Soluble antigen binds to the B-cell antigen receptor and is internalized for subsequent processing and the presentation of antigen-derived peptides to T cells. Many antigens are not soluble, however, but are integral components of membrane; furthermore, soluble antigens will usually be encountered in vivo in a membrane-anchored form, tethered by Fc or complement receptors. Here we show that B-cell interaction with antigens that are immobilized on the surface of a target cell leads to the formation of a synapse and the acquisition, even, of membrane-integral antigens from the target. B-cell antigen receptor accumulates at the synapse, segregated from the DC45 coreceptor which is excluded from the synapse, and there is a corresponding polarization of cytoplasmic effectors in the B cell. B-cell antigen receptor mediates the gathering of antigen into the synapse and its subsequent acquisition, therebypotentiating antigen processing and presentation to T cells with high efficiency. Synapse formation and antigen acquisition will probably enhance the activation of B cells at low antigen concentration, allow context-dependent antigen recognition and enhance the linking of B- and T-cell epitopes.

To investigate the B-cell response to antigen encountered as part of an immune complex tethered to a cell surface, immune complexes comprising hen-egg lysozyme (HEL) aggregated with specific immunoglobulin-\(\gamma\) (IgG) monoclonal antibodies were loaded on to the surface of an Fcy receptor (FcyR)-expressing myeloid cell line. The immune complexes had a patchy distribution over the myeloid cell surface (Fig. 1a, b); however, on incubation with antigen-specific B cells, cell aggregates were formed in which the immune complexes on the myeloid cell and the B-cell antigen receptor (BCR) on the B cell were gathered together into a region of synapsis (Fig. 1c, d).

Similar results were obtained using immune complexes loaded onto FcyRII-expressing I-cell transfectedants. Immunocytochemistry suggested that there is a gathering of tethered immune complexes, mediated by the BCR (possibly aided by the oligomeric nature of the BCR\(^2\)) and accompanied by an apparent reorganization of the B-cell surface, as judged by segregation of the BCR from CD45 in the region of synapsis (Fig. 1e).

This concentration of BCR is reminiscent of the capping of surface IgM that results from incubation of B cells with polyclonal anti-IgM antisera\(^6,7\). We therefore determined whether the reorganization of the B-cell membrane was also contributed by the BCR (possibly aided by the oligomeric nature of the BCR\(^2\)) and accompanied by an apparent reorganization of the B-cell surface, as judged by segregation of the BCR from CD45 in the region of synapsis (Fig. 1e).

Reorganization of components of the B-cell membrane was also evident from a depletion of CD22 from the centre of most synapses (although often concentrated at the edges), where there was a concentration of ganglioside GM1 (which is associated classically with many Src-family tyrosine kinases). There was also polarization of cytoplasmic components, as judged by a depletion of the signal-inhibitory phosphatase SHP1 in the region of the synapse (Fig. 2d–f), but a concentration of phosphotyrosine-containing proteins as well.
as actin and phospholipase C (PLC)-γ2 (Fig. 2d). This gross reorganization of a B cell after interacting with a target cell may reflect changes that also occur (but possibly on a smaller scale) after BCR crosslinking by soluble antigen, as such crosslinking\(^8\) leads to BCR translocation into a GM1-enriched fraction that is depleted of CD22 and CD45.

We wanted to determine whether reorganization of the B cell could be detected using targets displaying a lower density of a lower-affinity antigen. Transfectants expressing a membrane-integral form of a mutated HEL (mHEL\(^*\)) that exhibits a 100-fold reduced affinity for the H\(\text{yHEL10}\) BCR\(^10\) were screened for low-expressing clones: J[mHEL\(^*\)]\(^8\) displays only around \(9 \times 10^3\) mHEL\(^*\) molecules per cell (compared with J[mHEL]\(^6\), which expresses about \(4 \times 10^5\) mHEL per cell).

