Chemostratigraphy and provenance of clays and other non-carbonate minerals in chalks of Campanian age (Upper Cretaceous) from Sussex, southern England

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ABSTRACT: Geochemical analysis of acid-insoluble residues derived from white chalks and marl seams of Campanian age from Sussex, UK, has been undertaken. All display a broadly similar <2 μm mineralogical composition consisting of smectite or smectite-rich illite-smectite with subordinate illite and minor amounts of talc. Plots of K_2O/Al_2O_3 and TiO_2/Al_2O_3 indicate that most marl seams have an acid-insoluble residue composition which is slightly different to that of the over- and underlying white chalk, implying that marl seams are primary sedimentary features not formed through white chalk dissolution. On the basis of a negative Eu anomaly and trace element geochemistry one marl seam, the Old Nore Marl, is considered to be volcanically derived and best classified as a bentonite; it is considered to correlate with the bentonite M1 of the north German succession.

The aim of this paper is to investigate the geochemistry of the non-carbonate fraction of marly and white chalks as found at Peacehaven Steps on the Sussex coast. The data will be used to examine the previously published interpretation of the Old Nore Marl as representing a bentonite (Mortimore *et al.*, 2001; p. 214). It will also discuss the origin of clay-rich beds higher in the succession and the non-carbonate fraction of the interbedded white chalks.

The Peacehaven Steps succession is of Campanian age (*O. pilula* to *G quadrata* zones). It is a well known locality on the Sussex coast, in part because the steps allow a rapid traverse of the chalk sequence whose bedding is near-horizontal at this point. The stratigraphy of the succession and its relationship to the remainder of the Anglo-Paris Basin is well documented (Mortimore, 1986; Mortimore *et al.*, 2001). The early Campanian is distinctive in that chalks of this age in southern England contain a number of laterally persistent more clay-rich beds, often termed "marl seams" which are more abundant at this level than in the underlying beds of Santonian age. The origin and correlation of these Campanian marls was first discussed in a series of papers by Brydone (Brydone, 1914a,b) who recognized the widespread lateral continuity of individual marl seams and argued, based on their lateral persistence, that they represented altered volcanic ashfalls.

TABLE 1. Geochemical data used in this study; all data are derived from acid-insoluble residues.

	SiO_2	TiO_2	Al_2O_3	$Fe_2O_3(t)$	MnO	MgO	CaO	Na_2O	K_2O	P_2O_5
	Oxide wt.%									
PHS 65	48.77	0.63	14.40	11.12	0.026	2.81	5.76	0.13	2.31	2.17
PHS 64	56.49	0.62	15.20	4.18	0.008	3.13	4.56	0.12	2.47	1.31
PHS 63	55.70	0.61	15.22	4.55	0.012	2.99	4.47	0.12	2.43	1.57
PHS 62	55.81	0.65	16.31	4.08	0.464	3.21	5.78	0.13	2.45	2.04
PHS 61	54.70	0.67	16.80	8.16	0.330	3.25	4.20	0.12	2.47	1.57
PHS 60	54.74	0.67	16.91	4.17	0.007	3.31	5.51	0.14	2.59	1.92
PHS 59	55.87	0.67	17.20	4.98	0.011	3.46	4.43	0.13	2.46	1.41
PHS 58	55.43	0.72	16.69	3.76	0.007	3.33	6.78	0.17	2.52	2.83
PHS 57	54.04	0.64	17.01	3.86	0.006	3.85	7.14	0.12	1.77	2.75
PHS 56	50.66	0.64	14.96	4.25	0.007	3.16	8.66	0.17	2.44	3.86
PHS 55	50.48	0.67	15.38	7.35	0.011	3.24	6.16	0.14	2.49	2.62
PHS 54	54.81	0.67	17.28	3.90	0.006	3.82	5.90	0.12	2.09	2.10
PHS 53	51.14	0.64	15.04	3.61	0.007	3.17	7.64	0.16	2.34	3.06
PHS 52	57.22	0.72	17.49	4.67	0.007	3.36	5.07	0.15	2.69	2.23
PHS 51	57.53	0.72	17.49	4.91	0.007	3.31	4.72	0.14	2.74	2.08
PHS 50	60.26	0.72	18.44	3.94	0.006	3.44	2.63	0.12	2.78	0.65
PHS 49	57.96	0.66	16.93	4.61	0.006	3.34	4.28	0.13	2.75	1.80
PHS 48	57.43	0.66	16.82	4.70	0.006	3.29	4.60	0.14	2.66	1.98
PHS 47	57.48	0.67	17.32	4.83	0.007	3.41	4.19	0.14	2.73	1.76
PHS 46	59.14	0.67	18.03	5.19	0.007	3.50	3.22	0.12	2.71	1.05
PHS 45	61.23	0.67	19.11	3.92	0.006	3.59	2.33	0.11	2.66	0.30
PHS 44	61.46	0.63	19.31	4.34	0.007	3.64	2.83	0.10	2.49	0.66
PHS 43	58.59	0.67	18.28	3.57	0.007	3.40	5.03	0.15	2.63	2.15
PHS 42	61.20	0.75	18.96	3.87	0.006	3.54	2.66	0.13	2.88	0.67
PHS 41	56.20	0.64	16.68	4.34	0.007	3.30	5.72	0.16	2.50	2.78
PHS 40	54.57	0.63	16.28	4.29	0.006	3.17	6.60	0.17	2.43	3.36
PHS 39	54.77	0.62	16.26	5.19	0.006	3.24	5.11	0.15	2.51	2.42
PHS 38	55.17	0.64	16.45	5.53	0.007	3.24	4.93	0.14	2.54	2.30
PHS 37	57.37	0.64	17.16	3.71	0.006	3.44	4.42	0.14	2.55	1.80
PHS 36	56.44	0.63	17.30	3.78	0.007	3.53	5.10	0.14	2.51	2.24
PHS 35	57.41	0.65	18.17	4.72	0.007	3.52	4.69	0.13	2.52	1.95
PHS 34	56.35	0.60	17.15	4.59	0.008	3.55	5.44	0.14	2.54	2.50
PHS 33	57.32	0.64	17.44	3.92	0.006	3.41	5.24	0.15	2.55	2.29
PHS 32	57.36	0.70	17.55	4.81	0.006	3.48	3.91	0.13	2.68	1.51
PHS 31	58.82	0.68	18.11	4.19	0.006	3.42	3.69	0.13	2.66	1.29
PHS 30	57.06	0.65	16.80	5.58	0.006	3.28	3.75	0.12	2.51	1.45
PHS 29	58.23	0.63	17.17	3.54	0.006	3.39	4.67	0.15	2.45	1.95
PHS 28	53.97	0.58	15.20	4.95	0.006	3.04	6.46	0.16	2.27	3.34
PHS 27	55.17	0.61	15.90	5.35	0.006	3.15	5.12	0.14	2.46	2.46
PHS 26	57.08	0.64	16.83	5.39	0.006	3.34	3.61	0.13	2.57	1.41
PHS 25	54.94	0.62	16.45	4.40	0.006	3.23	5.84	0.15	2.41	2.81
PHS 24	59.31	0.67	17.43	4.00	0.007	3.77	2.54	0.11	2.71	0.59
PHS 23	61.12	0.68	18.14	4.44	0.005	3.55	2.12	0.10	2.57	0.24
PHS 22	57.78	0.64	17.18	3.74	0.006	3.44	4.37	0.13	2.49	1.79
PHS 21	59.05	0.66	17.61	4.92	0.006	3.58	2.99	0.12	2.52	0.92
PHS 20	64.39	0.55	14.60	3.65	0.005	2.83	3.64	0.12	2.17	1.49
PHS 19	54.40	0.59	16.14	3.86	0.006	3.20	6.65	0.16	2.32	3.28
PHS 18	56.28	0.61	16.76	4.85	0.006	3.30	4.75	0.14	2.40	2.04
PHS 17	58.15	0.66	17.24	4.54	0.007	3.53	3.84	0.13	2.63	1.48
PHS 16	60.23	0.69	18.11	3.82	0.006	3.62	3.71	0.14	2.68	1.29
PHS 15	63.22	0.62	18.15	3.53	0.005	3.85	2.49	0.11	2.37	0.49
PHS 14	59.32	0.65	18.19	5.48	0.005	3.36	2.72	0.11	2.43	0.69
PHS 13	59.26	0.67	18.00	4.13	0.006	3.32	3.55	0.13	2.60	1.23
PHS 12	55.39	0.59	15.22	3.55	0.006	2.97	7.68	0.18	2.27	3.91
PHS 11	58.44	0.64	16.15	4.39	0.005	3.13	4.51	0.14	2.38	1.93
PHS 10	64.81	0.52	12.83	3.66	0.005	2.50	5.25	0.14	1.86	2.55
PHS 9	60.18	0.66	17.92	3.63	0.006	3.48	3.79	0.13	2.37	1.30

TABLE 1 (*contd.*).

