

Supplementary data for the article:

Andrić, F.; Šegan, S.; Dramićanin, A.; Majstorović, H.; Milojković-Opsenica, D. Linear Modeling of the Soil-Water Partition Coefficient Normalized to Organic Carbon Content by Reversed-Phase Thin-Layer Chromatography. *Journal of Chromatography A* **2016**, *1458*, 136–144. <https://doi.org/10.1016/j.chroma.2016.06.063>

SUPPLEMENTARY MATERIAL

Linear modeling of the soil-water partition coefficient normalized to organic carbon content by reversed-phase thin-layer chromatography

Filip Andrić^{1*}, Sandra Šegan², Aleksandra Dramićanin³, Helena Majstorović⁴, Dušanka Milojković-Opsenica¹

¹Faculty of Chemistry, University of Belgrade, Studentski trg 12-16, P.O. Box 51, 11158 Belgrade, Serbia

²Institute of Chemistry, Technology and Metallurgy, Njegoševa 12, 11000 Belgrade, Serbia

³Innovation center Faculty of Chemistry Ltd, Studentski trg 12-6, 11000 Belgrade, Serbia

⁴AIC „Tamiš“, Novoseljski put 33, 26000 Pančevo, Serbia

*Corresponding author

E-mail: andric@chem.bg.ac.rs

Phone: +381 11 3336755

TABLE OF CONTENTS

TABLE S1 List of standard compounds and unknowns	(p. 2)
TABLE S2 Retention factors of calibration set of compounds and unknowns - given as R_M , and $\log k$ values in the case of TLC and HPLC experiments respectively	(p. 4)
TABLE S3 Simple univariate calibration models based on TLC experiments	(p. 6)
Figure S1 PCR model – variable statistics demonstrating different impact of chromatographic systems in overall $\log K_{OC}$ modelling	(p. 7)
Figure S2 PLS model – variable statistics demonstrating different impact of chromatographic systems in overall $\log K_{OC}$ modelling	(p. 7)
TABLE S4 Chromatographically determined and computationally estimated $\log K_{OC}$ values	(p. 9)
TABLE S5a Comparison of chromatographic and <i>in silico</i> methods – collection of normalized SRD values	(p. 11)
TABLE S5b Normalized SRD values obtained by sevenfold jack-knife resampling procedure	(p. 12)
TABLE S6 LSER regression coefficients of TLC systems	(p. 13)
REFERENCES	(p. 14)
A: List of literature sources of $\log K_{OC}$ values	
B: List of literature sources of pK_a values	

TABLE S1 List of standard compounds and unknowns accompanied with the list of $\log K_{OC}$ values measured by soil sorption experiments (U.S. EPA); Molar mass M ; Abraham's solvatochromic parameters (A – hydrogen bond proton donating ability, B – hydrogen bond proton accepting ability, S – dipolarity/polarizability, E – molar refractivity in excess, V – McGowan's molar molecular volume); Acidity constants pK_a ; Ionization degree α at $pH = 6$

No	Compound	$\log K_{OC}$	Ref	M (g/mol)	Solvatochromic parameters					pK_a^b	Ref	α (%) ^b $pH = 6$
					A	B	S	E	V			
<i>Standard compounds</i>												
1	Phenol	1.90	A1	94.12	0.60	0.30	0.89	0.81	0.775	9.99	B1	0.01
2	4-Nitrophenol	2.37	A1	139.12	0.82	0.26	1.72	1.07	0.949	7.18	B1	6.20
3	Benzyl Alcohol	1.10	A2	109.14	0.39	0.56	0.87	0.80	0.916	-	-	-
4	1-Naphthylamine	3.51	A1	143.20	0.20	0.57	1.26	1.67	1.185	3.92 ^a	B2	99.2 ^a
5	1-Naphthol	3.10	A1	144.18	0.60	0.37	1.05	1.52	1.144	9.34	B3	0.05
6	2,4-Dichlorophenol	2.81	A1	163.00	0.53	0.19	0.84	0.96	1.020	7.65	B1	2.19
7	Anthracene	4.31	A1	178.24	0.00	0.28	1.34	2.29	1.454	-	-	-
8	Acetophenone	1.80	A2	120.16	0.00	0.48	1.01	0.82	1.014	-	-	-
9	2,4,6-Trichlorophenol	3.03	A1	197.45	0.42	0.15	0.94	1.07	1.142	6.42	B1	27.55
10	Ethyl- <i>p</i> -hydroxybenzoate	2.21	A1	166.19	0.69	0.45	1.35	0.86	1.272	8.50	B4	0.32
11	<i>p</i> -Anisidine	1.93	A3	123.17	0.16	0.66	1.26	1.05	1.016	5.36 ^a	B5	81.36
12	1,2,3-Benzotriazole	1.69	A1	119.14	0.62	0.48	1.70	1.47	0.864	8.37	B3	0.42
13	Diphenylamine	2.78	A1	169.24	0.08	0.51	1.08	1.59	1.424	0.78 ^a	B6	>99.9 ^a
14	2,2'-Dipyridyne	1.60	A1	156.20	0.00	0.81	1.33	1.38	1.242	4.33 ^a	B7	>97.91 ^a
15	4-Bromoaniline	1.96	A1	172.03	0.31	0.30	1.19	1.19	0.991	3.86 ^a	B8	>99.2 ^a
16	Benzophenone	2.63	A1	182.23	0.00	0.50	1.50	1.45	1.481	-	-	-
17	4-Aminobenzoic acid	1.70	A4	137.15	0.94	0.69	1.61	1.08	1.032	2.38	B9	99.98
18	Pyrene	4.90	A1	202.26	0.00	0.25	1.52	2.60	1.585	-	-	-
19	Benzo[<i>a</i>]pyrene	5.95	A2	252.32	0.00	0.31	1.84	3.32	1.954	-	-	-
20	Fluorene	3.70	A1	166.23	0.00	0.25	1.06	1.59	1.357	-	-	-
21	Acenaphthene	3.59	A1	154.22	0.00	0.26	1.14	1.75	1.259	-	-	-
22	Naphthalene	2.96	A1	128.18	0.00	0.20	0.92	1.34	1.085	-	-	-
23	Phenanthrene	4.35	A1	178.24	0.00	0.29	1.29	2.06	1.454	-	-	-
24	Dibenz[<i>a,h</i>]anthracene	6.22	A1	278.36	0.00	0.35	1.99	3.43	2.192	-	-	-
25	Benz[<i>a</i>]anthracene	5.30	A1	228.30	0.00	0.29	1.66	2.71	1.823	-	-	-
26	Simazine	2.10	A1	201.69	0.33	0.95	1.20	1.55	1.479	1.62 ^a	B10	>99.9 ^a
27	Propazine	2.40	A1	229.75	0.30	0.97	1.14	1.47	1.761	1.85 ^a	B10	>99.9 ^a
28	Ametryn	2.59	A1	227.38	0.26	1.07	1.27	1.50	1.802	4.10 ^a	B11	>98.8 ^a
29	Prometryn	2.85	A1	241.41	0.26	1.10	1.26	1.51	1.940	4.05 ^a	B11	>98.9 ^a