After incubation of J[mHEL\(^*\)]\(^8\) with HEL-specific B cells, staining for mHEL\(^*\) (which was previously only dim) was now brightly focused in the synapse (Fig. 3a). This quantity of mHEL\(^*\) antigen is only sufficient to bring a small proportion of the total BCR into the synapse; however, there is nevertheless a local reorganization of the B cell as the BCR gathering is accompanied by local exclusion of CD45. Notably, the low abundance of mHEL\(^*\) on the target cell surface is sufficient to trigger B-cell activation, as judged by the upregulation of CD66. Indeed, surface-tethered mHEL\(^*\) is several orders of magnitude more efficacious than soluble HEL\(^*\) in this respect (Fig. 3b, c). This is consistent with the apparent increased potency of membrane (as opposed to soluble) antigens in determining B-cell fate \(\text{in vivo}^{11,12}\), which probably reflects the effective affinity increase resulting from restricting antigen mobility to two dimensions, coupled with the ability to concentrate antigen into the synapse\(^13\).

To follow synapse formation in real time and exclude the possibility of a fixation artefact, we established targets that displayed mHEL fused to green fluorescent protein (GFP). Incubation with HEL-specific B cells led to rapid aggregation of mHEL–GFP in the synapse (Fig. 3d), although the frequency of synapses decreased at later time points (data not shown). The B cells bind well to their targets even at \(13^\circ\text{C}\), but formation of a synapse at which BCR and antigen have visibly accumulated requires a higher temperature—consistent with a need for membrane reorganization (Fig. 3e). Similar kinetics of synapse formation were also observed in a different B-cell/target interaction system—that of B cells carrying the 3-83 BCR specific for major histocompatibility complex (MHC) class I molecules with targets that express H2-K\(^\beta\) (Fig. 3f; BCR/antigen affinity estimated\(^14:\) to be about \(10^6\times M^{-1}\)).

We wondered whether the decrease in observable synapses over time reflected internalization or degradation of antigen–BCR. In T cells, rapid internalization of the antigen receptor is observed after their interaction with antigen-presenting cells\(^15\). After culturing HEL-specific B cells with mHEL targets, flow cytometry showed that there was a rapid downregulation of BCR on the surface of the sorted B cells—more rapid than with soluble antigen (Fig. 4a, b).

This downregulation was peculiar to the BCR and did not apply to the CD19, CD22 or CD45 co-receptors; in addition, experiments performed using B-cell transfectants rather than splenic B cells showed that the downregulation occurring with the antigen-specific BCR did not extend to an irrelevant BCR expressed on the same cell (Fig. 4c). This loss of detectable surface BCR might reflect its internalization by the B cell, its acquisition by the target cell or some form of masking. Immunocytochemistry of cells sorted after co-culture showed that IgM was readily detectable in the sorted B cells (but not the sorted target cells), if they were permeabilized before staining (Fig. 4d). This IgM is unlikely to be \(\text{de novo}\) synthesized IgM that has not yet been transported to the cell surface, as protease stripping of surface IgM on splenic HEL-specific B cells showed that the intracellular pool of IgM was very small. Moreover, co-staining for IgM and HEL showed that, after B-cell–mHEL target interaction, the permeabilized, sorted B cells stained brightly for the