	SiO$_2$	TiO$_2$	Al$_2$O$_3$	Fe$_2$O$_3$(t)	MnO	MgO	CaO	Na$_2$O	K$_2$O	P$_2$O$_5$
	Oxide wt.%									
PHS 8	52.42	0.58	15.66	3.95	0.006	3.01	8.24	0.19	2.17	4.31
PHS 7	60.29	0.69	18.17	3.90	0.006	3.46	3.18	0.12	2.51	0.91
PHS 6	54.45	0.60	15.58	5.42	0.006	2.96	6.51	0.17	2.27	3.24
PHS 5	60.54	0.70	17.30	3.33	0.006	3.19	3.27	0.13	2.56	0.97
PHS 4	55.97	0.56	14.31	4.96	0.006	2.66	6.36	0.16	2.12	3.20
PHS 3	57.99	0.62	15.58	3.64	0.006	3.07	4.75	0.16	2.27	1.95
PHS 2	60.10	0.60	17.38	3.82	0.006	4.00	4.23	0.15	1.87	1.35
PHS 1	57.94	0.67	17.21	4.06	0.006	3.25	3.67	0.13	2.47	1.18

	Ba	Cr	Cs	Hf	Nb	Pb	Rb	Sc	Sr	Ta	Th	U	V	Y	Zr
	mg kg^{-1}														
PHS 65	173	65.6	7.0	3.9	13.5	35.8	86	10.8	172	1.02	10.5	10.3	282	112	139
PHS 64	183	57.6	7.6	4.0	14.0	21.7	91	9.5	138	1.03	8.9	6.3	211	108	135
PHS 63	192	59.8	7.8	3.7	13.6	26.9	94	9.6	148	1.04	9.4	8.5	221	106	126
PHS 62	205	71.1	8.3	4.1	14.8	31.0	101	9.9	182	1.11	11.1	7.0	218	106	142
PHS 61	218	74.7	8.3	4.3	15.0	43.7	103	10.8	142	1.11	11.9	5.7	227	85	146
PHS 60	215	72.8	8.4	4.4	17.0	22.8	103	10.2	181	1.22	11.1	9.2	228	126	148
PHS 59	196	70.4	8.1	6.0	25.1	31.6	99	9.7	130	2.37	12.5	7.5	268	104	190
PHS 58	190	69.2	8.2	4.9	19.8	23.5	98	10.1	242	1.51	11.6	9.0	231	136	155
PHS 57	135	56.1	5.4	4.7	14.1	34.1	67	9.1	209	1.05	10.7	7.5	211	181	148
PHS 56	173	61.2	7.9	4.3	14.8	31.9	93	10.1	305	1.06	10.6	9.4	202	165	144
PHS 55	177	64.7	8.1	4.2	14.8	38.8	97	9.7	214	1.03	10.2	7.9	209	138	142
PHS 54	145	54.2	5.9	4.1	12.8	22.7	77	8.6	186	0.91	9.6	6.0	178	114	134
PHS 53	184	61.6	7.5	3.9	13.4	19.9	92	10.1	262	0.97	10.7	11.8	211	150	140
PHS 52	200	70.0	8.1	4.1	14.4	20.5	106	11.4	196	1.04	10.1	6.1	190	106	144
PHS 51	198	67.8	7.6	3.8	14.2	24.0	105	11.3	179	1.02	10.5	9.7	227	145	135
PHS 50	218	65.2	8.1	3.8	14.8	19.5	107	11.6	85	1.08	10.3	5.2	208	110	131
PHS 49	201	67.4	8.0	11.8	13.8	18.5	104	10.8	158	0.99	9.6	7.0	203	126	457
PHS 48	195	65.9	7.6	3.6	13.6	21.7	99	10.3	175	1.01	9.2	8.6	198	136	128
PHS 47	205	69.7	8.0	3.5	13.7	20.8	105	10.9	167	1.01	9.7	6.5	199	108	128
PHS 46	218	72.6	8.2	4.1	14.0	23.5	108	10.6	119	1.07	10.5	6.0	216	95	144
PHS 45	221	78.3	8.0	3.4	13.8	18.2	110	11.1	75	1.07	10.9	3.0	193	55	111
PHS 44	203	79.0	7.6	3.1	13.3	25.9	105	10.2	96	1.03	11.6	3.4	183	57	103
PHS 43	218	65.7	7.5	3.4	14.0	18.0	105	11.4	190	1.06	11.2	7.5	188	135	119
PHS 42	252	79.2	8.2	3.8	15.2	18.5	114	12.0	94	1.14	11.1	5.7	309	72	130
PHS 41	195	64.1	7.3	3.9	13.5	22.3	96	10.4	221	1.00	9.5	8.4	178	156	142
PHS 40	194	64.2	7.3	3.6	13.6	21.8	95	10.7	264	1.02	9.7	9.2	196	157	133
PHS 39	184	66.1	7.4	3.8	13.7	21.7	95	10.6	233	1.04	8.9	11.8	231	183	139
PHS 38	181	66.4	7.4	3.6	13.7	22.6	96	10.6	196	1.01	9.6	9.9	219	164	131
PHS 37	188	70.3	7.5	3.5	13.7	15.4	97	10.9	174	1.08	10.0	9.9	224	136	124
PHS 36	183	66.5	7.3	3.2	14.0	14.0	97	10.7	218	1.09	9.8	7.9	185	131	114
PHS 35	194	67.0	7.5	3.3	14.0	21.1	101	10.9	188	1.09	10.0	5.1	182	102	115
PHS 34	190	70.7	7.6	3.3	12.7	17.4	100	10.4	220	0.94	9.2	6.9	205	116	115
PHS 33	209	71.2	7.6	3.4	13.8	18.9	101	11.2	194	1.04	10.2	7.2	191	132	122
PHS 32	211	74.1	7.7	3.7	15.1	21.1	106	11.3	163	1.18	10.6	7.3	214	126	129
PHS 31	218	75.9	7.8	3.5	13.8	17.6	103	11.2	134	1.04	10.0	6.4	207	108	124
PHS 30	196	70.8	7.5	3.5	13.4	34.5	99	10.4	126	1.02	9.4	6.5	202	109	124
PHS 29	195	68.9	7.4	3.4	12.7	15.6	95	10.8	184	0.97	9.4	5.6	174	122	122
PHS 28	175	62.2	6.8	3.4	12.2	23.8	88	9.9	254	0.95	9.0	12.2	213	179	124
PHS 27	186	68.7	7.5	3.3	12.7	26.7	97	11.3	223	0.96	9.7	10.5	220	175	121
PHS 26	207	70.3	7.9	3.4	13.0	21.1	104	10.8	141	0.98	9.7	5.8	203	120	123
PHS 25	199	65.0	7.3	3.3	12.8	19.2	97	10.6	241	0.96	9.6	9.0	201	144	118
PHS 24	229	76.3	7.8	3.4	13.7	14.2	118	10.9	86	1.00	9.3	4.5	195	76	115
PHS 23	212	77.2	7.9	3.4	14.0	18.6	115	10.8	50	1.06	10.2	6.5	220	96	120
PHS 22	212	81.5	8.0	3.3	13.6	15.0	112	11.5	177	1.02	10.6	8.1	214	123	116
PHS 21	221	78.8	8.2	3.5	14.0	21.0	108	12.0	115	1.04	10.7	7.8	233	119	126

TABLE 1. (*contd.*).

