Virus-Neutralizing Monoclonal Antibody Expressed in Milk of Transgenic Mice Provides Full Protection against Virus-Induced Encephalitis

ANDREAS F. KOLB,1,2* LECIA PEWE,3 JOHN WEBSTER,4 STANLEY PERLMAN,3 C. BRUCE A. WHITELEAF,4 AND STUART G. SIDDELL2

Cell Physiology Group, Hannah Research Institute, Ayr,1 and Department of Gene Expression and Development, Roslin Institute, Roslin,4 United Kingdom; Institute of Virology, University of Würzburg, Würzburg, Germany2; and Department of Pediatrics and Microbiology, University of Iowa, Iowa City, Iowa3

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Neutralizing antibodies represent a major host defense mechanism against viral infections. In mammals, passive immunity is provided by neutralizing antibodies passed to the offspring via the placenta or the milk as immunoglobulin G and secreted immunoglobulin A. With the long-term goal of producing virus-resistant livestock, we have generated mice carrying transgenes that encode the light and heavy chains of an antibody that is able to neutralize the neurotropic JHM strain of murine hepatitis virus (MHV-JHM). MHV-JHM causes acute encephalitis and acute and chronic demyelination in susceptible strains of mice and rats. Transgene expression was targeted to the lactating mammary gland by using the ovine b-lactoglobulin promoter. Milk from these transgenic mice contained up to 0.7 mg of recombinant antibody/ml. In vitro analysis of milk derived from different transgenic lines revealed a linear correlation between antibody expression and virus-neutralizing activity, indicating that the recombinant antibody is the major determinant of MHV-JHM neutralization in murine milk. Offspring of transgenic and control mice were challenged with a lethal dose of MHV-JHM. Litters suckling nontransgenic dams succumbed to fatal encephalitis, whereas litters suckling transgenic dams were fully protected against challenge, irrespective of whether they were transgenic. This demonstrates that a single neutralizing antibody expressed in the milk of transgenic mice is sufficient to completely protect suckling offspring against MHV-JHM-induced encephalitis.

Coronaviruses are a group of enveloped viruses with a single-stranded RNA genome of positive polarity (37). They are frequently associated with respiratory and gastrointestinal disorders in both animals and humans. Many coronavirus infections are mild in adult animals, whereas they often cause severe and sometimes lethal diseases in neonates (9, 32). To a large extent, this is due to the immature immune system of the newborn host. Maternal antibodies supplied via the placenta and milk efficiently protect newborn animals against the fatal consequences of acute coronavirus infections during this critical phase (14, 15). Cross-fostering experiments have shown that milk-borne antibodies (immunoglobulin A [IgA] and IgG) are sufficient to completely protect newborn mice against lethal doses of murine hepatitis virus (MHV) (15).

Vaccination against coronavirus infections has been employed with various degrees of success (23, 25, 36). The vaccines are usually highly strain specific (16), but they are also dependent on specific routes of infection and often short-lived. Live-virus vaccines are also associated with the danger of in vivo recombination, leading to novel viruses with increased pathogenicity.

Neutralizing monoclonal antibodies generated in response to coronavirus infections have been isolated in many laboratories (12, 35, 42), and it has been shown that antibodies which inhibit virus entry into susceptible cells in vitro can also effectively prevent acute coronavirus-induced disease in vivo (26, 42). Coronavirus infections cause a high mortality only during a short time period (up to 20 days postpartum in mice), which largely coincides with the suckling period. We and others (3, 39) have therefore reasoned that the recombinant expression of neutralizing antibodies in the milk of transgenic animals may provide an effective strategy to protect animals during this critical phase. To provide a proof of principle, we have generated transgenic mice expressing a highly neutralizing monoclonal antibody directed against the neurotropic MHV strain JHM (MHV-JHM). The recombinant antibody was secreted into the milk at yields of up to 0.7 mg/ml. The biological activity of the milk-borne antibody was demonstrated by virus neutralization assays in vitro, and a linear correlation between antibody expression and neutralization was found. When litters suckling transgenic dams were infected with a lethal dose of MHV-JHM, they were completely protected against virus-induced disease, irrespective of whether the newborn mice were transgenic. These results provide the first example of transgene-mediated lactogenic immunity in vivo.