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**Figure 1** Synapse formation between B cell and antigen-displaying target cell. a, Loading of HEL–IgG immune complexes (ICs) onto the 279.AC1 myeloid cell line\(^2\) or L-cell transfectants expressing human Fc\(\gamma\)RI. ICs were detected by staining for HEL and IgG1. L, L cells; AC, 279.AC1 cells. b, ICs are patchily distributed on the surface of 279.AC1 (top) or L[Fc\(\gamma\)RI] cells (bottom) that have been loaded at 4°C, fixed and stained for mouse IgG1. Projection views are shown on right. c, Synapse formation after incubation (10 min, 37°C) of IC-loaded targets with excess HEL-specific splenic B cells. For both myeloid and L cells, a single optical section of a single B-cell/target cell interaction is shown in either single or merged view. Whereas most B cells were in contact with a target when IC-loaded L[Fc\(\gamma\)RI] cells were used, only 10% of the B cells formed aggregates with IC-loaded 279.AC1 cells, correlating with internalization of the ICs at 37°C by the myeloid cells—reflecting their expression of Fc\(\gamma\)RII and Fc\(\gamma\)RIII (not shown). d, Projection views of HEL-specific splenic B cells interacting with IC-loaded L[Fc\(\gamma\)RI] cells. Cells were double stained with anti-mouse IgG1 (for ICs; green) and anti-mouse IgM (for BCR; red). e, Comparison of BCR (red) and CD45R(B220) (green) distribution on HEL-specific B cells interacting with IC-loaded 279.AC1 or L[Fc\(\gamma\)RI] targets; a single optical section is shown in each case.
Figure 2  Polarization of the B cell after encountering membrane-immobilized antigen. a, Abundance of surface HEL in the cloned J558 transfectants (J[mHEL]6, J[mHEL+]8 and J[mHEL–GFP]) monitored by staining with anti-HEL IgG1-κ (HyHEL5) and PE-conjugated anti-κ. b, Distribution of BCR (IgM or IgD), CD22 and CD45 on HEL-specific splenic B cells interacting with J[mHEL]6 targets viewed as single optical sections. Cells were fixed after a 10-min incubation at 37 °C. c, Segregation of IgM from CD45 and CD22. Samples were prepared as in a but co-stained for CD45 or CD22 (green) and IgM (red). d, Polarization of HEL-specific B cells interacting with J[mHEL]6 targets analysed by double staining, in each case showing a single confocal section in single or merged view. Ganglioside GM1 was detected using cholera toxin B subunit (CTB). See Supplementary Information for additional staining for actin and PLC-γ2. e, Views along the plane of synapsis co-staining for IgM (red) and HEL, phosphotyrosine or SHP1 (green), showing coincidence of HEL–IgM and phosphotyrosine–IgM at the synapse but exclusion of SHP1. f, Quantification of the depletion of CD45, CD22 and SHP1 at synapses. Left, the mean extent of depletion of the appropriate marker at the synapse (from analysis of an average of 50 synapses) is expressed relative to the abundance of that marker adjacent to the synapse but in the same confocal plane. Right, the percentage of antigen-specific synapses (‘HEL+’) in which such depletion was noted (CD45, below 80%; CD22 and SHP1, below 60%) is compared with depletion in contacts between B cells and antigen-lacking targets (HEL–).
cells with upregulated expression of CD86. numbers of J558[mHEL+] targets, and activation was monitored as the percentage of B

\[ a \]

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**Figure 3** Parameters affecting synapse formation. **a**, Synapse formation between HEL-specific splenic B cells and J[mHEL+] targets (see Fig. 2a), which express antigen of reduced affinity at low density. Top, single or merged view of cells stained for HEL and IgM. Bottom, staining for CD45 and IgM, showing a single optical section. **b**, Culture of HEL-specific splenic B cells (20 h) with either J[mHEL+] (twofold excess) or soluble HEL (0.1 μg ml⁻¹) causes similar B-cell activation as judged by upregulation of CD86. **c**, Comparison of B-cell activation by membrane-tethered and soluble HEL. HEL-specific splenic B cells were cultured with varying concentrations of soluble HEL+ or varying numbers of J558[mHEL+] targets, and activation was monitored as the percentage of B cells with upregulated expression of CD86. **d**, Real-time microscopy of an HEL-specific splenic B cell interacting at 20 °C with an HEL→GFP-expressing target. **e**, Temperature dependence of synapse formation, as judged by the percentage of B cells bound to a target in which 50% of the IgM in one optical section was concentrated in the region of synapsis. **f**, Synapse frequency as a function of time. The experiment shown (as in **e** but at 37 °C) was performed using H2Kβ-specific splenic B cells from 3-83 mice and H2Kβ hybridoma targets. Shown is a single-plane confocal image of such a synapse (co-stained for IgM, red; H2Kβ, green).

 antigen, indicating that downregulation of the BCR might reflect acquisition of the antigen from the target cell and internalization of the mHEL→BCR complex by the B cell (Fig. 4e).