	Ba	Cr	Cs	Hf	Nb	Pb	Rb	Sc	Sr	Ta	Th	U	V	Y	Zr
	mg kg^{-1}														
PHS 20	178	59.2	6.7	2.7	11.1	17.4	88	9.5	134	0.81	8.1	7.0	174	100	99
PHS 19	185	64.1	7.5	3.0	12.2	18.9	95	10.4	255	0.92	9.0	11.2	196	144	106
PHS 18	190	75.5	7.9	3.3	13.0	22.0	98	10.2	169	0.98	8.9	9.5	208	139	121
PHS 17	204	76.5	8.5	3.4	14.1	16.9	106	11.5	135	0.99	9.6	5.1	201	95	123
PHS 16	219	89.7	8.7	3.6	14.7	14.3	111	12.6	128	1.04	11.0	6.6	237	108	132
PHS 15	190	84.5	7.8	3.2	13.5	13.0	99	10.3	80	0.96	10.2	4.2	199	64	115
PHS 14	206	61.9	7.4	3.2	14.8	21.2	98	10.5	75	0.98	9.1	5.9	200	83	119
PHS 13	239	70.5	7.6	3.4	15.1	15.5	104	11.1	120	0.99	9.6	7.0	201	92	125
PHS 12	179	63.1	6.9	1.3	13.9	17.4	89	9.6	292	0.91	8.6	16.4	220	155	63
PHS 11	198	72.4	7.5	3.0	15.7	21.6	98	10.2	165	0.99	9.3	10.3	243	118	128
PHS 10	157	64.5	6.0	2.8	12.8	15.2	76	9.5	204	0.82	8.5	8.1	181	111	114
PHS 9	207	95.3	8.0	3.5	17.2	15.8	103	10.9	137	1.04	11.0	4.5	229	70	139
PHS 8	176	68.2	6.8	1.2	15.5	18.1	89	11.6	324	0.93	10.0	11.1	187	140	66
PHS 7	215	87.5	8.1	3.8	19.4	17.9	108	10.6	110	1.11	11.2	3.7	232	54	155
PHS 6	186	71.4	7.4	1.4	20.9	30.0	96	11.1	263	1.04	10.5	9.8	205	134	79
PHS 5	222	78.3	8.5	4.2	23.2	12.2	109	10.9	104	1.21	10.8	3.9	199	71	166
PHS 4	175	55.4	6.4	1.0	25.0	27.5	83	9.3	243	1.16	9.8	11.5	179	172	57
PHS 3	173	63.7	7.0	8.7	48.0	18.0	92	10.0	175	4.20	11.6	7.1	201	138	238
PHS 2	132	66.9	4.3	16.8	47.9	32.3	68	7.0	134	8.93	24.8	4.3	186	82	385
PHS 1	206	73.0	7.9	6.9	34.6	15.2	104	10.6	122	2.08	11.0	5.1	172	114	216

	La	Ce	Pr	Nd	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu
	mg kg^{-1}													
PHS 65	85.7	68.5	16.9	68.7	12.6	2.97	14.0	1.95	11.2	2.39	5.70	0.70	3.48	0.47
PHS 64	83.5	66.1	16.0	65.3	11.9	2.85	13.5	1.87	10.9	2.28	5.33	0.65	3.19	0.43
PHS 63	79.5	65.6	15.4	62.7	11.5	2.74	12.9	1.80	10.5	2.23	5.22	0.64	3.10	0.42
PHS 62	82.8	72.0	16.5	67.1	12.5	2.90	13.5	1.90	10.8	2.29	5.41	0.66	3.29	0.45
PHS 61	73.8	68.2	14.9	59.0	11.0	2.45	11.4	1.61	9.2	1.92	4.62	0.59	3.08	0.42
PHS 60	95.1	80.7	18.9	76.7	14.3	3.37	15.8	2.24	12.7	2.68	6.28	0.76	3.74	0.51
PHS 59	82.6	74.5	16.6	66.5	12.5	2.73	13.5	1.90	10.8	2.28	5.37	0.68	3.42	0.48
PHS 58	100.6	84.1	19.8	81.2	15.1	3.51	16.8	2.34	13.5	2.85	6.72	0.81	3.93	0.53
PHS 57	145.8	119.6	28.3	110.6	19.8	4.51	21.5	3.07	17.6	3.68	8.65	1.04	4.93	0.64
PHS 56	111.8	87.9	21.7	89.0	16.7	3.95	19.2	2.68	15.5	3.28	7.68	0.92	4.37	0.59
PHS 55	93.2	75.9	18.1	74.3	14.0	3.33	15.7	2.21	12.8	2.73	6.46	0.78	3.79	0.51
PHS 54	77.8	66.4	15.5	63.4	11.9	2.85	13.5	1.89	10.8	2.28	5.41	0.64	3.08	0.42
PHS 53	95.1	78.1	18.2	74.8	14.0	3.37	16.2	2.29	13.5	2.89	6.84	0.80	3.85	0.52
PHS 52	76.3	67.8	15.1	62.7	11.6	2.77	12.7	1.79	10.5	2.23	5.30	0.64	3.24	0.45
PHS 51	106.1	90.4	21.2	88.7	16.7	3.93	18.2	2.54	14.5	3.04	6.95	0.82	3.87	0.52
PHS 50	82.8	73.8	16.5	67.0	12.3	2.86	13.3	1.90	11.1	2.34	5.59	0.69	3.53	0.48
PHS 49	91.2	74.6	17.8	74.6	13.9	3.29	15.3	2.13	12.3	2.57	6.00	0.72	3.45	0.47
PHS 48	96.7	77.2	18.4	76.0	14.3	3.32	16.1	2.23	13.0	2.76	6.48	0.76	3.69	0.49
PHS 47	82.0	71.7	16.2	67.1	12.6	2.94	13.8	1.87	10.8	2.27	5.37	0.65	3.20	0.43
PHS 46	75.8	71.6	15.1	61.8	11.7	2.70	12.4	1.74	9.9	2.07	4.89	0.61	3.04	0.41
PHS 45	52.5	56.8	10.7	41.9	7.8	1.72	7.7	1.11	6.3	1.30	3.20	0.43	2.28	0.32
PHS 44	55.9	60.1	11.6	46.0	8.6	1.88	8.5	1.20	6.7	1.38	3.31	0.42	2.24	0.31
PHS 43	104.0	91.2	19.5	80.4	15.1	3.47	16.3	2.32	13.6	2.87	6.75	0.83	4.03	0.53
PHS 42	59.9	58.5	11.9	47.6	8.9	1.96	9.2	1.31	7.6	1.63	3.96	0.53	2.76	0.38
PHS 41	107.3	84.3	20.5	85.7	16.0	3.76	18.1	2.54	14.8	3.18	7.47	0.87	4.15	0.57
PHS 40	106.5	82.5	20.5	85.3	15.8	3.78	18.0	2.51	14.8	3.16	7.53	0.89	4.24	0.57
PHS 39	120.4	88.2	22.2	92.6	17.3	4.14	19.9	2.82	16.6	3.59	8.31	0.98	4.52	0.61
PHS 38	108.2	82.4	20.4	85.5	15.7	3.73	18.2	2.59	15.2	3.29	7.73	0.92	4.31	0.58
PHS 37	94.2	75.4	18.0	75.2	14.1	3.29	16.0	2.23	13.0	2.79	6.57	0.78	3.82	0.51
PHS 36	92.2	76.9	17.5	72.6	13.6	3.20	15.3	2.14	12.5	2.67	6.34	0.76	3.68	0.48
PHS 35	75.4	66.9	14.8	61.6	11.6	2.69	12.6	1.74	10.2	2.15	5.07	0.61	3.10	0.41
PHS 34	82.4	69.4	15.8	65.9	12.3	2.92	13.8	1.91	11.1	2.35	5.52	0.66	3.25	0.43
PHS 33	96.4	81.5	18.9	78.3	14.6	3.42	16.4	2.28	13.1	2.75	6.44	0.77	3.71	0.50

Table 1. (*contd.*).