No	Compound	log <i>K</i> _{OC}	Ref	<i>M</i> (g/mol)	Solvatochromic parameters					p <i>K</i> _a ^b	Ref	α (%) ^b pH = 6
					<i>A</i>	<i>B</i>	<i>S</i>	<i>E</i>	<i>V</i>			
<i>Unknowns</i>												
30	3-Nitrophenol	-		139.12	0.79	0.23	1.57	1.05	0.949	8.36	B1	0.43
31	2-Naphthol	-		144.18	0.61	0.40	1.08	1.52	1.144	9.67	B12	0.02
32	4-Hydroxybenzaldehyde	-		122.13	0.85	0.37	1.54	1.01	0.932	7.61	B3	2.40
33	2-Aminophenol	-		109.14	0.60	0.66	1.10	1.11	0.875	9.44 (-OH)	B1	0.00
34	4- <i>t</i> -Butylphenol	-		150.24	0.56	0.41	0.89	0.81	1.339	10.31	B1	0.00
35	2,6-Dimethylphenol	-		122.18	0.39	0.39	0.79	0.86	1.057	10.59	B1	0.00
36	4-Methoxyphenol	-		124.15	0.57	0.48	1.17	0.90	0.975	10.27	B1	0.01
37	Methyl- <i>p</i> -hydroxybenzoate	-		152.16	0.69	0.45	1.37	0.90	1.131	8.47	B4	0.34
38	2-Nitrobenzaldehyde	-		151.13	0.00	0.38	1.59	1.12	1.047	-	-	-
39	3-Nitrobenzaldehyde	-		151.13	0.00	0.48	1.49	1.10	1.047	-	-	-
40	Phthalimide	-		147.14	0.39	0.44	1.97	1.18	1.021	8.3	B13	0.50
41	Oxazepam	-		286.73	0.45	1.60	1.10	2.35	1.992	10.9 (-OH)	B14	0.00
42	Lorazepam	-		321.18	0.45	1.63	1.28	2.51	2.114	11.5 (-OH)	B15	0.00
43	Clonazepam	-		315.73	0.33	1.50	1.75	2.46	2.107	10.5 (-OH)	B16	0.00
44	Bromazepam	-		316.17	0.33	1.62	1.38	2.31	1.940	11.0 (-OH)	B17	0.00
45	Diazepam	-		284.76	0.00	1.25	1.57	2.08	2.074	3.40 ^a	B18	99.75 ^a
46	Nitrazepam	-		281.29	0.47	1.10	2.17	2.21	1.985	10.8	B15	0.00
47	Chlordiazepoxide	-		299.78	0.13	0.94	1.65	1.95	2.174	4.80 ^a	B18	94.06 ^a
48	Clobazam	-		300.76	0.00	1.47	2.49	2.29	2.133	6.65 ^a	B19	18.30 ^a
49	Medazepam	-		270.78	0.00	0.67	1.40	1.92	2.058	6.20 ^a	B20	38.69 ^a
50	Chrysene	-		228.30	0.00	0.29	1.66	2.71	1.823	-	-	-

^aDissociation degree is referred to the positively charged (protonated) form

^bCompounds marked in bold are present in negatively charged form under studied chromatographic conditions

TABLE S2 Retention factors of calibration set of compounds and unknowns - given as R_M , and $\log k$ values in the case of TLC and HPLC experiments respectively

Comp. no.	HPLC $\log k$	TLC																			
		RP18 MeOH					RP18 ACN					CN MeOH					CNACN				
		40%	50%	60%	70%	80%	40%	50%	60%	70%	80%	40%	50%	60%	70%	40%	45%	50%	55%	60%	
1	-0.040	0.580	0.267	0.057	-0.175	-0.367	0.475	0.261	-0.276	-0.690	-0.738	0.010	-0.285	-0.409	-0.479	-0.076	-0.192	-0.318	-0.460	-0.558	
2	-0.064	0.706	0.395	0.184	-0.059	-0.267	0.513	0.232	-0.187	-0.632	-0.738	0.337	0.005	-0.181	-0.302	0.044	-0.042	-0.220	-0.344	-0.460	
3	-0.265	0.629	0.329	0.064	-0.161	-0.324	0.347	0.109	-0.295	-0.632	-0.660	-0.151	-0.402	-0.494	-0.592	-0.180	-0.233	-0.369	-0.486	-0.586	
4	0.487	1.252	0.868	0.533	0.199	-0.064	0.916	0.617	0.204	-0.131	-0.285	0.564	0.179	0.034	-0.130	0.268	0.233	0.057	-0.114	-0.202	
5	0.443	1.444	1.007	0.589	0.240	-0.040	0.916	0.591	0.152	-0.235	-0.389	0.623	0.192	-0.034	-0.183	0.220	0.199	-0.038	-0.172	-0.291	
6	0.248	1.358	1.183	0.710	0.323	0.017	0.876	0.644	0.169	-0.196	-0.345	0.536	0.146	-0.097	-0.285	0.325	0.172	-0.064	-0.192	-0.313	
7	0.664	1.688	1.721	1.612	1.092	0.641	1.555	1.466	0.973	0.602	0.322	1.284	0.822	0.490	0.143	0.654	0.548	0.396	0.239	-0.019	
8	0.187	0.979	0.725	0.363	0.088	-0.150	0.668	0.457	0.107	-0.238	-0.361	0.125	-0.105	-0.299	-0.418	0.108	0.003	-0.180	-0.337	-0.436	
9	0.382	1.521	1.130	0.777	0.458	0.209	0.985	0.703	0.384	0.062	-0.137	0.649	0.252	0.009	-0.233	0.440	0.333	0.092	-0.076	-0.306	
10	0.219	1.127	0.809	0.388	0.088	-0.174	0.596	0.314	-0.043	-0.457	-0.606	0.187	-0.099	-0.327	-0.467	0.025	-0.003	-0.213	-0.359	-0.521	
11	-0.431	0.575	0.348	0.113	-0.161	-0.324	0.418	0.239	0.020	-0.398	-0.490	0.406	0.265	0.145	0.016	0.318	0.219	0.097	-0.088	-0.205	
12	-0.390	0.502	0.239	-0.009	-0.244	-0.446	0.194	-0.017	-0.368	-0.845	-1.038	-0.138	-0.339	-0.481	-0.518	-0.207	-0.261	-0.362	-0.529	-0.558	
13	0.317	1.521	1.227	0.929	0.538	0.199	1.474	1.098	0.640	0.183	-0.053	0.987	0.543	0.161	0.000	0.673	0.645	0.303	0.120	-0.142	
14	0.366	1.179	0.823	0.444	0.182	-0.034	0.679	0.526	0.287	0.016	-0.027	0.051	-0.153	-0.338	-0.402	0.083	0.035	-0.095	-0.185	-0.306	
15	0.228	1.127	0.789	0.528	0.138	-0.115	0.805	0.555	0.215	-0.183	-0.341	0.382	0.077	-0.148	-0.216	0.193	0.126	-0.019	-0.133	-0.320	
16	0.624	1.730	1.310	0.692	0.501	0.177	1.344	1.007	0.597	0.163	-0.060	0.665	0.208	-0.081	-0.213	0.053	0.032	-0.100	-0.275	-0.438	
17	-0.341	-0.309	-0.419	-0.579	-0.586	-0.763	-0.251	-0.329	-0.634	-1.028	-1.071	-0.245	-0.462	-0.570	-0.597	0.199	0.086	-0.198	-0.331	-0.502	
18	0.787	2.035	1.721	1.693	1.225	0.790	1.736	1.371	1.129	0.741	0.455	1.549	0.872	0.490	0.243	0.910	0.815	0.499	0.294	0.058	
19	0.909	2.035	2.026	1.693	2.024	1.241	2.041	1.522	1.398	1.028	0.764	1.648	1.245	0.807	0.446	1.048	0.956	0.625	0.452	0.135	
20	0.815	1.730	1.541	1.168	0.958	0.562	1.555	1.273	0.861	0.461	0.185	1.206	0.580	0.250	0.035	0.651	0.604	0.241	0.146	-0.077	
21	0.716	2.035	2.026	1.113	0.787	0.387	1.736	1.178	0.823	0.368	0.140	1.126	0.516	0.224	0.006	0.625	0.479	0.213	0.127	-0.116	
22	0.514	2.035	2.026	1.040	0.611	0.309	1.242	1.074	0.668	0.249	0.020	1.086	0.468	0.207	-0.009	0.558	0.462	0.197	0.106	-0.126	
23	1.026	2.035	2.026	1.512	1.036	0.635	1.736	1.350	0.983	0.546	0.464	1.319	0.733	0.356	0.082	0.767	0.614	0.265	0.205	-0.064	
24	1.198	2.035	2.026	1.524	1.383	0.940	2.041	1.522	1.297	1.091	0.826	1.601	1.253	0.768	0.417	1.053	0.922	0.567	0.436	0.168	
25	0.967	2.035	2.026	1.693	1.457	1.041	2.041	1.702	1.297	0.897	0.527	1.493	1.201	0.670	0.327	0.998	0.829	0.448	0.359	0.142	