If the B cell acquires the antigen from the target cell, this should be visible in living cells using mHEL→GFP targets, without the risk of flow cytometry or fixation-associated artefacts. This is indeed the case. Following the interaction at 20 °C in real time shows that synapse formation is followed by the appearance of GFP fluorescence at points in the B cell that are distant from the synapse (Fig. 5a); this effect is more marked if the cells are incubated at 37 °C (Fig. 5b). Furthermore, if these cells are fixed and permeabilized, antigen transfer is evident from the fact that serial sections reveal co-localization of mHEL→GFP and IgM at sites in the B cell that are both removed from, and in a different plane to that of the synapse (Fig. 5c; Supplementary Information movie 2). The mHEL→GFP that has transferred to the B cell seems to undergo degradation, as the GFP fluorescence diminishes with time (Fig. 5d). The ability of the B cell to acquire an integral membrane antigen from the target cell and then internalize and degrade it should presumably allow the B cell to present antigen-derived peptides to T cells. HEL-specific B-cell transfectants co-cultured with H2Kβ-negative mHEL targets are very effective in presenting HEL-derived peptides to T cells (Fig. 5e). This presentation does not simply reflect BCR-mediated uptake of soluble mHEL fragments that have been released spontaneously by the target cells, as presentation is abolished if the B cells and target cells are separated by a mesh.

This presentation of membrane-tethered antigens is impressively efficient: co-incubation of 10⁵ HEL-specific B cells with only 3×
$10^5$ mHEL-expressing target cells (which correspond to a total of some $10^8$–$10^9$ surface-expressed mHEL molecules) is sufficient to yield detectable interleukin (IL)-2 production in the antigen-presentation assay. In contrast, presentation of soluble HEL under similar assay conditions requires more than $10^{11}$ molecules of antigen. The increased efficacy of presentation of tethered (as opposed to soluble) antigen is even more striking when using mutated HELs exhibiting lower affinity for the BCR (data not shown).

The precise nature of the synapse is at present unclear, although co-staining with clathrin and early endosomal antigen suggests that much of the BCR–antigen complex may be internalized at the synapse (data not shown). Nevertheless, the synapse formed between the B cell and the antigen-displaying target cell shares some obvious similarities to the synapse formed between T cells and their targets, such as a concentration of antigen receptor but exclusion of CD45 from the region of synapsis. But whereas T cells interacting with their targets, such as a concentration of antigen receptor but exclusion of CD45 from the region of synapsis, such a pinch-pinching of membrane vesicles from the target cell. Our results show that membrane tethering of antigen also leads to great enhancement of B-cell signalling, as judged by the upregulation of CD86 (Fig. 3c), it seems that the focusing of antigen into the synapse may be important in B-cell responses to antigen—especially at low antigen abundance. With respect to membrane-integral antigens on foreign target cells, the gathering of antigen into the synapse and its consequent acquisition will presumably enhance the extent to which cognate B cells will present peptides to T cells that are derived from proteins linked to the antigen recognized by the BCR (rather than from irrelevant proteins in the foreign target). This might favour efficient recruitment of T-cell help and bias against promiscuous T-cell activation and the development of autoimmunity.

**Methods**

**Loading and detection of immune complexes**

To load immune complexes onto 279.AC1 myeloid cells or L-cell transfectants expressing human FcγRI (a gift from M. Clark), we incubated cells as described with a mixture of biotinylated HEL, HEL-specific IgG1 HyHEL5 and F10 (which recognize HEL epitopes distinct from that recognized by the HyHEL10 BCR expressed by B cells from MD4 transgenic mice) and rabbit anti-mouse IgG. Loaded immune complexes were detected using fluorescein isothiocyanate (FITC)–streptavidin and PE-conjugated goat anti-mouse IgG. We detected SHP1 and phosphorylated tyrosine (P-Tyr) using rabbit polyclonal antisera (SantaCruz) and biotinylated monoclonal antibody 4G10 (Upstate Biotechnol-
egg), respectively. Staining specificity was controlled by single staining, as well as by using secondary antibodies in the absence of the primary stain.