	La	Ce	Pr	Nd	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu
	mg kg^{-1}													
PHS 32	95.5	79.2	18.6	76.9	14.3	3.31	15.7	2.19	12.8	2.66	6.24	0.76	3.73	0.50
PHS 31	79.5	68.7	15.5	64.4	12.1	2.82	13.3	1.87	10.9	2.31	5.45	0.67	3.32	0.46
PHS 30	79.5	66.7	15.3	64.0	11.9	2.85	13.3	1.87	10.8	2.29	5.45	0.67	3.25	0.45
PHS 29	87.2	71.3	17.2	71.7	13.3	3.15	14.9	2.06	12.2	2.55	6.08	0.73	3.61	0.49
PHS 28	116.4	86.8	21.4	91.1	17.0	4.09	19.7	2.80	16.8	3.62	8.47	0.99	4.67	0.63
PHS 27	119.5	89.3	22.2	92.8	17.3	4.15	19.7	2.81	16.6	3.56	8.47	1.02	4.84	0.64
PHS 26	84.8	71.8	16.7	68.9	13.0	3.03	14.3	2.05	12.0	2.53	5.98	0.73	3.60	0.49
PHS 25	99.1	81.1	19.0	79.1	15.0	3.58	16.9	2.40	14.0	2.97	6.94	0.84	4.07	0.54
PHS 24	61.7	57.8	12.3	50.3	9.4	2.16	10.0	1.41	8.1	1.69	4.06	0.54	2.69	0.38
PHS 23	72.7	66.5	14.7	60.2	11.3	2.67	12.2	1.74	10.2	2.15	5.10	0.64	3.27	0.44
PHS 22	94.6	84.3	19.3	79.5	14.9	3.52	16.1	2.27	13.0	2.71	6.33	0.77	3.76	0.50
PHS 21	85.7	74.9	17.0	70.1	13.1	3.13	14.5	2.09	12.2	2.56	6.05	0.76	3.75	0.50
PHS 20	72.8	61.9	14.5	59.8	11.2	2.61	12.4	1.75	10.1	2.16	5.13	0.63	3.13	0.44
PHS 19	94.8	74.6	17.5	74.0	14.1	3.37	16.0	2.26	13.5	2.90	6.89	0.80	3.85	0.52
PHS 18	93.5	75.1	17.8	74.3	13.8	3.39	16.0	2.23	13.3	2.83	6.58	0.79	3.75	0.50
PHS 17	70.6	62.8	14.1	58.2	10.8	2.55	11.9	1.65	9.7	2.07	5.00	0.62	3.10	0.44
PHS 16	81.1	73.3	16.3	66.2	12.3	2.85	13.3	1.91	11.1	2.35	5.70	0.72	3.64	0.50
PHS 15	53.2	55.3	11.2	45.3	8.4	1.91	8.6	1.24	7.1	1.50	3.62	0.45	2.38	0.33
PHS 14	61.1	58.8	12.6	51.3	9.6	2.28	10.4	1.48	8.6	1.82	4.41	0.54	2.82	0.38
PHS 13	69.5	64.6	14.1	58.2	11.0	2.63	12.1	1.71	9.7	2.05	4.70	0.57	2.86	0.39
PHS 12	101.8	81.2	19.5	81.4	15.3	3.74	17.8	2.50	14.7	3.17	7.32	0.87	4.14	0.55
PHS 11	82.8	70.4	16.2	68.2	12.8	3.04	14.3	2.01	12.0	2.55	6.03	0.72	3.55	0.49
PHS 10	79.1	67.3	15.3	63.6	11.9	2.87	13.3	1.89	11.1	2.37	5.61	0.67	3.32	0.44
PHS 9	60.2	60.4	12.3	49.0	9.1	2.01	9.3	1.34	7.8	1.65	3.95	0.53	2.81	0.39
PHS 8	95.5	82.8	18.1	75.4	14.3	3.41	16.3	2.34	13.7	2.93	6.87	0.81	3.94	0.53
PHS 7	45.6	51.0	9.5	37.8	7.1	1.58	7.1	1.03	6.1	1.27	3.17	0.44	2.35	0.34
PHS 6	91.2	78.0	17.4	72.8	13.5	3.18	15.5	2.24	13.1	2.86	6.67	0.81	3.93	0.53
PHS 5	53.5	54.7	10.9	43.9	8.2	1.86	8.8	1.26	7.5	1.63	3.98	0.52	2.78	0.40
PHS 4	108.9	89.8	20.8	88.0	16.5	3.83	19.4	2.79	16.4	3.59	8.26	0.97	4.52	0.60
PHS 3	102.0	102.4	21.4	88.1	17.1	3.41	18.3	2.67	15.3	3.16	7.30	0.90	4.31	0.58
PHS 2	69.9	101.4	18.1	73.2	14.7	2.23	13.7	2.00	11.0	2.16	4.94	0.61	3.10	0.42
PHS 1	79.3	77.4	15.9	66.1	12.7	2.81	14.2	2.04	12.0	2.53	5.96	0.72	3.61	0.50

In the white chalk succession of the Anglo-Paris Basin and eastern England marl seams are most abundant in beds of Turonian and early Coniacian age and here they have been extensively studied (Pacey, 1984; Deconinck & Chamley, 1995; Wray & Wood, 1998; Wray, 1999). Previous work has demonstrated that Turonian−early Coniacian marl seams have either a detrital origin, or are bentonites derived from the alteration of distal volcanic ash (Wray, 1999). This work built on earlier mineralogical studies on Turonian successions in the Lower Saxony area of Germany where bentonites were first described (Dorn & Bräutigam, 1959; Seibertz & Vortisch, 1979; Ernst *et al.*, 1983; Wray, 1995; Wray *et al.*, 1995; Wray & Wood, 1995; Wray *et al.*, 1996; Wray & Wood, 2002). Wray (1999) demonstrated that it is possible to correlate all the Turonian marl seams of volcanogenic origin in the Anglo-Paris Basin with ones in the

successions of both eastern England and northern Germany, facilitating a field applicable chronostratigraphic framework for the Turonian and Coniacian Stages. In contrast, whilst it is possible to correlate detrital marl seams within a region, it has not been possible to confidently correlate them between regions.

Although data have been generated from younger marl seams by Wray, none of this information has been formally published; instead, specific pieces of information have entered the literature in the form of personal communications (e.g. Mortimore *et al.*, 2001; p. 214) which is somewhat unsatisfactory. One aim of this piece of work is to formally publish the underlying evidence for some of these statements.