TABLE S2 Continues

Comp. no.	HPLC log <i>k</i>	TLC																			
		RP18 MeOH					RP18 ACN					CN MeOH					CN ACN				
		40%	50%	60%	70%	80%	40%	50%	60%	70%	80%	40%	50%	60%	70%	40%	45%	50%	55%	60%	
26	0.084	1.421	1.007	0.518	0.171	-0.067	0.742	0.427	0.124	-0.147	-0.296	0.214	-0.086	-0.254	-0.418	0.089	-0.093	-0.173	-0.280	-0.357	
27	0.277	2.035	1.412	0.868	0.429	0.129	1.145	0.788	0.380	0.062	-0.100	0.454	0.027	-0.135	-0.387	0.543	0.393	-0.013	-0.063	-0.236	
28	0.380	1.733	1.419	1.033	0.473	0.208	1.029	0.585	0.375	0.094	-0.085	0.406	0.101	-0.166	-0.383	0.169	0.140	0.051	-0.053	-0.110	
29	0.461	1.733	1.548	1.175	0.638	0.307	1.341	0.746	0.543	0.190	0.034	0.538	0.177	-0.127	-0.337	0.224	0.172	0.076	0.003	-0.078	
30	0.038	0.830	0.540	0.290	-0.113	-0.237	0.612	0.049	-0.190	-0.490	-0.768	0.240	-0.027	-0.278	-0.427	-0.122	-0.172	-0.340	-0.403	-0.492	
31	0.426	1.424	0.950	0.593	0.120	-0.081	0.880	0.243	0.043	-0.285	-0.529	0.430	0.101	-0.176	-0.341	-0.036	-0.033	-0.200	-0.294	-0.373	
32	-0.134	0.476	0.146	-0.027	-0.415	-0.455	0.288	-0.179	-0.439	-0.834	-1.037	-0.073	-0.330	-0.501	-0.568	-0.278	-0.339	-0.494	-0.572	-0.625	
33	-0.147	0.255	0.024	-0.119	-0.456	-0.546	0.243	-0.141	-0.391	-0.654	-0.965	0.063	-0.085	-0.223	-0.296	-0.020	-0.096	-0.141	-0.253	-0.377	
34	0.383	1.733	1.356	0.850	0.349	0.087	1.061	0.489	0.269	-0.104	-0.306	0.402	0.024	-0.292	-0.490	0.527	0.230	-0.004	-0.208	-0.104	
35	0.117	1.083	0.811	0.566	0.113	-0.078	0.851	0.352	0.154	-0.197	-0.430	0.215	-0.043	-0.317	-0.439	0.341	0.162	-0.029	-0.157	-0.291	
36	-0.123	0.555	0.203	0.068	-0.364	-0.425	0.364	-0.061	-0.287	-0.593	-0.957	-0.066	-0.294	-0.406	-0.503	-0.089	-0.252	-0.319	-0.412	-0.605	
37	0.214	0.913	0.526	0.258	-0.190	-0.297	0.467	-0.012	-0.200	-0.533	-0.811	0.040	-0.224	-0.467	-0.512	-0.102	-0.238	-0.336	-0.439	-0.595	
38	0.098	1.051	0.453	0.201	-0.196	-0.320	0.726	0.250	0.082	-0.244	-0.521	0.095	-0.111	-0.357	-0.387	0.115	-0.070	-0.147	-0.256	-0.312	
39	0.064	0.874	0.540	0.290	-0.113	-0.250	0.709	0.220	0.069	-0.244	-0.546	0.185	-0.037	-0.285	-0.348	0.102	-0.089	-0.157	-0.277	-0.326	
40	-0.006	0.633	0.332	0.104	-0.301	-0.373	0.320	-0.104	-0.245	-0.579	-0.831	-0.047	-0.238	-0.475	-0.578	-0.135	-0.350	-0.366	-0.483	-0.576	
41	0.470	1.553	1.164	0.690	0.196	-0.054	0.680	0.157	-0.040	-0.345	-0.564	0.243	-0.046	-0.331	-0.477	0.062	-0.238	-0.272	-0.384	-0.549	
42	0.590	1.553	1.235	0.665	0.173	-0.124	0.709	0.185	-0.027	-0.320	-0.573	0.303	-0.027	-0.313	-0.473	0.115	-0.185	-0.224	-0.347	-0.513	
43	0.445	1.553	1.056	0.593	0.097	-0.109	0.802	0.263	0.037	-0.264	-0.582	0.480	0.231	-0.130	-0.311	0.246	-0.083	-0.112	-0.243	-0.392	
44	0.362	1.424	0.959	0.562	0.081	-0.109	0.592	0.160	0.037	-0.214	-0.333	0.112	-0.147	-0.387	-0.469	0.030	-0.238	-0.258	-0.333	-0.496	
45	0.721	1.733	1.318	1.013	0.553	0.260	1.258	0.729	0.502	0.190	0.037	0.534	0.245	-0.118	-0.296	0.388	0.036	0.022	-0.138	-0.229	
46	0.405	1.424	1.046	0.640	0.120	-0.109	0.795	0.243	0.005	-0.299	-0.546	0.476	0.217	-0.095	-0.264	0.217	-0.127	-0.131	-0.305	-0.377	
47	0.522	1.553	1.101	0.690	0.113	-0.094	0.775	0.220	-0.018	-0.299	-0.555	0.476	0.197	-0.095	-0.275	0.204	-0.039	-0.141	-0.298	-0.355	
48	0.627	1.733	1.266	0.717	0.183	-0.048	1.019	0.430	0.183	-0.107	-0.322	0.480	0.221	-0.102	-0.293	0.303	-0.020	-0.051	-0.236	-0.291	
49	1.076	2.068	2.038	2.033	1.314	0.847	2.037	1.488	1.417	0.951	0.727	1.719	1.136	0.795	0.440	1.135	0.760	0.490	0.316	0.144	
50	0.742	2.068	2.038	2.033	1.724	1.204	2.037	1.570	1.316	0.842	0.610	2.024	1.547	0.697	0.232	1.305	0.931	0.595	0.443	0.236	

