flexibility of methylcelluloses of differing degree of substitution by combined sedimentation and viscosity analysis. *Macromol Biosci* 8:1108–1115. doi:10.1002/mabi.200800064
- Pavlov GM, Korneeva EV, Michailova NA, Ananyeva EP (1992) Hydrodynamic properties of the fractions of mannan formed by *Rhodotorula rubra* yeast. *Carbohydr Polym* 19:243–248
- Peng Y, Zhang L, Zeng F, Xu Y (2003) Structure and antitumor activity of extracellular polysaccharides from mycelium. *Carbohydr Polym* 54:297–303
- Pujol C, Daniels KJ, Lockhart SR et al (2004) The closely related species *Candida albicans* and *Candida dubliniensis* can mate. *Eukaryot Cell* 3:1015–1027
- Rowe AJ (1992) The concentration dependence of sedimentation. In: Harding SE, Rowe AJ, Horton JC (eds) Analytical ultracentrifugation in biochemistry and polymer science. Royal Society of Chemistry, Cambridge, pp 394–406
- Schuck P, Rossmanith P (2000) Determination of the sedimentation coefficient distribution by least-squares boundary modeling. *Biopolymers* 54:328–341
- Schuck P, Gillis RB, Besong TMD et al (2014) SEDFIT–MSTAR: molecular weight and molecular weight distribution analysis of polymers by sedimentation equilibrium in the ultracentrifuge. *Analyst* 139:79–92. doi:10.1039/C3AN01507F
- Shibata N, Arai M, Haga E et al (1992) Structural identification of an epitope of antigenic factor 5 in mannans of *Candida albicans* NIH B-792 (serotype B) and J-1012 (serotype A) as beta-1, 2-linked oligomannosyl residues. *Infect Immun* 60:4100–4110

- Shibata N, Ikuta K, Imai T et al (1995) Existence of branched side chains in the cell wall mannan of pathogenic yeast, *Candida albicans* structure-antigenicity relationship between the cell wall mannans of *Candida albicans* and *Candida parapsilosis*. J Biol Chem 270:1113–1122
- Shibata N, Akagi R, Hosoya T et al (1996) Existence of novel branched side chains containing-1, 2 and-1, 6 linkages corresponding to antigenic factor 9 in the mannan of *Candida guilliermondii*. J Biol Chem 271:9259–9266
- Shibata N, Kobayashi H, Okawa Y, Suzuki S (2003) Existence of novel β -1, 2 linkage-containing side chain in the mannan of *Candida lusitanae*, antigenically related to *Candida albicans* serotype A. Eur J Biochem 270:2565–2575
- Shibata N, Suzuki A, Kobayashi H, Okawa Y (2007) Chemical structure of the cell-wall mannan of *Candida albicans* serotype A and its difference in yeast and hyphal forms. Biochem J 404:365–372
- Sullivan DJ, Westerneng TJ, Haynes KA et al (1995) *Candida dubliniensis* sp. nov.: phenotypic and molecular characterization of a novel species associated with oral candidosis in HIV-infected individuals. Microbiology 141:1507–1521
- Suzuki S, Suzuki M, Hatsukaiwa H et al (1969) Antitumor activities of polysaccharides. III. Growth-inhibitory activity of purified mannan and glucan fractions from baker's yeast against sarcoma-180 solid tumor. Gann 60:273–277
- Suzuki A, Shibata N, Suzuki M et al (1997) Characterization of β -1, 2-mannosyltransferase in *Candida guilliermondii* and its utilization in the synthesis of novel oligosaccharides. J Biol Chem 272:16822–16828
- Tombs M, Harding S (1998) An introduction to polysaccharide biotechnology. Taylor and Francis, London
- Ukai S, Kiho T, Hara C et al (1983) Polysaccharides in Fungi. XIII. Antitumor activity of the various polysaccharides isolated from *Dictyophora indusiata*, *Ganoderma japonicum*, *Cordyceps cicadae*, *Auricularia auricula-judae*, and *Auricularia* species. Chem Pharmaceut Bull 31:741–744