**Generation of target cells**

Target cells displaying a membrane-integral version of either wild-type HEL or a mutant \(^{16}\) exhibiting reduced affinity for HEL10 (\([R^8], D^{47}, G^{48}, N^{50}\) designated HEL \(^*\)) were generated by transfecting mouse J558L plasmacytoma cells with constructs analogous to those used \(^{18}\) for expression of soluble HEL/HEL \(^*\), except that 14 Ser/Gly codons, the H2K \(\beta\)-transmembrane region, and a 23-codon cytoplasmic domain were inserted immediately upstream of the termination codon by polymerase chain reaction. For mHEL–GFP, we included the EGFP coding domain in the Ser/Gly linker. Abundance of surface HEL was monitored by flow cytometry and radiolabelled-antibody binding using HYHELs and DI.3 HEL-specific monomeric antibodies, for which the mutant HELs used in this work showed unaltered affinities \(^{18}\).

**Interaction assays**

For B-cell/target interaction assays, splenic B cells from 3–83 or MD4 transgenic mice \(^{28,29}\) and HEL-specific monoclonal antibodies, for which the mutant HELs used in this work exhibited reduced affinity for HyHEL10 (\([R^21, D^{101}, G^{102}, N^{103}\] designated HEL \(*\)) were used for expression of soluble HEL/HEL \(^*\) before being applied to polylysine-coated slides. Cells were fixed in 4% paraformaldehyde/PBS or methanol and permeabilized with PBS/0.1% Triton X-100 before immunofluorescence. We acquired confocal images using a Nikon E800 microscope attached to BioRad Radiance Plus scanning system equipped with 488-nm and 543-nm lasers, as well as differential interference contrast for transmitted light. GFP fluorescence in living cells in real time was visualized using a Radiance 2000 and Nikon E800 inverted microscope. Images were processed using BioRad Lasersharp 1024 or 2000 software to provide single plane images, confocal projections or slicing.

**Antigen presentation**

Presentation of HEL epitopes to T-cell hybridomas 2G7 (specific for I-\(\beta\)2[HEL18–20]) and IEE (specific for I-E\(\beta\)[HEL14–15]) by transfectants of the LK35.2 B-cell hybridoma expressing an HEL–specific IgM BCR was monitored as described \(^{19}\).

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**Duplexes of 21–nucleotide RNAs mediate RNA interference in cultured mammalian cells**

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RNA interference (RNAi) is the process of sequence-specific, post-transcriptional gene silencing in animals and plants, initiated by double-stranded RNA (dsRNA) that is homologous to sequence in the silenced gene \(^{4,5}\). The mediators of sequence-specific messenger RNA degradation are 21- and 22-nucleotide small interfering RNAs (siRNAs) generated by ribonuclease III cleavage from longer dsRNAs \(^{6,10–12}\). Here we show that 21-nucleotide siRNA duplexes specifically suppress expression of endogenous and heterologous genes in different mammalian cell lines, including human embryonic kidney (293) and HeLa cells. Therefore, 21-nucleotide siRNA duplexes provide a new tool for studying gene function in mammalian cells and may eventually be used as gene-specific therapeutics.

Uptake of dsRNA by insect cell lines has previously been shown to ‘knock-down’ the expression of specific proteins, owing to sequence-specific, dsRNA-mediated mRNA degradation \(^{10–12}\). However, it has not been possible to detect potent and specific RNA interference in commonly used mammalian cell culture systems, including 293 (human embryonic kidney), NIH/3T3 (mouse fibroblast), BHK-21 (Syrian baby hamster kidney), and CHO–K1 (Chinese hamster ovary) cells, applying dsRNA that varies in size between 38 and 1,662 base pairs (bp) \(^{16,17}\). This apparent lack of RNAi in mammalian cell culture was unexpected, because RNAi exists in mouse oocytes and early embryos \(^{18,19}\), and because RNAi-related, transgene-mediated co-suppression was also observed in cultured Rat-1 fibroblasts \(^{20}\). But it is known that dsRNA in the cytoplasm of mammalian cells can trigger profound physiological