MATERIALS AND METHODS

DNA cloning. Monoclonal antibody (MAb) A1 was selected for these studies because it is highly potent with regard to virus neutralization and inhibition of virus-induced cell-to-cell fusion (42). The isolation and cloning of cDNAs encoding the variable regions of MAb A1 have been described previously (21). In brief, mRNA was isolated from the A1 hybridoma cell line and reverse transcribed. The resulting wx and vy cDNAs were amplified by PCR, using primers which bind in the framework of the variable regions (21). The variable region-
digested and religated with EcoRV linker

β-LG gene

HindIII

EcoRV

XbaI

PvuII

PvuII

pBJ41

HindIII

EcoRV/NcoI

EagI/EcoRV

pBJ41-A1H

HindIII

EcoRV/NcoI

EcoRI/EcoRV

pBJ41-A1L

FIG. 1. Schematic representation of the ovine β-LG gene, the expression vector pBJ41, and the transgene constructs pBJ41-A1H and pBJ41-A1L. Exons of the β-LG gene and the human IgG genes are represented by shaded boxes and filled boxes, respectively. Exons encoding the heavy chain constant region are marked as 1, 2, 3, and 5; the exon encoding the light chain constant region is marked as Cx. The inserted variable regions (V_y and V_s) isolated from the A1 mouse hybridoma cell line are represented as hatched boxes. The mammary gland-specific expression vector pBJ41 contains sequences from the first and fifth exons (17 and 8 bp, respectively) and the entire sixth and seventh exons of the β-LG gene. A linker carrying a singular EcoRV site was inserted in between the two PvuII sites in exons 1 and 5, a, b, and c, β-LG translational initiation site, stop codon, and polyadenylation signal, respectively; d and e, IgG translational start and stop codons, respectively. The A1 heavy chain coding region was inserted as a blunt-ended NcoI-Eagl fragment into the singular EcoRV site of pBJ41. The A1 light chain coding region was inserted as a blunt-ended NcoI-EcoRI fragment into the same site.

Animals. Transgenic mice (F1, CBA × C57BL/6) were produced by pronuclear injection at the animal facility of the Roslin Institute as described previously (4, 43). The plasmids pBJ41-A1L and pBJ41-A1H and the genomic β-LG vector pBS11gXS (1) were all linearized by digestion with SfiI-XbaI and microinjected at a 1:1:3 ratio. pSS1lgXS carries the entire β-LG coding region, with 4.3 kb of 5′ flanking region and 1.9 kb of 3′ flanking region. An excess of genomic β-LG vector over the expression vector biases for increased frequency of transgene expression (4). Transgenic mice were identified by PCR and Southern blot analysis of genomic DNA. The pBJ41-A1H transgene was detected by PCR, using the primers pBJup (5′-AGC CGT CCT GTC TCA GCC CT 3′) and A1Hsp (5′-TGC ATG TGA TGG ACA GGC-3′). The primer pair gives rise to a 264 bp product. The pBJ41-A1L transgene was detected by Southern blot analysis of BamHI-digested genomic DNA. The pBJ41-A1H transgene was detected by PCR, using the primers pBJup (5′-AGC CGT CCT GTC TCA GCC CT 3′) and A1Hsp (5′-TGC ATG TGA TGG ACA GGC-3′). The primer pair gives rise to a 264 bp product. The pBJ41-A1L transgene was detected by Southern blot analysis of BamHI-digested genomic DNA. The blot was hybridized with a pBJ41-A1L-specific 281-bp PCR product amplified with the primer pair pBJup and A1Lsp (5′-CTA CTA AGG TTT TTG CAT TA-3′) using plasmid DNA as template. Transgene copy number was determined by Southern blot analysis on liver DNA from G1 mice. Aliquots (20 μg) of DNA were digested with BamHI, separated on a 1% agarose gel, and blotted to a nylon membrane. The blot was subsequently probed with a 1.6-kb BamHI-SphI fragment of the β-LG promoter which is present in all three constructs. Eleven transgenic founder mice were generated, eight of which transmitted the transgenes. These eight lines were used for further analysis.