TABLE S3 Simple – the least square – univariate calibration models based on individual TLC experiments; Models are given in a form of $\log K_{OC} = a + bR_M$; statistically satisfactory models are marked in bold.

Chromatography system	<i>a</i>	<i>b</i>	$RMSE_{Cal}$	$RMSE_{Pred}$	R^2_{Cal}	R^2_{Pred}	<i>n</i>	<i>F</i>	<i>p</i>
RP-18 MeOH_40	0.74±0.52	1.61±0.34	0.916	0.974	0.5163	0.4538	29	28.81	<1·10 ⁻⁵
RP-18 MeOH_50	1.12±0.36	1.62±0.27	0.795	0.864	0.6354	0.5699	29	47.06	<1·10 ⁻⁵
RP-18 MeOH_60	1.47±0.24	1.97±0.25	0.659	0.718	0.7494	0.7033	29	80.74	<1·10 ⁻⁵
RP-18 MeOH_70	2.00±0.14	2.11±0.19	0.500	0.527	0.8557	0.8397	29	160.17	<1·10⁻⁵
RP-18 MeOH_80	2.57±0.12	2.60±0.25	0.516	0.557	0.8465	0.8210	29	148.90	<1·10⁻⁵
RP-18 ACN_40	0.87±0.29	2.00±0.24	0.629	0.710	0.7721	0.7094	29	91.45	<1·10 ⁻⁵
RP-18 ACN_50	1.28±0.25	2.26±0.28	0.642	0.687	0.7625	0.7282	29	86.66	<1·10 ⁻⁵
RP-18 ACN_60	2.08±0.16	2.24±0.25	0.589	0.637	0.8001	0.7664	29	108.07	<1·10 ⁻⁵
RP-18 ACN_70	2.92±0.13	2.19±0.25	0.597	0.641	0.7948	0.7635	29	104.61	<1·10 ⁻⁵
RP-18 ACN_80	3.33±0.15	2.40±0.30	0.642	0.680	0.7624	0.7333	29	86.64	<1·10 ⁻⁵
CN-MeOH_40	1.55±0.16	2.21±0.19	0.471	0.506	0.8722	0.8522	29	184.31	<1·10⁻⁵
CN-MeOH_50	2.27±0.10	2.67±0.19	0.403	0.441	0.9062	0.8878	29	260.93	<1·10⁻⁵
CN-MeOH_60	2.94±0.10	3.36±0.26	0.439	0.474	0.8892	0.8706	29	216.65	<1·10⁻⁵
CN-MeOH_70	3.69±0.13	4.13±0.40	0.531	0.556	0.8373	0.8220	29	138.92	<1·10⁻⁵
CN-ACN_40	1.75±0.18	3.33±0.35	0.568	0.614	0.8142	0.7831	29	118.34	<1·10 ⁻⁵
CN-ACN_45	2.00±0.16	3.50±0.36	0.557	0.598	0.8214	0.7938	29	124.17	<1·10 ⁻⁵
CN-ACN_50	2.72±0.12	4.43±0.44	0.537	0.568	0.8340	0.8140	29	135.68	<1·10⁻⁵
CN-ACN_55	3.29±0.13	4.39±0.46	0.562	0.600	0.8182	0.7926	29	121.53	<1·10 ⁻⁵
CN-ACN_60	4.26±0.20	5.37±0.63	0.615	0.643	0.7819	0.7618	29	96.81	<1·10 ⁻⁵

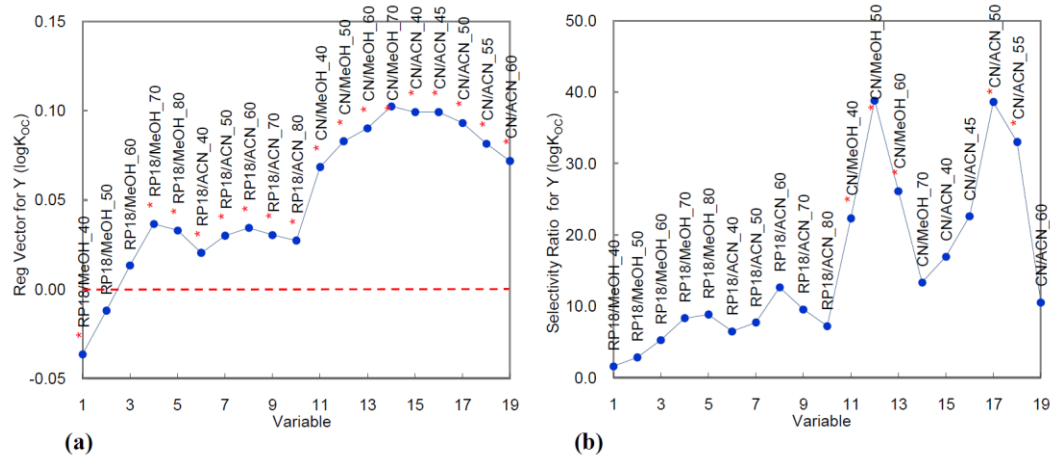


Figure S1. PCR model – variable statistics demonstrating different impact of chromatographic systems in overall logK_{OC} modelling; Regression vector plot (a) and sensitivity ratio diagram (b); Regression coefficients statistically significantly different from zero and points with high sensitivity ratio are marked by asterisk.

