

**Supplementary Table 1.** Comparison of patient demographics and disease characteristics between patients in and not in clinical remission at time of therapeutic drug monitoring for vedolizumab. FDR, false discovery rate.

	Clinical Remission (n=57)	Not in Clinical Remission (n=16)	p-value	FDR p-value
Median age, years (IQR)	37.8 (27.3-57.9)	32.8 (27.2-53.4)	0.59	0.79
Male gender, n (%)	32 (56.1)	7 (43.8)	0.38	0.67
Median disease duration, years (IQR)	11.9 (7.0-18.9)	15 (8.8-26.3)	0.31	0.64
Diagnosis of CD, n (%)	31 (54.4)	12 (75.0)	0.50	0.78
Active smoking, n (%)	6 (10.5)	2 (12.5)	0.82	0.96
Previous anti-TNF exposure, n (%)	27 (47.4)	13 (81.3)	<b>0.02</b>	0.14
Concomitant immunomodulator, n (%)	11 (19.3)	4 (25.0)	0.62	0.79
Median duration on vedolizumab, years (IQR)	1.6 (0.8-2.1)	1.4 (0.9-2.5)	0.97	0.97
4-weekly dosing, n (%)	7 (12.3)	7 (43.8)	<b>&lt;0.01</b>	<b>&lt;0.01</b>
Median BMI (IQR)	25.4 (22.7-29.0)	25.7 (18.7-32.8)	0.97	0.97
Median Albumin, g/dL (IQR)	36.0 (34.0-38.0)	34.0 (31.3-36.8)	<b>0.04</b>	0.19
Median CRP, mg/L (IQR)	3.0 (1.0-4.5)	4.0 (1.3-11.0)	0.17	0.60
Median FC, µg/g (IQR)	62.0 (20.0-447.0)	130.0 (55.5-529.3)	0.32	0.64
Biologic remission, n (%)	33 (57.9)	7 (43.8)	0.32	0.64

**Supplementary Table 2.** Comparison of patient demographics and disease characteristics between patients in and not in biologic remission at time of therapeutic drug monitoring for vedolizumab. FDR, false discovery rate.

	<b>Biologic Remission (n=40)</b>	<b>Not in Biologic Remission (n=33)</b>	<b>p-value</b>	<b>FDR p-value</b>
Median age, years (IQR)	34.5 (25.8-49.6)	40.8 (29.0-62.1)	0.32	0.48
Male gender, n (%)	19 (47.5)	20 (60.6)	0.26	0.48
Median disease duration, years (IQR)	12.5 (8.2-20.8)	8.9 (6.4-18.9)	0.44	0.54
Diagnosis of CD, n (%)	22 (55.0)	21 (63.6)	0.45	0.54
Active smoking, n (%)	3 (7.5)	5 (15.2)	0.30	0.48
Previous anti-TNF exposure, n (%)	22 (55.0)	18 (54.5)	0.97	0.97
Concomitant immunomodulator, n (%)	5 (12.5)	10 (30.3)	0.06	0.24
Median duration on vedolizumab, years (IQR)	1.6 (1.0-2.3)	1.3 (0.7-1.9)	0.12	0.36
4-weekly dosing, n (%)	3 (7.5)	11 (33.3)	<0.01	<0.01
Median BMI (IQR)	25.0 (22.5-29.0)	26.2 (22.3-31.7)	0.82	0.90
Median Albumin, g/dL (IQR)	37.5 (35.3-39.0)	35.0 (32.0-36.0)	<0.01	<0.01
Clinical remission, n (%)	33 (82.5)	24 (72.7)	0.32	0.48

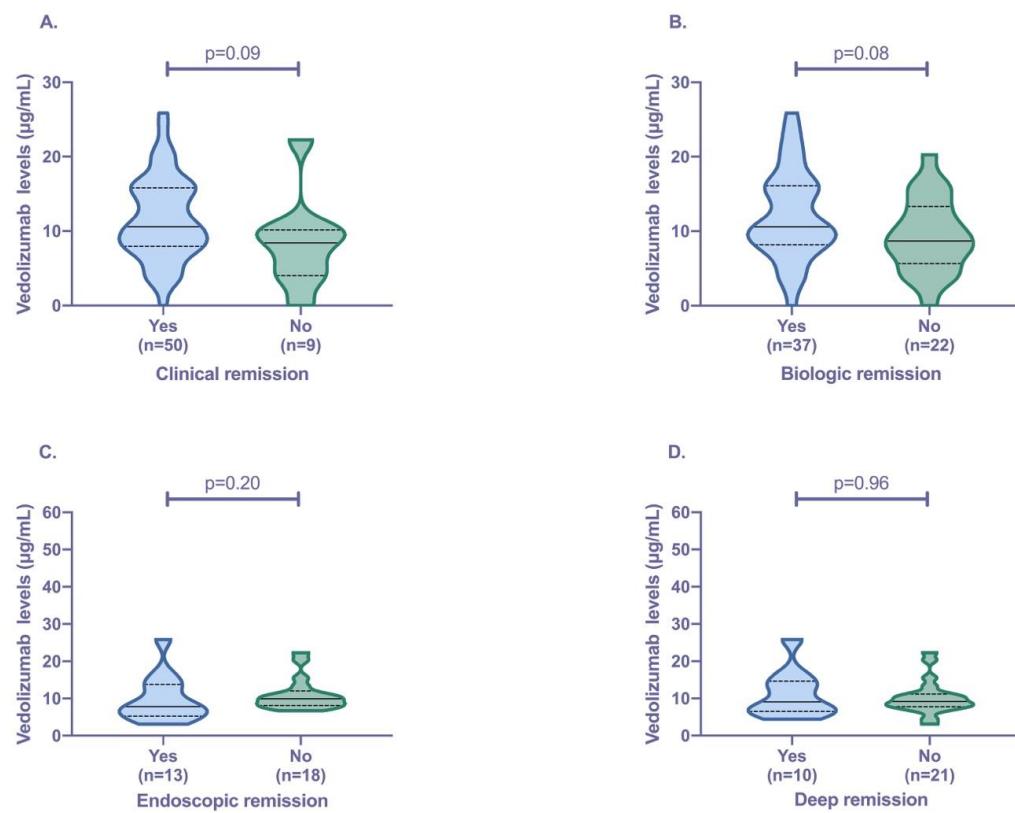
**Supplementary Table 3.** Comparison of patient demographics and disease characteristics between patients in and not in endoscopic remission +/- 8 weeks from therapeutic drug monitoring for vedolizumab. FDR, false discovery rate.