The virus challenge experiments were carried out with pathogen-free C57BL/6 (B6) mice obtained from the National Cancer Institute (Bethesda, Md.). These mice were seronegative for MHV-JHM. Transgenic females were crossed back with B6 males for five generations, and offspring were screened for the presence of the transgenes as described above. For cross-fostering experiments, suckling mice born to B6 females mated with B6 males and transgenic females mated with B6 males were switched within 24 h of delivery.

Protein analysis. Western blot analyses were done essentially as described previously (20). The milk of transgenic mice (isolated at peak lactation, day 10 postpartum) was diluted 1:5 with water and centrifuged at 14,000 × g for 5 min. Three phases were separated: a layer of fat above an aqueous phase and a pellet. The aqueous phase containing the whey fraction was isolated, and aliquots were mixed with reducing and denaturing sample buffer and separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis. The electrophoretically separated proteins were transferred to a nitrocellulose membrane by semidry electrophotlotting. The constant regions of the chimeric A1 antibody were detected by using a rabbit anti-human IgG-horseradish peroxidase (HRP)-linked antiserum (Dianova). The results were quantified by densitometric scanning on a Molecular Dynamics Densitometer.

Cells and viruses. DBT (delayed brain tumor) cells (24) were cultivated at 37°C in minimum essential medium (Life Technologies) supplemented with 10% fetal bovine serum (Sigma), nonessential amino acids, glutamine, and antibiotics. The MHV-JHM strain used for the in vitro experiments was described previously (35). Virus neutralization was quantified in synctium focus reduction assays. MHV-JHM infections lead to extensive cell-to-cell fusion without inducing cytotoxic effects in the first 24 h postinfection. However, synctium formation (as is the case with plaque formation) indicates the presence of an infection center and is therefore equivalent to plaque formation in its diagnostic value. In keeping with the literature, the number of infectious centers is referred to as PFU. Defatted and diluted milk samples were further diluted in phosphate-buffered saline. Aliquots (500 μl) of a 10−4 dilution of MHV-JHM, equivalent to about 70 PFU of virus, were mixed with 500 μl of different dilutions of milk samples isolated from transgenic lines or nontransgenic control mice and incubated for 1 h at 37°C. Subsequently, the virus-antibody mixture was added to confluent DBT cells for 1 h at 37°C. The virus-containing supernatant was then removed. The cells were washed twice with phosphate-buffered saline, and fresh medium was added. Foci of synctium formation were counted after the infected cells were incubated for 16 h at 37°C. The virus strain used for the in vivo studies was grown as described previously (31). To determine the protective efficacy of breast milk-expressed antibody, 10-day-old mice were challenged by intranasal or intracerebral inoculation, as described previously (31). Intranasal challenge was done by inoculation of 2 × 104 to 4 × 105 PFU of MHV-JHM. Intracerebral challenge was done by inoculation of 7 × 105 or 7 × 106 PFU of MHV-JHM.
Mice were monitored daily for mortality and morbidity.

In all, eight lines of transgenic mice were generated. The expression vectors pBJ41-A1L and pBJ41-A1H were microinjected into fertilized oocytes at equimolar levels. However, the transgene copy numbers seem to indicate that consistently more copies of the light than the heavy chain construct were incorporated into the host genome. The reason for this is unknown, but it may be due to a leaky expression of the transgenes during early developmental stages which, in turn, leads to a cytotoxic overproduction of heavy chain protein and the subsequent loss of embryos.

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RESULTS

MHV-JHM infections can be effectively prevented in vitro and in vivo by neutralizing antibodies, CD4 T cells, and CD8 T cells (10, 40, 42). The majority of neutralizing antibodies are directed against the viral surface (S) glycoprotein. MAAb A1 (42), which binds to the S1 subunit of the MHV-JHM S protein, is one of the most potent antibodies with regard to virus neutralization and the inhibition of virus-induced cell-to-cell fusion. We have isolated the variable regions of MAAb A1 and transferred them into different eukaryotic expression vectors. The recombinantly expressed version of MAAb A1 displayed the same biological activity as the parental MAAb secreted from the A1 hybridoma cell line (21, 22).