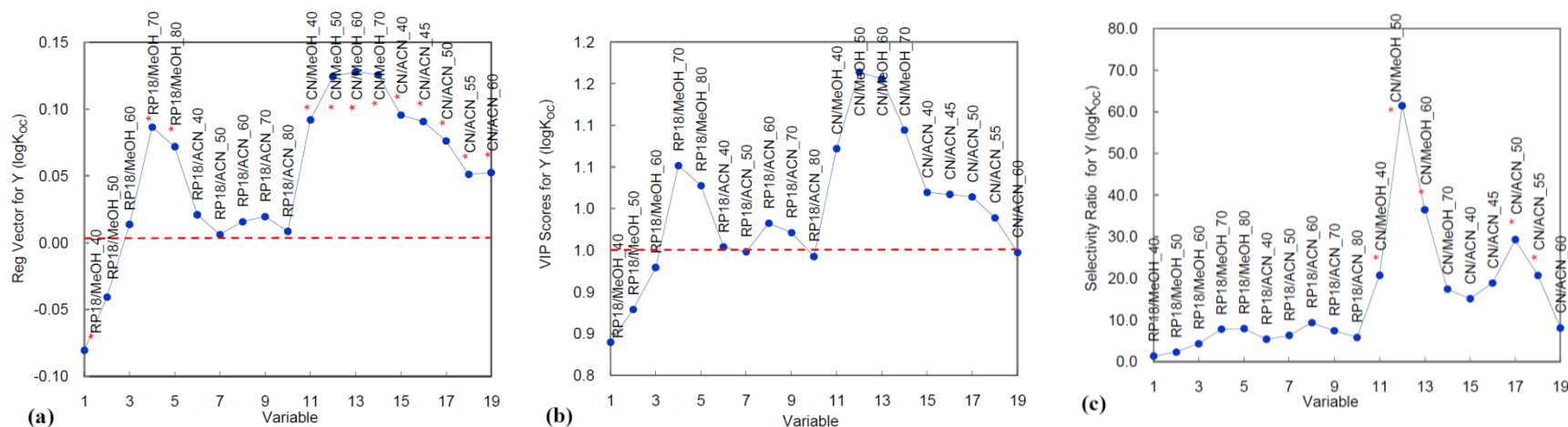


Figure S2. PLS model – variable statistics demonstrating different impact of chromatographic systems in overall logK_{OC} modelling; Regression vector plot (a), VIP score graph (b) and sensitivity ratio diagram (c); Regression coefficients statistically significantly different from zero ($p = 0.05$) and points with high sensitivity ratio are marked by asterisk.

PLS and PCR variable diagnostic plots demonstrate differences of individual chromatographic systems in contribution to the overall model. In the case of PCR 17 regression coefficients are statistically different from zero (tested by a t-test of cross-validated values, at the significance level $p = 0.05$). The exceptions are only two coefficients associated with chromatographic systems: RP18/MeOH_50 and RP18/MeOH_60 (Figure S1a). According to the values of sensitivity ratio (cut-off value = 20) five out of seventeen coefficients, corresponding to the chromatographic systems based on CN-silica in combination with methanol content of 40 – 60% v/v, and acetonitrile content of 50 – 55% v/v, may be considered as the mostly influential ones (**Figure S1b**).

In the case of the PLS model there are 13 statistically significant regression coefficients. The exceptions are 6 coefficients associated with chromatographic systems mostly obtained on RP18-silica in combination with acetonitrile-water mixtures (**Figure S2a**). Since the VIP scores describe the average contribution of variables to the overall model, it is easy to perceive those ones that contribute better than the average ($VIP > 1$). In that sense, thirteen chromatographic systems contributes to the overall PLS model better than the average, most of them are obtained on CN-silica layers in combination with methanol or acetonitrile as organic modifier (Figure S2b). Selectivity ratio plot (**Figure S2c**) indicates that the most influential chromatographic systems (cut-off value = 20) are those based on CN-silica in combination with methanol content of 40 – 60% v/v, and acetonitrile of 50 – 55% v/v (**Figure S2c**).

Clearly, the use of CN-silica layers has a stronger impact to the overall modeling of $\log K_{OC}$ than RP18-modified beds. This could lead to a presumption that CN-modified layers should be more suitable for chromatographic determination of $\log K_{OC}$ than hydrophobic, long alkyl-chain modified silica. This is in accordance with OECD guideline recommendations regarding the achieved accuracy.

TABLE S4 Chromatographically determined and computationally estimated log K_{OC} values

