Supplementary Table S1: Analysis of protein N-glycosylation in UGP- and UGP+ cell using lectin affinity chromatography.

			RCA+,c	Complex <sup>b</sup>	Hybrid	Polymannose	[2- <sup>3</sup> H]GP <sup>a</sup>
			% total [2-³H]GP				cpm x 10 <sup>-5</sup>
UGP⁺	Cells		22.7	48.6	11.3	40.1	5.02
	Medium		23.0	83.3	8.6	8.2	1.48
UGP-	Cells		3.5	35.1	13.1	51.8	5.79
	Medium		5.4	70.2	9.6	20.2	1.65
UGP-	Cells	+ Gal	26.2	35.6	18.0	46.3	3.07
	Medium	+ Gal	37.7	65.6	24.2	10.2	1.93

<sup>&</sup>lt;sup>a</sup>Cells were rdiolabelled with [2-<sup>3</sup>H]Man for 18 h as described in Materials and Methods and glycopeptides ([2-<sup>3</sup>H]GP) were prepared from the cells and media.

<sup>&</sup>lt;sup>b</sup>Glycopeptides were subjected to ConA-Sepharose chromatography and eluted as described in Materials and Methods to yield fractions containing glycopeptides possessing complex-, hybrid- and polymannose-type N-glycans.

<sup>&</sup>lt;sup>c</sup>Similarly, glycopeptides were loaded onto RCA-I-agarose columns. After washing the columns, bound glycopeptides (RCA<sup>+</sup>) were eluted as described in Materials and Methods.