White chalks of lower Campanian age are predominantly composed of calcium carbonate and usually only comprise at most 1−2% non-carbonate
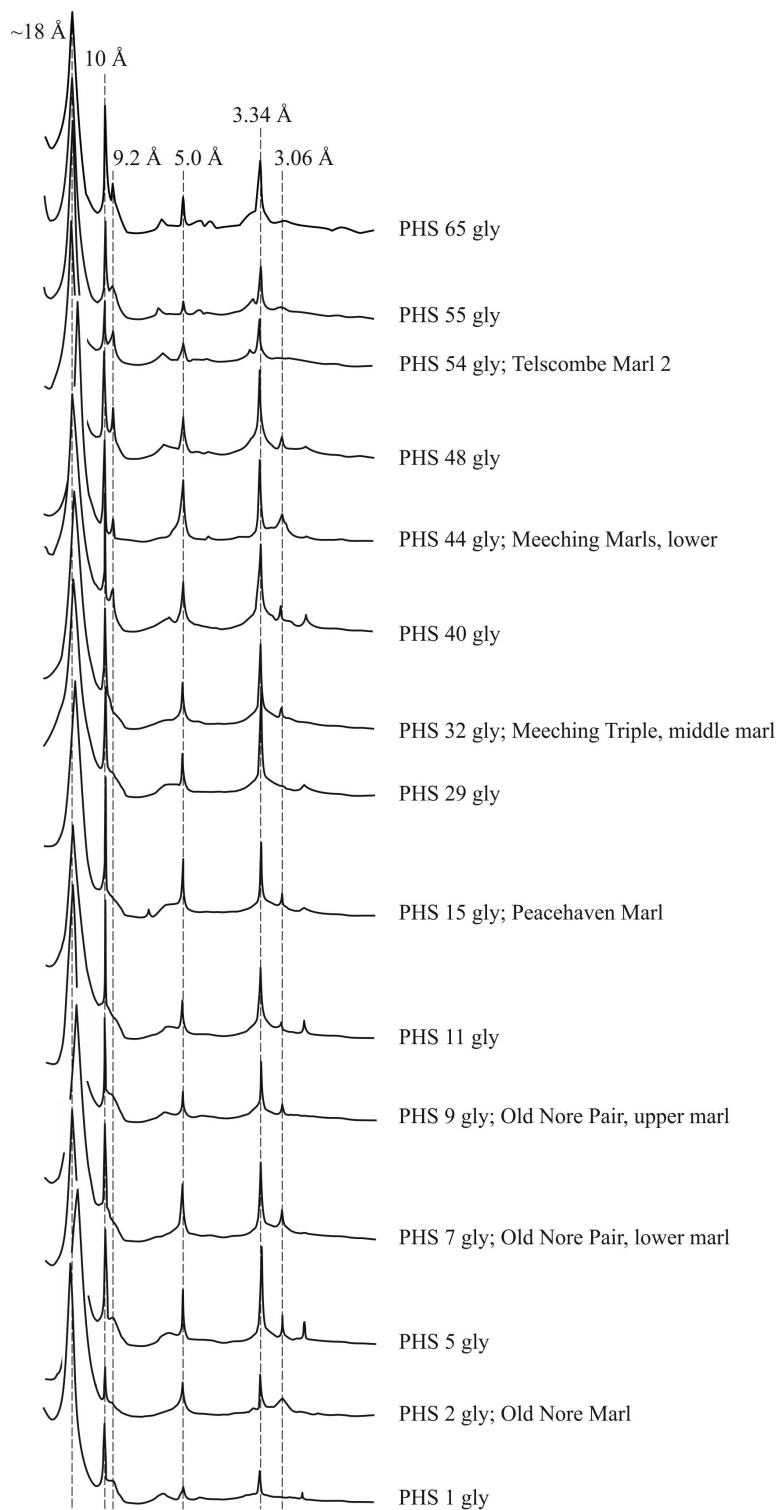
minerals; even many of the marl seams only contain in the region of 3−10% non-carbonate minerals. Such a low proportion of non-carbonate material tests the limits of detection of even the most modern analytical systems and hence we elected to dissolve away the carbonate fraction of all samples and subject the residual non-carbonate fraction to geochemical testing. This approach enables the direct comparison of the non-carbonate fraction present in different lithologies without the need to compensate for variable carbonate content.

Previous work has investigated the non-carbonate mineralogy of white chalks of comparable age from elsewhere in Sussex (Weir & Catt, 1965; Perrin, 1971; Jeans, 2006). All samples were found to have a non-carbonate fraction dominated by smectite with subordinate illite. Also reported by Weir and Catt was the presence of euhedral alkali feldspar crystals in the fine silt fraction which were interpreted as being of authigenic origin. The samples examined in the present study have undergone mineralogical examination and some aspects of the data are presented in Jeans *et al.* (2014); additional data are presented below. Jeans *et* al. (2014) reported a similar mineralogy to that described by Weir and Catt and Perrin with a predominance of smectite or smectite-rich illite/ smectite but they also noted the presence of minor amounts of talc along with illite. Jeans *et al.* (2014) also reported the presence of minute euhedral alkali feldspar crystals in a small number of white chalk samples located between the Peacehaven and Meeching Triple Marls.

METHODOLOGY

Acid-insoluble residues of 65 samples were obtained by dissolving roughly crushed samples of chalk and marls in cold 1m acetic acid followed by washing of the residue with deionized water. Residues were ground in an agate ball mill and dried at 105ºC. All powders were then dissolved using a lithium metaborate fusion by the method described in Wray & Wood (1998). Quantification of major and abundant trace metals was achieved using a Thermo iCAP 6500 inductively coupled plasma-optical emission spectrometer; quantification of less abundant trace metals and the rare-earth elements (*REE*) was achieved using a Thermo X Series inductively coupled plasma-mass spectrometer. Calibration for both instruments was carried out using matrix matched multi-element standards constructed from traceable single element standards solutions. The laboratory in which this work was performed is accredited to ISO 17025 for this test (Testing Lab 2180). Five certified reference materials were tested in the same units of work as the samples and one of these reference materials, USGS Cody Shale (SCo-1), was used as the shale normalising standard for the rare-earth element plots (see discussion in Jarvis & Jarvis, 1985). The expanded uncertainty of all measurements is less than 10% (relative).

The <2 μm acid-insoluble residue fraction was also separated and subjected to mineralogical analysis by X-ray diffraction (XRD). For details of the methodology adopted see Jeans *et al.* (2014).

RESULTS AND DISCUSSION

The results obtained for the acid insoluble residue major, trace and rare-earth elements are presented in Table 1. The data reveal a number of important features which provide additional insights into the development of marl seams; these would not have been readily seen from bulk rock samples due to their high, diluting carbonate content. In order to make chemical comparison between samples, elemental data were normalized to the Al_2O_3 value of the sample. This approach assumes that aluminium is primarily associated with clay minerals in the non-carbonate fraction. Normalisation of data to the aluminium value removes the inter-sample variability caused by

FIG. 1 (*facing page*). X-ray diffraction traces of glycerolated acid-insoluble residues from the <2 μm fraction of representative white chalks and marl seams. All samples are dominated by smectite or smectite-rich illite/smectite (peak centred on ~18 Å) with subordinate illite (peaks at 10 Å and 5 Å) and the low, variable presence of talc (peaks at 9.2 Å and 3.06 Å). The 3.34 Å peak is a mixture of illite and quartz. Visual analysis of peak shape and height indicate that the relative abundance of illite and smectite remain broadly constant throughout the section with the exception of the Old Nore Marl where the illite peak is notably smaller. Tescombe Marl 2 also has a slightly smaller illite peak than that seen in the white chalks and marls above and below. The remaining marl seams showed no apparent difference to the residue from the white chalks above and below.
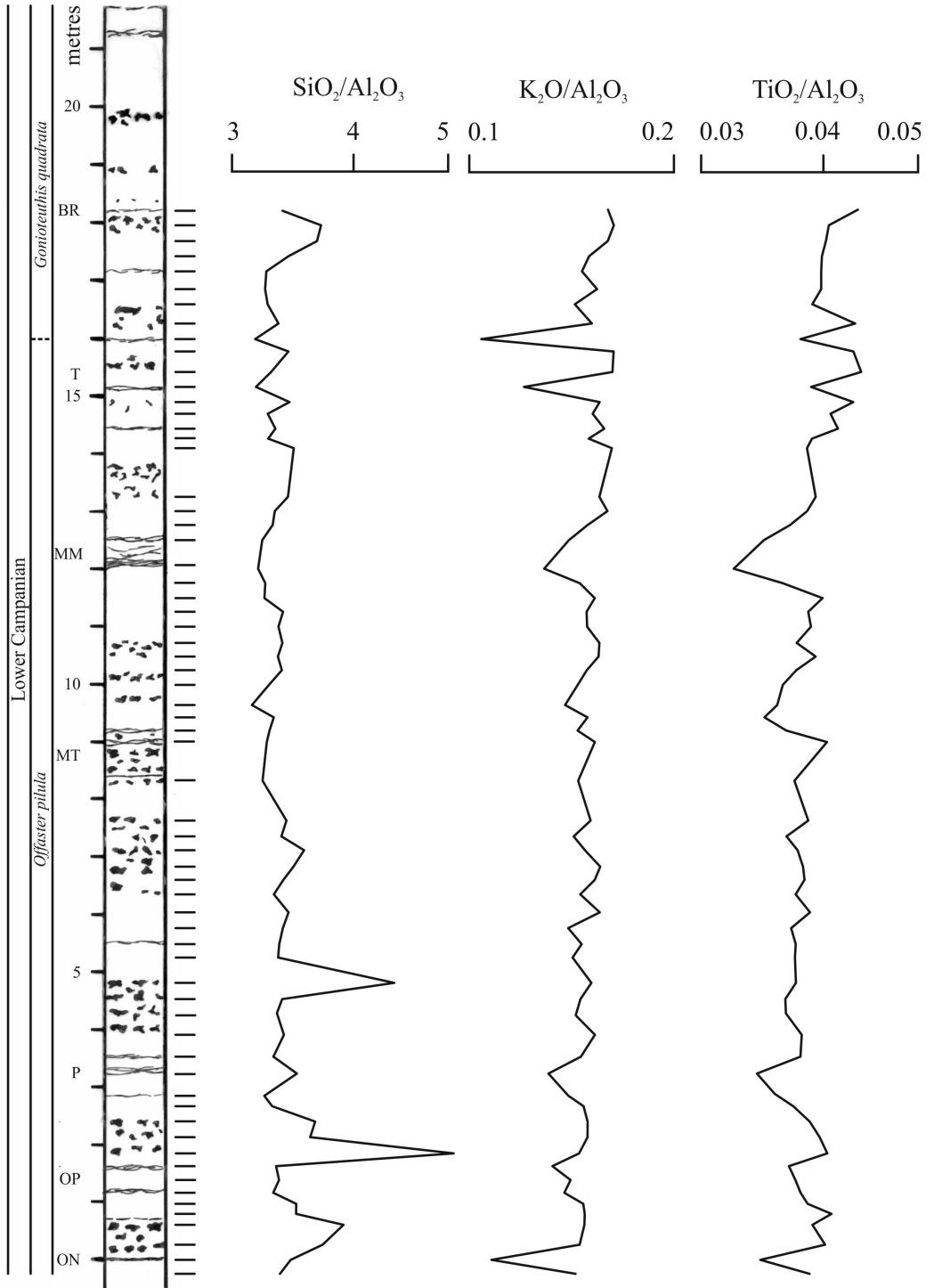
factors such as dilution by quartz which would otherwise hinder interpretation.