No.	Compound	TLC									HPLC	LSER		EPI Suite	
		PCR	PLS	RP18/ MeOH 70% v/v	RP18/ MeOH 80% v/v	CN/ MeOH 40% v/v	CN/ MeOH 50% v/v	CN/ MeOH 6 % v/v	CN/ MeOH 70% v/v	CN/ ACN 50% v/v		Poole	Nguyen	MCI	KOW
<i>Standard compounds</i>															
1	Phenol	1.41	1.47	1.63	1.62	1.57	1.50	1.57	1.71	1.31	1.49	1.65	1.58	2.27	1.90
2	4-Nitrophenol	1.94	2.07	1.88	1.88	2.30	2.28	2.33	2.44	1.75	1.40	1.85	2.18	2.46	2.37
3	Benzyl Alcohol	1.17	1.20	1.66	1.73	1.22	1.19	1.28	1.24	1.08	0.66	1.44	1.32	1.33	1.12
4	Naphthylamine	2.83	2.83	2.42	2.41	2.80	2.74	3.06	3.15	2.98	3.41	2.67	2.53	3.40	1.95
5	1-Naphthol	2.63	2.64	2.51	2.47	2.93	2.78	2.83	2.94	2.55	3.25	3.02	2.71	3.31	2.67
6	2,4-Dichlorophenol	2.56	2.56	2.68	2.62	2.74	2.66	2.61	2.51	2.44	2.54	2.62	2.48	2.69	2.78
7	Anthracene	4.38	4.42	4.31	4.24	4.39	4.46	4.59	4.28	4.48	4.06	4.46	4.27	4.21	3.86
8	Acetophenone	1.97	1.96	2.19	2.18	1.83	1.99	1.94	1.96	1.92	2.31	1.67	1.81	1.71	1.99
9	2,4,6-Trichlorophenol	2.97	2.96	2.97	3.12	2.99	2.94	2.97	2.73	3.13	3.02	3.01	2.94	3.25	3.28
10	Ethyl- <i>p</i> -hydroxybenzoate	1.77	1.79	2.19	2.12	1.97	2.00	1.84	1.76	1.78	2.43	2.23	2.30	2.20	2.39
11	<i>p</i> -Anisidine	2.87	2.96	1.66	1.73	2.45	2.97	3.43	3.76	3.15	0.06	1.42	1.52	1.65	1.48
12	1,2,3-Benzotriazole	1.13	1.21	1.49	1.41	1.25	1.36	1.32	1.55	1.12	0.21	1.65	1.82	1.72	1.79
13	Diphenylamine	3.80	3.76	3.14	3.09	3.73	3.72	3.48	3.69	4.07	2.79	3.35	3.14	2.92	2.82
14	2,2'-Dipiridylum	2.16	2.06	2.39	2.49	1.67	1.86	1.80	2.03	2.30	2.97	1.93	1.93	3.09	2.11
15	4-Bromoaniline	2.50	2.49	2.29	2.27	2.40	2.47	2.44	2.80	2.64	2.46	2.30	2.38	2.05	2.15
16	Benzophenone	2.51	2.50	3.06	3.04	3.02	2.82	2.67	2.81	2.28	3.91	3.04	3.21	3.06	2.88
17	4-Aminobenzoic acid	1.40	1.46	0.77	0.59	1.01	1.03	1.02	1.22	1.84	0.39	1.29	1.33	0.90	0.59
18	Pyrene	4.81	4.83	4.59	4.63	4.97	4.59	4.59	4.70	4.93	4.51	5.02	4.84	4.74	4.24
19	Benzo[<i>a</i>]pyrene	5.57	5.73	6.28	5.81	5.19	5.59	5.66	5.54	5.50	4.95	6.31	6.00	5.77	5.32
20	Fluorene	4.00	4.00	4.02	4.04	4.22	3.81	3.78	3.83	3.79	4.61	3.72	3.63	3.96	3.63
21	Acenaphthene	3.77	3.69	3.66	3.58	4.04	3.64	3.69	3.72	3.67	4.25	3.60	3.51	3.70	3.40
22	Naphthalene	3.60	3.51	3.29	3.38	3.95	3.52	3.64	3.65	3.60	3.51	3.03	2.99	3.19	2.86
23	Phenanthrene	4.23	4.20	4.19	4.23	4.47	4.22	4.14	4.03	3.90	5.38	4.21	4.08	4.22	3.87
24	Dibenz[<i>a,h</i>]anthracene	5.42	5.47	4.92	5.02	5.09	5.61	5.53	5.41	5.24	6.01	6.79	6.49	6.28	5.68
25	Benz[<i>a</i>]anthracene	5.16	5.23	5.08	5.29	4.85	5.47	5.19	5.04	4.71	5.16	5.51	5.33	5.25	5.00
26	Simazine	2.01	1.95	2.36	2.40	2.03	2.04	2.09	1.96	1.95	1.94	2.52	2.15	2.17	2.18
27	Propazine	2.78	2.61	2.91	2.91	2.56	2.34	2.49	2.09	2.67	2.64	3.07	2.65	2.54	2.34
28	Ametryn	2.65	2.53	3.00	3.12	2.45	2.54	2.38	2.11	2.95	3.02	2.90	2.54	2.63	2.36
29	Prometryn	2.89	2.77	3.35	3.37	2.74	2.74	2.51	2.30	3.06	3.31	3.18	2.77	2.82	3.51

TABLE S4 Continues

No.	Compound	TLC									HPLC	LSER		EPI Suite		
		PCR	PLS	RP18/ MeOH 70% v/v	RP18/ MeOH 80% v/v	CN/ MeOH 40% v/v	CN/ MeOH 50% v/v	CN/ MeOH 60% v/v	CN/ MeOH 70% v/v	CN/ ACN 50% v/v		Poole	Nguyen	MCI	KOW	
	<i>Unknowns</i>															
30	3-Nitrophenol	2.33	2.43	1.76	1.96	2.08	2.19	2.01	1.93	1.22	1.77	1.99	2.24	2.46	2.42	
31	2-Naphthol	2.81	2.82	2.26	2.36	2.50	2.54	2.35	2.28	1.84	3.19	2.94	2.64	3.30	2.59	
32	4-Hydroxybenzaldehyde	1.58	1.69	1.13	1.39	1.39	1.38	1.26	1.34	0.53	1.14	1.66	1.83	1.15	1.61	
33	2-Aminophenol	2.56	2.66	1.04	1.15	1.69	2.04	2.19	2.47	2.10	1.10	1.35	1.17	1.96	1.41	
34	4- <i>t</i> -Buthylphenol	3.30	3.13	2.74	2.80	2.44	2.33	1.96	1.66	2.71	3.03	2.71	2.53	3.11	2.92	
35	2,6-Dimethylphenol	3.09	3.00	2.24	2.37	2.03	2.15	1.88	1.88	2.60	2.06	2.21	2.05	2.70	2.40	
36	4-Methoxyphenol	1.99	2.04	1.23	1.47	1.41	1.48	1.57	1.61	1.31	1.18	1.65	1.65	2.08	2.02	
37	Methyl- <i>p</i> -hydroxybenzoate	2.01	2.03	1.60	1.80	1.64	1.67	1.37	1.57	1.23	2.41	1.94	2.03	1.94	2.11	
38	2-Nitrobenzaldehyde	2.69	2.61	1.59	1.74	1.76	1.97	1.74	2.09	2.07	1.99	1.86	2.33	1.24	1.88	
39	3-Nitrobenzaldehyde	2.77	2.75	1.76	1.92	1.96	2.17	1.98	2.25	2.03	1.86	1.71	2.08	1.24	1.73	
40	Phthalimide	1.85	1.90	1.37	1.60	1.45	1.63	1.34	1.30	1.10	1.61	1.54	2.09	0.15	1.10	
41	Oxazepam	2.36	2.35	2.42	2.43	2.09	2.14	1.83	1.72	1.52	3.35	3.37	2.27	2.76	1.73	
42	Lorazepam	2.48	2.43	2.37	2.25	2.22	2.19	1.89	1.74	1.73	3.79	3.64	2.58	2.98	1.81	
43	Clonazepam	2.99	2.98	2.21	2.29	2.61	2.88	2.50	2.41	2.23	3.26	3.47	2.86	4.06	2.45	
44	Bromazepam	2.37	2.30	2.17	2.29	1.80	1.87	1.64	1.75	1.58	2.95	2.95	2.12	3.56	2.21	
45	Diazepam	3.57	3.48	3.17	3.25	2.73	2.92	2.55	2.47	2.82	4.26	3.55	3.17	3.88	2.44	
46	Nitrazepam	2.98	3.01	2.26	2.29	2.61	2.85	2.62	2.60	2.14	3.11	3.43	3.32	3.85	2.36	
47	Chlordiazepoxide	2.99	3.00	2.24	2.33	2.61	2.79	2.62	2.56	2.10	3.54	4.21	3.98	5.48	2.23	
48	Clobazam	3.20	3.13	2.39	2.45	2.61	2.85	2.60	2.48	2.50	3.92	2.82	2.93	2.54	2.03	
49	Medazepam	6.33	6.37	4.78	4.78	5.35	5.30	5.62	5.51	4.89	5.56	4.61	4.37	4.48	3.30	
50	Chrysene	6.66	6.79	5.64	5.71	6.02	6.39	5.29	4.65	5.36	4.34	5.51	5.33	5.26	5.04	

TABLE S5a Comparison of chromatographic and *in silico* methods;

