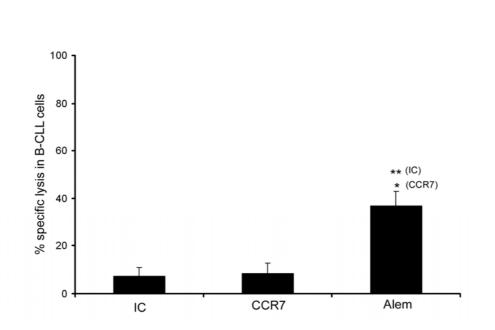
Supplementary 7	Table 1.	Characteristics of the studied	patient samples
-----------------	----------	--------------------------------	-----------------

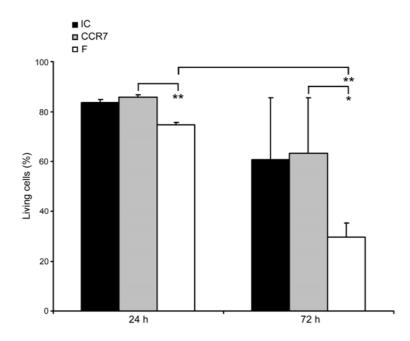
Patient	Rai			Hb	Leukocye	Platelet
number	Stage	Sex	Age (y)	(g/dL)	count	count
number	Stage			(g/uL)	(10 <sup>3</sup> /mm <sup>3</sup> )	(10 <sup>3</sup> /mm <sup>3</sup> )
1	Ι	F	80	11.9	17.7	231
2	0	М	55	14.2	15.5	155
3	III	F	81	11.8	97.2	93
4	IV	М	40	10.3	19.22	3
5	IV	М	81	10.1	10.6	2
6	0	F	89	13.7	11.6	190
7	Ι	F	76	na	na	na
8	III	М	58	11.3	12.4	122
9	IV	F	84	10.9	23.3	24
10	IV	М	73	12.8	19.9	145
11	IV	М	59	na	na	na
12	IV	М	60	12.0	7.0	75
13	IV	М	53	14.2	6.9	70
14	IV	М	74	12.3	74.2	176
15	0	М	50	15.9	70.5	150
16	IV	М	60	12.3	6.2	83
17	Ι	М	60	15.3	57.7	179
18	Ι	М	48	16.3	14.1	136
19	IV	F	79	12.2	101.7	93
20	0	F	73	13.9	5.5	226
21	Ι	F	57	16	17.45	161
22	0	F	72	15.2	11.84	297
23	0	М	55	14.2	15.13	155

The table presents the clinical features of the studied patients. F, female; M, male; y, years; Hb, hemoglobin; na, not available.



Supplementary Figure 1. Mouse anti-human CCR7 antibodies do not mediate antibodydependent cell-mediated cytotoxicity (ADCC)

CLL cells were incubated with media alone or in the presence of an isotype control (IC), anti-CD52 (alemtuzumab, Alem) or anti-CCR7 antibodies. Isolated and IL-2-stimulated NK cells from healthy donors were used as effector cells at a effector to target (E:T) ratio of 6:1. After 4h, the percentage of CLL cells killed by ADCC was determined by FCM as described in the Material and Methods section. Data were normalized to the media control. Columns represent the mean of three independent experiments; bars represent mean standard error. \*, p<0.05; \*\*, p<0.01.



Supplementary Figure 2. Mouse anti-human CCR7 antibodies do not mediate direct proapoptotic effects in CLL cells

CLL cells were incubated in the presence of either an isotype control antibody (IC), anti-CCR7 antibody (CCR7) or fludarabine as positive control (F). Cell viability was determined after 24 and 72 hours by double staining with FITC-labelled Annexin-V (AV) and 7-AAD. Flow cytometry analysis and quantification of living cells (7-AAD<sup>-</sup> AV<sup>-</sup> cells) was performed. The percentage of viable living cells is shown. Results are expressed as mean of three different experiments  $\pm$  mean standard error. \*, p<0.05; \*\*, p<0.01.