A representative selection of the <2 µm glycerolated acid insoluble residue mineralogical XRD tracings are presented in Fig. 1. All samples are dominated by smectite or smectite-rich illite/smectite (peak centred on ~18 Å) with subordinate illite (peak at 10 Å) and the low, variable presence of talc (peak at 9.2 Å). Visual analysis of peak shape and height indicate that the relative abundance of illite and smectite remain broadly constant throughout the section with the exception of the Old Nore Marl where the illite peak is notably smaller. Telscombe Marl 2 also has a slightly smaller illite peak than that seen in the white chalks and marls above and below. The remaining marl seams showed no apparent difference to the residue from the white chalks above and below at the resolution of our X-ray diffraction measurements.

Major element data

Stratigraphic plots of selected aluminium normalized elements are presented in Fig. 2. A plot of SiO_2/Al_2O_3 is interpreted as reflecting variation in the quartz/clay ratio. It shows isolated, single sample spikes and also low magnitude increases, both of which have a broad association with the flint bands within the chalks. There is no increase in the SiO_2/Al_2O_3 ratio associated with the presence of marl seams.

A plot of K_2O/Al_2O_3 is instructive in illustrating changes in the proportion of K-rich minerals in the non-carbonate fraction. Earlier mineralogical studies discussed above have demonstrated that potassium feldspar is rare or absent in the non-carbonate fraction and that the dominant K-bearing phase is the clay mineral illite. The reported presence of rare, small alkali feldspar crystals between the Peacehaven and Meeching Triple Marls (Jeans *et al.*, 2014) does not influence the K_2O/Al_2O_3 plot.

Examination of the stratigraphic profile for K_2O/Al_2O_3 shows that all prominent clay-rich marl beds with the exception of Telscombe Marl 1 and the Black Rabbit Marl (Fig. 2) are associated with a decrease in the K_2O/Al_2O_3 ratio. This is mimicked by a decrease in the TiO_2/Al_2O_3 ratio. This pattern is interpreted as indicating that most if not all the marl seams have a slightly more smectite-rich non-carbonate mineralogy than that found in the over- and underlying white chalks. This is partly supported by XRD mineralogical traces of the Old Nore Marl and Telscombe Marl 2, but for most of the other marls the XRD data are inconclusive (Fig. 1). Jeans *et al.* (2014) presented semi-quantitative insoluble residue data for a number of fractions (<0.2 µm, 0.2−1 µm and 1−2 µm) from selected samples which also adds some support to our conclusion in that the relative abundance of illite in the volumetrically dominant 0.2−1.0 µm fraction is higher in the white chalk samples studied when compared to the marls. In the <0.2 µm fraction smectite dominates all samples whilst in the volumetrically minor 1−2 µm fraction illite is more abundant. The absence of this feature in some of the very thin, minor marls is probably due to the relatively low abundance of non-carbonate material in these beds and hence a greater incorporation of 'white chalk' derived non-carbonate material at the preparation/laboratory dissolution stage. Bulk rock geochemical data from these samples typically gave Al_2O_3 values of 0.5 to 0.75 wt.%, reflecting their low clay content. The change in K_2O and TiO_2/Al_2O_3 ratios between marl seams and white chalks implies that it is very unlikely that the marl seams were formed by syn-sedimentary or post depositional dissolution of the white chalk carbonate fraction, and that they are primary sedimentary features. The question as to whether the marl seams are derived from a different source is more difficult to answer as the slight change in elemental (and mineralogical) composition may be a reflection of the mode of transportation and/or the speed of transport from the source region to the site of

FIG. 2 (*facing page*). Stratigraphic plot of variation in SiO_2/Al_2O_3, K_2O/Al_2O_3 and TiO_2/Al_2O_3 as found in acid-insoluble residues through the Peacehaven Stairs section. SiO_2/Al_2O_3 data show subtle increases associated with many of the flint bands. K_2O/Al_2O_3 and TiO_2/Al_2O_3 show subtle decreases within most marl seams; this is interpreted as a slight reduction in the proportion of potassium- and titanium-bearing minerals (predominantly illite) within the marl seams. ON: Old Nore Marl (PHS2), OP: Old Nore Pair (PHS 7, 9), PP: Peacehaven Marl (PHS15), MT: Meeching Triple Marls (PHS31, 32, 33), MM: Meeching Marls (pair) (PHS 44, 45), T: Telscombe Marls (PHS 54), BR: Black Rabbit Marl (PHS 65).
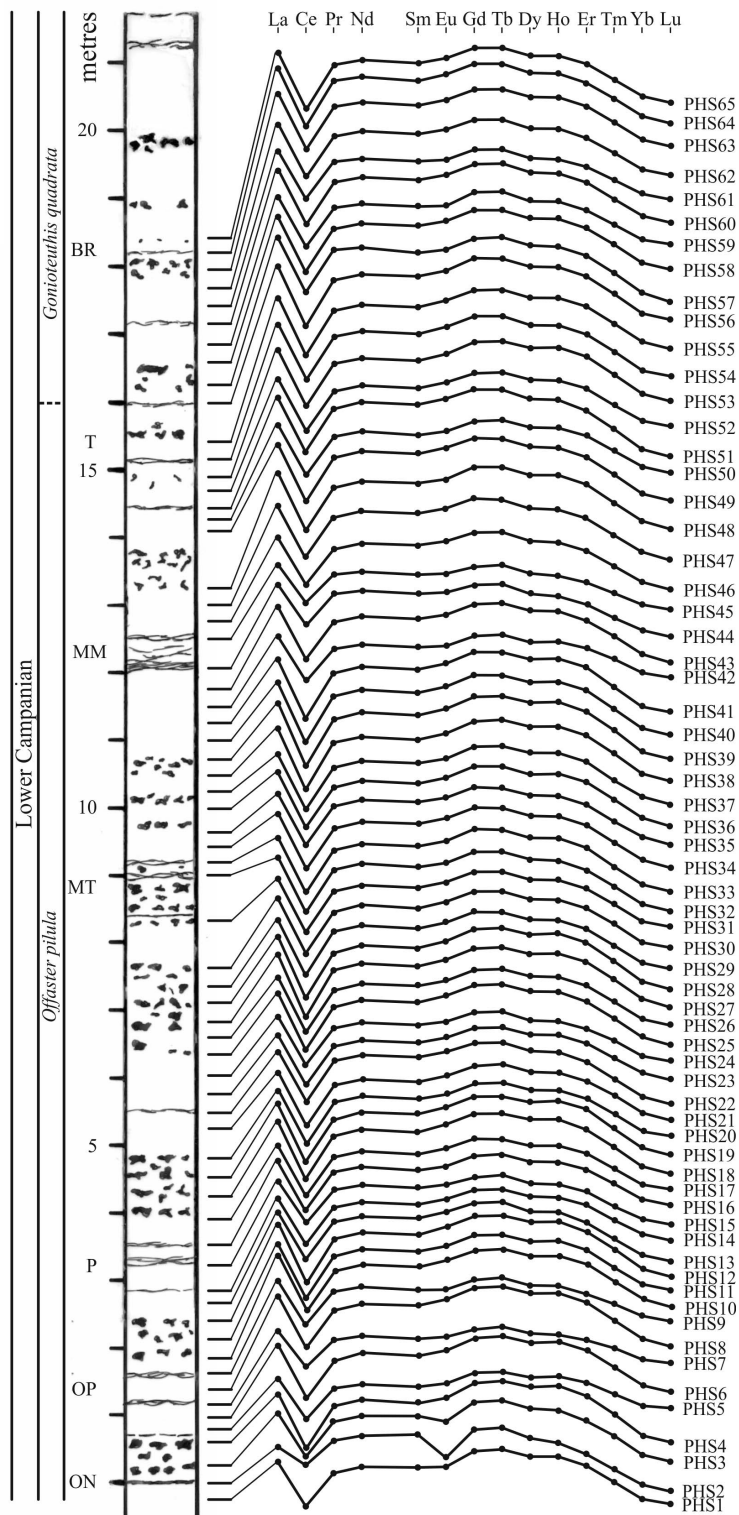
deposition. The elevated smectite content may also reflect the preferential formation of authigenic smectite within the marl layers, but we have no supporting evidence for this proposal. The only key exception to this discussion is the Old Nore Marl which is considered below.