Collection of normalized SRD values

Method	SRD
TLC (CN/MeOH_40)	9.60
TLC (CN/MeOH_50)	10.72
LSER (Poole)	12.80
TLC (RP18/MeOH_80)	13.60
TLC (RP18/MeOH_70)	13.76
TLC (CN/MeOH_60)	14.08
LSER (Nguyen)	15.36
TLC (PCR)	16.48
TLC (PLS)	17.60
HPLC	18.08
TLC (CN/MeOH_70)	19.04
EPI (MCI)	19.84
TLC (CN/ACN_50)	20.00
EPI (KOW)	21.60

TABLE S5b Normalized SRD values obtained by the sevenfold jack-knife resampling procedure

Method	SRD1	SRD2	SRD3	SRD4	SRD5	SRD6	SRD7	Median
TLC (CN/MeOH_40)	8.8	9.7	11.7	10.0	10.0	10.6	7.6	9.96
TLC (CN/MeOH_50)	11.3	8.7	13.2	11.5	10.6	10.4	10.4	10.61
LSER (Poole)	13.6	11.3	14.7	14.3	11.7	11.9	13.2	13.20
TLC (RP18/MeOH_80)	13.4	12.6	18.2	13.4	13.4	13.2	12.6	13.38
TLC (RP18/MeOH_70)	13.6	13.4	18.0	13.2	13.4	13.2	12.1	13.42
TLC (CN/MeOH_60)	13.8	13.2	18.4	15.2	12.6	12.8	14.1	13.83
LSER (Nguyen)	16.6	14.1	17.3	16.2	15.4	12.3	15.8	15.80
TLC (PCR)	17.0	16.0	19.3	18.2	14.3	13.4	17.5	17.01
TLC (PLS)	18.6	16.5	20.3	17.7	14.5	15.2	19.3	17.75
HPLC	18.4	16.5	20.6	19.3	19.0	16.0	18.6	18.61
TLC (CN/MeOH_70)	19.5	17.3	24.2	19.7	17.3	17.3	19.3	19.26
EPI (MCI)	20.4	16.7	21.6	20.8	19.9	19.3	19.5	19.91
TLC (CN/ACN_50)	21.5	18.4	22.3	22.3	18.6	18.0	18.8	18.83
EPI (KOW)	21.1	20.6	26.8	23.8	19.9	20.8	20.1	20.78

TABLE S6 LSER regression coefficients of TLC systems given as the value \pm standard deviation; I – intercept, a – hydrogen bond basicity, b – hydrogen bond acidity, s – dipolarity/polarizability, e – molar refraction in excess, and ν – ability for formation of a vacant space; Stationary phase octadecyl (RP18)-modified silica and cyano (CN)-modified silica; Mobile phases: mixtures of methanol (MeOH) and acetonitrile (ACN) with water in different proportions (v/v); Statistical parameters: R^2 – squared determination coefficient, $S.D.$ – standard error of the model, F – Fisher’s statistic, P – significance of F , n – number of calibration points (compounds)

No.	Chrom.system	I	a	b	s	e	ν	R^2	$S.D.$	F	P	n
1	MeOH 40%	0.72 \pm 0.19	-0.50 \pm 0.15	-0.660 \pm 0.13	-0.52 \pm 0.12	0.030 \pm 0.12	1.30 \pm 0.20	0.8314	0.248	43.39	6.36 \cdot 10 ⁻¹⁶	50
2	MeOH 50%	0.47 \pm 0.18	-0.52 \pm 0.15	-0.876 \pm 0.13	-0.59 \pm 0.14	0.161 \pm 0.11	1.29 \pm 0.19	0.8671	0.241	57.40	3.60 \cdot 10 ⁻¹⁸	50
3	MeOH 60%	0.04 \pm 0.15	-0.40 \pm 0.12	-0.942 \pm 0.10	-0.461 \pm 0.094	0.181 \pm 0.085	1.21 \pm 0.16	0.8934	0.200	73.75	2.92 \cdot 10 ⁻²⁰	50
4	MeOH 70%	-0.26 \pm 0.14	-0.37 \pm 0.11	-1.00 \pm 0.09	-0.417 \pm 0.084	0.352 \pm 0.076	0.94 \pm 0.14	0.9159	0.178	95.84	1.64 \cdot 10 ⁻²²	50
5	MeOH 80%	-0.41 \pm 0.10	-0.32 \pm 0.08	-0.81 \pm 0.07	-0.307 \pm 0.064	0.245 \pm 0.057	0.77 \pm 0.11	0.9220	0.135	104.09	3.12 \cdot 10 ⁻²³	50
6	ACN 40%	0.49 \pm 0.13	-0.57 \pm 0.10	-1.00 \pm 0.09	-0.352 \pm 0.079	0.156 \pm 0.072	1.05 \pm 0.13	0.9185	0.168	99.20	8.24 \cdot 10 ⁻²³	50
7	ACN 50%	0.49 \pm 0.14	-0.67 \pm 0.11	-0.93 \pm 0.09	-0.409 \pm 0.084	0.226 \pm 0.075	0.75 \pm 0.14	0.9032	0.177	82.13	3.54 \cdot 10 ⁻²¹	50
8	ACN 60%	0.01 \pm 0.12	-0.67 \pm 0.09	-0.875 \pm 0.08	-0.318 \pm 0.073	0.193 \pm 0.066	0.81 \pm 0.12	0.9230	0.154	105.41	2.42 \cdot 10 ⁻²³	50
9	ACN 70%	-0.40 \pm 0.11	-0.665 \pm 0.09	-0.789 \pm 0.07	-0.313 \pm 0.066	0.198 \pm 0.060	0.80 \pm 0.11	0.9331	0.140	122.83	1.08 \cdot 10 ⁻²⁴	50
10	ACN 80%	-0.50 \pm 0.13	-0.68 \pm 0.10	-0.725 \pm 0.09	-0.347 \pm 0.079	0.177 \pm 0.071	0.75 \pm 0.13	0.8986	0.167	77.95	9.90 \cdot 10 ⁻²¹	50
11	MeOH 40%	-0.01 \pm 0.15	-0.35 \pm 0.12	-1.02 \pm 0.10	-0.298 \pm 0.090	0.349 \pm 0.081	0.80 \pm 0.15	0.8957	0.191	75.56	1.82 \cdot 10 ⁻²⁰	50
12	MeOH 50%	-0.39 \pm 0.12	-0.30 \pm 0.10	-0.820 \pm 0.08	-0.138 \pm 0.075	0.313 \pm 0.067	0.62 \pm 0.13	0.8983	0.158	77.74	1.04 \cdot 10 ⁻²⁰	50
13	MeOH 60%	-0.40 \pm 0.11	-0.273 \pm 0.09	-0.632 \pm 0.07	-0.142 \pm 0.068	0.262 \pm 0.061	0.41 \pm 0.11	0.8648	0.143	56.28	5.23 \cdot 10 ⁻¹⁸	50
14	MeOH 70%	-0.406 \pm 0.094	-0.271 \pm 0.08	-0.452 \pm 0.06	-0.081 \pm 0.058	0.227 \pm 0.052	0.19 \pm 0.10	0.8351	0.123	44.56	3.92 \cdot 10 ⁻¹⁶	50
15	ACN 40%	-0.038 \pm 0.14	-0.28 \pm 0.11	-0.661 \pm 0.09	-0.222 \pm 0.084	0.160 \pm 0.076	0.62 \pm 0.14	0.8037	0.178	36.03	1.71 \cdot 10 ⁻¹⁴	50
16	ACN 45%	0.040 \pm 0.13	-0.26 \pm 0.10	-0.710 \pm 0.09	-0.269 \pm 0.077	0.192 \pm 0.070	0.49 \pm 0.13	0.8218	0.164	40.58	2.13 \cdot 10 ⁻¹⁵	50
17	ACN 50%	-0.143 \pm 0.093	-0.296 \pm 0.07	-0.448 \pm 0.06	-0.166 \pm 0.057	0.160 \pm 0.052	0.327 \pm 0.10	0.8338	0.121	44.15	4.64 \cdot 10 ⁻¹⁶	50
18	ACN 55%	-0.242 \pm 0.089	-0.279 \pm 0.07	-0.431 \pm 0.06	-0.188 \pm 0.055	0.168 \pm 0.049	0.306 \pm 0.10	0.8375	0.116	45.35	2.85 \cdot 10 ⁻¹⁶	50
19	ACN 60%	-0.409 \pm 0.088	-0.260 \pm 0.07	-0.371 \pm 0.06	-0.116 \pm 0.054	0.066 \pm 0.049	0.340 \pm 0.089	0.7761	0.115	30.50	2.95 \cdot 10 ⁻¹³	50