	Endoscopic Remission (n=14)	Not in Endoscopic Remission (n=26)	p-value	FDR p-value
Median age, years (IQR)	43.2 (31.0-60.8)	35.9 (28.7-61.7)	0.66	0.78
Male gender, n (%)	7 (50.0)	13 (50.0)	0.99	0.99
Median disease duration, years (IQR)	15.9 (10.0-19.9)	12.0 (5.9-29.0)	0.43	0.77
Diagnosis of CD, n (%)	6 (42.9)	16 (61.5)	0.26	0.56
Active smoking, n (%)	1 (7.1)	6 (23.1)	0.21	0.56
Previous anti-TNF exposure, n (%)	8 (57.1)	16 (61.5)	0.79	0.86
Concomitant immunomodulator, n (%)	5 (35.7)	5 (19.2)	0.25	0.56
Median duration on vedolizumab, years (IQR)	1.1 (0.6-2.0)	1.4 (0.8-2.3)	0.63	0.78
4-weekly dosing, n (%)	1 (7.1)	8 (30.8)	0.09	0.56
Median BMI (IQR)	26.7 (24.0-40.0)	25.0 (22.2-30.0)	0.13	0.56
Median Albumin, g/dL (IQR)	35.5 (33.8-37.3)	34.5 (32.8-37.0)	0.48	0.77
Clinical remission, n (%)	10 (71.4)	16 (61.5)	0.53	0.77
Biologic remission, n (%)	12 (85.7)	7 (26.9)	<0.01	<0.01

**Supplementary Table 4.** Comparison of patient demographics and disease characteristics between patients in and not in deep remission +/- 8 weeks from therapeutic drug monitoring for vedolizumab. FDR, false discovery rate.

	Deep Remission (n=10)	Not in Deep Remission (n=30)	p-value	FDR p-value
Median age, years (IQR)	52.0 (31.0-67.6)	35.9 (29.0-60.3)	0.38	0.65
Male gender, n (%)	6 (60.0)	7 (23.3)	<b>0.03</b>	0.15
Median disease duration, years (IQR)	16.3 (10.9-19.9)	12.0 (6.7-28.3)	0.49	0.65
Diagnosis of CD, n (%)	3 (30.0)	20 (66.7)	<b>0.04</b>	0.15
Active smoking, n (%)	1 (10.0)	3 (10.0)	0.99	0.99
Previous anti-TNF exposure, n (%)	4 (40.0)	20 (66.7)	0.14	0.34
Concomitant immunomodulator, n (%)	3 (30.0)	7 (23.3)	0.67	0.80
Median duration on vedolizumab, years (IQR)	1.3 (0.6-2.0)	1.3 (0.7-2.3)	0.94	0.99
4-weekly dosing, n (%)	0	9 (30.0)	<b>0.049</b>	0.15
Median BMI (IQR)	25.8 (23.1-41.4)	25.1 (22.4-30.7)	0.48	0.65
Median Albumin, g/dL (IQR)	35.0 (33.8-38.0)	35.0 (32.8-36.3)	0.38	0.65
Biologic remission, n (%)	9 (90.0)	10 (33.3)	<b>&lt;0.01</b>	<b>0.02</b>

**Supplementary Figure 1.** Association of trough vedolizumab levels with (A) clinical remission; (B) biologic remission; (C) endoscopic remission; and (D) deep remission after omitting patients on 4-weekly dosing. Violin plots show median (solid line), interquartile range (dotted line), maximum and minimum.



**Supplementary Figure 2.** Association of trough vedolizumab levels with (A) biologic remission defined as CRP< 5mg/L plus faecal calprotectin <150 µg/g and (B) biologic remission defined as CRP< 5mg/L plus faecal calprotectin <50 µg/g. Violin plots show median (solid line), interquartile range (dotted line), maximum and minimum.