Rare-earth element profiles and trace element composition

Previous studies of clay-rich beds in the Chalk have demonstrated the ability of the rare earth elements (*REE*) to differentiate between those of detrital provenance and those of a volcanogenic provenance (Wray, 1995, 1999; Wray & Wood, 1998, 2002). The key differentiating factor has been the presence or absence of a negative europium (Eu) anomaly in plots generated by normalising *REE* data to that derived from an average shale (in this case USGS Reference Material Cody Shale, SCo-1). The anomaly is generated by the preferential uptake of Eu into feldspar minerals within the magma chamber and their subsequent loss from the melt during fractional crystallization. Rocks such as rhyolites derived from evolved magmas are invariably characterized by a significant negative Eu anomaly (Rollinson, 1993). Shale-normalized *REE* patterns of the acid-insoluble residue of all samples are presented stratigraphically in Fig. 3. Most samples display a very similar pattern comprising of a negative Ce anomaly, a slight enrichment in the middle *REE* and a gradual depletion in the heavy *REE*. Sample PHS 2 from within the Old Nore Marl displays broadly the same pattern but with a reduced Ce anomaly and, importantly, a marked negative Eu anomaly; a slight, negative Eu anomaly is also present in sample PHS 3, a chalk sample just above the Old Nore Marl. Using experience gleaned from previous studies of marls of Turonian and Coniacian age (Wray, 1999) we propose that the Old Nore Marl is volcanically derived; its higher smectite content

means that it is best classified as a bentonite. The slight Eu anomaly present in sample PHS 3 either represents a residual tail of activity or, more likely, the upward reworking of non-carbonate material from the Old Nore Marl by bioturbating organisms. These are the only two marly samples which display these traits; all other marls display a profile which has previously been interpreted as indicating a detrital origin (Wray, 1999).

The use of acid-insoluble residues also enables the investigation of the *REE* profiles of the non-carbonate fraction of white chalks, something which has not been possible in previous studies. The white chalk residue profiles are near identical to that of the detrital marls and none of the white chalk profiles display a negative Eu anomaly (with the exception of PHS 3, as discussed above). This implies that there was little or no low-level or background volcanism occurring during the accumulation of the studied white chalk succession − the Old Nore event was a distinct, time limited event.

Differentiation of the Old Nore Marl from the other insoluble residues can also be demonstrated using trace element plots. The igneous petrology plot of Pearce (1996) is often used to classify bentonites (e.g. Wray, 1999; Batchelor *et al*., 2003). Figure 4a shows a clear distinction between the Old Nore Marl (PHS 2) and other samples, with the adjacent sample PHS 3 plotting at an intermediate position. If the elements plotted are considered immobile then the plot implies that the Old Nore Marl is of rhyodacite to dacite composition, a broadly comparable composition to that of the Turonian-age bentonites of the Anglo-Paris Basin as discussed by Wray (1999). A plot of Yb *vs.* Ta (Pearce *et al.*, 1984) in Fig. 4b indicates that the Old Nore bentonite plots within the WPG field and thus has a within plate source. In both plots all the other detrital marl seams plot close to the mean value for Continental Crust (Rudnick & Gao, 2003). This is coincidentally within the andesite/basalt

FIG. 3 (*facing page*). Stratigraphic plot of variation in shale-normalized rare earth element (*REE*) patterns as found in acid-insoluble residues through the Peacehaven Stairs section. Most patterns are very similar and display a negative Ce anomaly, a slight enrichment in the middle *REE*s and a gradual depletion in the heavy *REE*s. Notably different is sample PHS 2 (Old Nore Marl) which displays a reduced Ce anomaly and a distinct negative Eu anomaly, and PHS3 which has a reduced but still observable negative Eu anomaly. With reference to previous studies in older chalks the Old Nore Marl is considered to be a bentonite. ON: Old Nore Marl, OP: Old Nore Pair, PP: Peacehaven Marl, MT: Meeching Triple Marls, MM: Meeching Marls (pair), T: Telscombe Marls; BR: Black Rabbit Marl.

field but no inference is made as to this being their origin, it is an artefact of the use of an igneous-petrology-derived diagram to demonstrate the likely composition of the Old Nore Marl/bentonite. Their inclusion on the plot reinforces the observation that data obtained from the Old Nore Marl are atypical for a detrital sediment.

Distant correlation and source of the Old Nore Marl

The presence of a bentonite close to the Santonian-Campanian boundary is also observed in northern Germany. The marl M1 found in the Lagerdorf section is also considered to be a bentonite (Schonfeld *et al.*, 1996) and the natural correlation compliant with the biostratigraphy is to correlate the Old Nore Marl to M1 (C.J. Wood *pers. comm.*; Mortimore *et al.*, 2001). Whilst trace element data show that the two beds have broadly the same composition there is an offset between M1 and Old Nore Marl (Figs 3a,b). This is not surprising given the geographic distance between the two sites and this phenomenon was also observed when comparing north German and southern England bentonites of Turonian age (Wray, 1999).

The source of bentonites in Cretaceous chalks was discussed by Wray (1999) and remains problematic with no clear candidate. Wray proposed that the source for bentonites of Turonian and Coniacian age was most likely to be associated with the early stages of rifting of the North Atlantic to the north west of the UK and this remains the case for the Campanian Old Nore Marl.

CONCLUSIONS

This work demonstrates that the provenance of non-carbonate material in early Campanian chalks varies. Non-carbonate material found in white chalks and most marl seams is considered to be of detrital origin, whilst the Old Nore Marl is volcanically derived and is best classified as a bentonite. This division is based primarily on chemical criteria that have been successfully applied to marl seams of Turonian and Coniacian
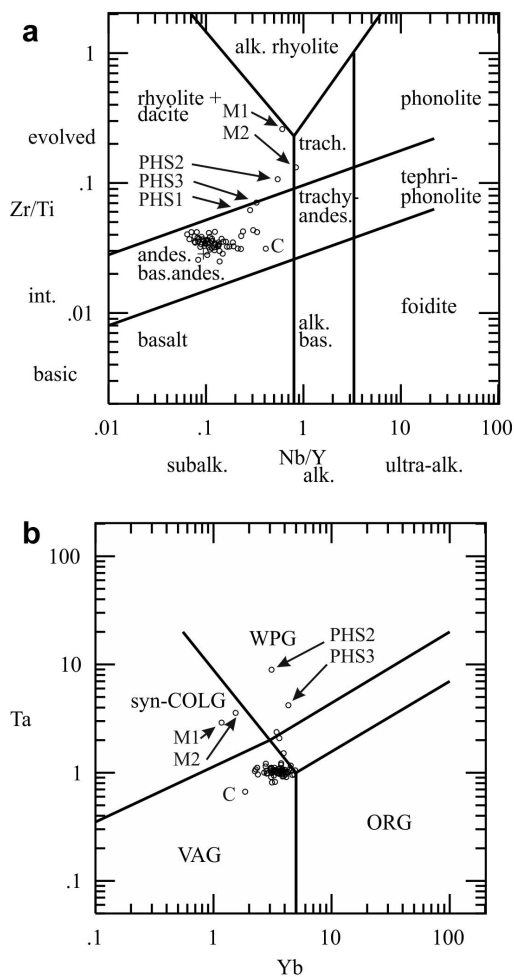