REFERENCES

A: List of literature sources of $\log K_{OC}$ values

- [A1] G. Schüürmann, R. Ebert, R. Kühne, Prediction of the sorption of organic compounds into soil organic matter from molecular structure, *Environ. Sci. Technol.* 40 (22) (2006) 7005–7011.
- [A2] W. Meylan, P. H. Howard, R. S. Boethling, Molecular topology/fragment contribution method for predicting soil sorption coefficients, *Environ. Sci. Technol.* 26 (1992) 1560–1567.
- [A3] T. H. Nguyen, K. Goss, P. W. Ball, Polyparameter linear free energy relationships for estimating the equilibrium partition of organic compounds between water and the natural organic matter in soils and sediments, *Environ. Sci. Technol.* 39 (2005) 913–924.
- [A4] B. Von Oepen, W. Kördel, W. Klein, G. Schüürmann, Predictive QSPR models for estimating soil sorption coefficients: potential and limitations based on dominating processes, *Sci. Total Environ.* 109/110 (1991) 343–354.

B: List of literature sources of pK_a values

- [B1] J. Jover, R. Bosque, J. Sales, Neural Network Based QSPR Study for Predicting pK_a of Phenols in Different Solvents, *QSAR Comb. Sci.* 26(3) (2007) 385–397.
- [B2] R.C. Weast (Ed.), *Handbook of Chemistry and Physics*. 69th ed., CRC Press Inc., Boca Raton, FL, 1988-1989, pp. D-160.
- [B3] E.P. Serjeant, B. Dempsey, *Ionisation Constants of Organic Acids in Aqueous Solution*, International Union of Pure and Applied Chemistry (IUPAC). IUPAC Chemical Data Series No. 23, Pergamon Press, Inc., New York, 1979.
- [B4] M. Boyce, E. Spickett, Separation and quantification of preservatives using ion Pair HPLC and CZE, *J. Chem. Educ.* 77 (6) (2000) 740–742.
- [B5] D.R. Lide, *CRC Handbook of Chemistry and Physics* 88th Edition 2007-2008. CRC Press, Taylor & Francis, Boca Raton, FL 2007, pp. 8-47.
- [B6] J. Sangster, Octanol-water partition coefficients of simple organic compounds, *J. Phys. Chem.* 18 (1989) 1111-1230.
- [B7] D.D. Perrin, The effect of temperature on pK values of organic bases, *Aust. J. Chem.* 17 (1964) 484-488.
- [B8] D.D. Perrin, *Dissociation constants of organic bases in aqueous solution*. IUPAC Chem Data Set, Butterworth, London, 1965.

- [B9] G. Kortüm, W. Vogel, K. Andrussov, Dissociation Constants of Organic Acids in Aqueous Solution, International Union of Pure and Applied Chemistry, Butterworth, London, 1961.
- [B10] C.D.S. Tomlin, The Pesticide Manual - World Compendium, 11th ed., British Crop Protection Council, Surrey, England, 1997.
- [B11] C.D.S. Tomlin, The e-Pesticide Manual, 13th Edition Version 3.1, British Crop Protection Council, Surrey, UK, 2004.
- [B12] D. Bhattacharyya, C.E. Hamrin Jr., R.P. Northey, Oxidation of Hazardous Organics in a Two-Phase Fluorocarbon-Water System, Haz. Waste Haz. Mater. 3 (1986) 405-429.
- [B13] S. Budavari, The Merck Index - Encyclopedia of Chemicals, Drugs and Biologicals, Rahway, Merck and Co., Inc., New York, 1989, pp. 1170.
- [B14] S.H. Hilal, L.A. Carreira, S.W. Karickhoff, Estimation of chemical reactivity parameters and physical properties of organic molecules using SPARC, in: P. Politzer, J.S. Murray, Quantitative Treatments of Solute/Solvent Interactions: Theoretical and Computational Chemistry Vol. 1, Elsevier, New York, 1994, pp. 291-353.
- [B15] J. Barrett, W. Franklin Smith, I.E. Davidson, An examination of acid-base equilibria of 1,4-benzodiazepines by spectrophotometry, J. Pharm. Pharmacol. 25 (1973) 387-393.
- [B16] S.A. Kaplan, K. Alexander, M.L. Jack, C.V. Puglisi, J.A.F. de Silva, T.L. Lee, R.E. Weinfeld, Pharmacokinetic profiles of clonazepam in dog and humans and of flunitrazepam in dog. J. Pharm. Sci. 63 (1974) 527-532.
- [B17] D.L. Sorby, E.M. Plein, J.D. Benmaman, Adsorption of phenothiazine derivatives by solid adsorbents, J. Pharm. Sci. 58 (1966) 785-794.
- [B18] E. Van der Kleijn, Protein binding and lipophilic nature of ataractics of the meprobamate- and diazepam-group, Arch. Int. Pharmacodyn. Ther. 179 (1969) 225-250.
- [B19] E. Souri, A.D. Farahani, R. Ahmadkhaniha, M. Amini, A stability indicating HPLC method for the determination of clobazam and its basic degradation product characterization, DARU, J. Pharm. Sci. 22 (2014) 49.
- [B20] G.F. le Pettit, Medazepam pKa determined by spectrophotometric and solubility methods, J. Pharm. Sci. 65 (1976) 1094-1095.