FIG. 4. (a) Acid-insoluble residue data plotted on the discrimination plot of Pearce (1996). Note that PHS 2 (Old Nore Marl) plots separately from the remaining samples and that samples PHS 1 and 3 immediately below and above the Old Nore Marl plot at a location between PHS 2 and the remaining samples. Although factors such as modification during transport and post depositional alteration must be considered, sample PHS 2 plots in the rhyolite/dacite field. Samples PHS 1 and 3 are interpreted as containing a proportion of bentonite due to bioturbation and reworking. There is no suggestion that the detrital beds are derived from an andesitic source and they plot close to the value for mean Continental Crust (*C*) (Rudnick & Gao, 2003). Also included are the results for bentonite M1 and the younger bentonite M2 from the north German succession (see Schonfeld *et al.*, 1996). (b) Granite provenance plot after (Pearce *et al.*, 1984). A plot of data from the Old Nore Marl (PHS2) indicates that it came from a within-plate source. VAG: volcanic-arc granite; ORG: ocean-ridge granite; WPG: within plate granite; syn-COLG: syn-collisional granite.

age. Chemical analysis of acid-insoluble residues demonstrates that the detrital marls have a subtly different composition to that of white chalks, implying that they were not formed through the penecontemporaneous or diagenetic dissolution of white chalks. Trace element analysis of the Old Nore Marl implies that the original ash had an acidic, rhyodacite to dacite composition and is derived from a within plate source, most likely associated with the early stages of rifting of the North Atlantic, to the northwest of the UK.

REFERENCES

Batchelor R.A., Harper D.A.T. & Anderson T.B. (2003) Geochemistry and potential correlation of Silurian (Telychian) metabentonites from Ireland and SW Scotland. *Geological Journal*, **38**, 161−174.

Brydone R.M. (1914a) The Zone of *offaster pilula* in the south English Chalk: Part 1. *Geological Magazine (Decade VI)*, **1**, 359−369.

Brydone R.M. (1914b) The Zone of *offaster pilula* in the south English Chalk: Part 4, general conclusions. *Geological Magazine (Decade VI)*, **1**, 509−513.

Deconinck J.-F. & Chamley H. (1995) Diversity of smectite origins in late Cretaceous sediments: Examples of chalks from northern France. *Clay Minerals*, **30**, 365−379.

Dorn P. & Bräutigam F. (1959) Hinweise auf oberkrei-devulkanismus in NW-Deutschland. *Abhandlungen der Braunschweigischen Wissenschaftlichen Gesellschaft*, **11**, 1−4.

Ernst G., Schmid F. & Seibertz E. (1983) Event-stratigraphie im Cenoman und Turon von NW-Deutschland. *Zitteliana*, **10**, 531−554.

Jarvis I. & Jarvis K.E. (1985) Rare-earth element geochemistry of standard sediments - a study using inductively coupled plasma spectrometry. *Chemical Geology*, **53**, 335−344.

Jeans C.V. (2006) Clay mineralogy of the Cretaceous strata of the British Isles. *Clay Minerals*, **41**, 47−150.

Jeans C.V., Tosca N., Hu X.F. & Boreham S. (2014) Clay mineral-grain size-calcite cement relationships in the Upper Cretaceous chalk, UK: A preliminary investigation. *Clay Minerals*, **49**, xxx-xxx.

Mortimore R.N. (1986) Stratigraphy of the Upper Cretaceous White Chalk of Sussex. *Proceedings of the Geologists' Association*, **97**, 97−140.

Mortimore R.N., Wood C.J. & Gallois R.W. (2001) *British Upper Cretaceous Stratigraphy*. Joint Nature Conservation Committee, Peterborough, 558 pp.

Pacey N.R. (1984) Bentonites in the Chalk of central eastern England and their relation to the opening of the northeast Atlantic. *Earth and Planetary Science Letters*, **10**, 48−60.

Pearce J.A. (1996) A user's guide to basalt discrimination diagrams. Pp. 79−113 in: *Trace element Geochemistry of Volcanic rocks; Applications for Massive Sulphide Exploration, Short Course Notes* (A.H. Bailes, E.H. Christiansen, A.G. Galley, G.A. Jenner, J.D. Keith, R. Kerrich, D.R. Lentz, C.M. Lesher, S.B. Lucas, J.N. Ludden, J.A. Pearce, S.A. Peloquin, R.A. Stern, W.E. Stone, E.C. Syme, H.S. Swinden, & D.A. Wyman, editors). Geological Association of Canada, **12**.

Pearce J.A., Harris N.B.W. & Tindle A.G. (1984) Trace element discrimination diagrams for the tectonic interpretation of granitic rocks. *Journal of Petrology*, **25**, 956−983.

Perrin R.M.S. (1971) *The Clay Mineralogy of British Sediments*. Mineralogical Society (Clay Minerals Group), London, 247 pp..

Rollinson H.R. (1993) *Using Geochemical Data: Evaluation, Presentation, Interpretation*. Longman Scientific & Technical, Harlow, 352 pp.

Rudnick R.L. & Gao S. (2003) Composition of the continental crust. Pp. 1−64 in: *The Crust; Treatise on Geochemistry* (R.L. Rudnick, editor), **3**,Elsevier.

Schonfeld J., Schulz M.-G., McArthur J.M., Burnett J., Gale A.S., Hambach U., Hansen H.J., Kennedy W.J., Rasmussen K.L., Thirlwall M.F. & Wray D.S. (1996) New results on biostratigraphy, palaeomagnetism, geochemistry and correlation from the standard section for the Upper Cretaceous White Chalk of northern Germany (Lägerdorf-Kronsmoor-Hemmoor). *Mitteilungen aus dem Geologisch-Paläontologischen Institut der Universität Hamburg*, **77**, 545−575.

Seibertz E. & Vortisch W. (1979) Zur stratigraphie, petrologie und genese einer bentonit-lage aus dem oberen mittel-Turon (Oberkreide) des südöstlichen Münsterlandes. *Geologische Rundschau*, **68**, 649−679.

Weir A.H. & Catt J.A. (1965) The mineralogy of some Upper Chalk samples from the Arundel area, Sussex. *Clay Minerals*, **6**, 97−110.

Wray, D.S. (1995) Origin of clay-rich beds in Turonian chalks from Lower Saxony, Germany − a rare earth element study. *Chemical Geology*, **119**, 161−173.

Wray D.S. (1999) Identification and long-range correlation of bentonites in Turonian - Coniacian (Upper Cretaceous) chalks of northwest Europe. *Geological*

Magazine, **136**, 361−371.

Wray D.S. & Wood C.J. (1995) Geochemical identification and correlation of tuff layers in Lower Saxony, Germany. *Berliner Geowissenschaftliche Abhandlungen*, **E16.1**, 215−226.

Wray D.S. & Wood C.J. (1998) Distinction between detrital and volcanogenic clay-rich beds in Turonian-Coniacian Chalks of eastern England. *Proceedings of the Yorkshire Geological Society*, **52**, 95−105.

Wray D.S. & Wood C.J. (2002) Identification of a new bentonite in sediments of mid-Turonian age from Lower Saxony, Germany and its correlation within NW Europe. *Austrian Academy of Science Series: Schriftenreihe der Erdwissenschaftlichen Kommissionen*, **15**, 47−58.

Wray D.S., Kaplan U. & Wood C.J. (1995) Tuffvorkommen und ihre bio- und eventstratigraphie im Turon des Teutoburger Waldes, Der Egge und Des Haarstrangs. *Geologie und Paläontologie in Westfalen*, **37**, 1−53.

Wray D.S., Wood C.J., Ernst G. & Kaplan U. (1996) Geochemical subdivision and correlation of clay-rich beds in Turonian sediments of northern Germany. *Terra Nova*, **8**, 603−610.