In the studies described here, the MAAb A1 variable regions were linked to human constant regions of the IgG isotype to facilitate their identification against the background of murine antibodies. The resulting chimeric open reading frames were inserted into a mammary gland-specific expression cassette based on the ovine β-LG gene (Fig. 1). After confirming expression of the antibody genes in cell culture (data not shown), the two expression vectors (pBJ41-A1L and pBJ41-A1H) were used for the generation of transgenic animals.

In all, eight lines of transgenic animals were analyzed for the expression of recombinant antibody by Western blotting. To do this, the human constant regions of the MHV-neutralizing antibody were detected by using an HRP-linked rabbit anti-human IgG antiserum. Of the eight transgenic lines transmitting all three transgenes, five expressed the heavy and the light chains of the recombinant antibody. In two transgenic lines, only light chain protein but no heavy chain protein was detectable (Table 1). The seven lines which expressed the light chain gene expressed the genomic β-LG construct as well. The light chain was always expressed in excess of the heavy chain (Table 1 and Fig. 2) and is also readily detected in a Coomassie blue-stained protein gel (data not shown). The transgene copy number was estimated in seven of the eight lines obtained (Table 1) and varied from 1 to 10 copies (heavy chain) and 4 to 15 copies (light chain). The transgenes encoding the antibody light chain and the heavy chain were microinjected into fertilized oocytes at equimolar levels. However, the transgene copy numbers seem to indicate that consistently more copies of the light than the heavy chain construct were incorporated into the host genome. The reason for this is unknown, but it may be due to a leaky expression of the transgenes during early developmental stages which, in turn, leads to a cytotoxic overproduction of heavy chain protein and the subsequent loss of embryos.

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Table 1. Transgene copy number and expression

<table>
<thead>
<tr>
<th>Transgenic line</th>
<th>pSS1tgXS</th>
<th>pBJ41-A1H</th>
<th>pBJ41-A1L</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No. of copies</td>
<td>Level of β-LG expression</td>
<td>No. of copies</td>
</tr>
<tr>
<td>HEP3</td>
<td>16</td>
<td>++ + + + +</td>
<td>2</td>
</tr>
<tr>
<td>HEP10</td>
<td>20</td>
<td>++ + + + +</td>
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<td>ND†</td>
<td>ND</td>
<td>ND</td>
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<tr>
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<td>12</td>
<td>++ + + + +</td>
<td>1</td>
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<td>HEP30</td>
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<td>2</td>
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</tr>
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<td>10</td>
</tr>
<tr>
<td>HEP50</td>
<td>2</td>
<td>++</td>
<td>2</td>
</tr>
</tbody>
</table>

† Transgene copy number was determined as described in Materials and Methods. Relative levels of transgene expression were determined by Western blotting and Coomassie brilliant blue protein staining after polyacrylamide gel electrophoresis.

‡ +, transgene expression could be detected by Western blotting and protein staining. The number of plus signs indicates the relative level of expression. (±), transgene expression could not be detected by Western blotting.

§ The concentration of heavy and light chain protein as quantified by densitometric scanning of Western blots (as described in Materials and Methods).

FIG. 2. Western blot analysis of milk samples. Aliquots of defatted milk samples (corresponding to 1 μl of milk) isolated from mice of six transgenic lines were separated by 15% polyacrylamide gel electrophoresis alongside a human IgG standard (1,000, 300, and 100 ng of IgG) and blotted onto nitrocellulose. The human IgG portion of the chimeric recombinant antibody was detected by using an HRP-linked rabbit-antihuman IgG antiserum. The blot was developed using a chemiluminescent detection system (Pierce). The positions of the antibody chains and the sizes of the molecular marker proteins are indicated.
No correlation between the transgene copy numbers and the levels of antibody expression could be detected (Table 1). We interpret this to indicate that the site of transgene integration had a dominant effect on gene expression. Although expression of the light chain protein was consistently higher than expression of the heavy chain protein, this did not directly correlate with transgene copy number. Additionally, no correlation could be detected between the levels of β-LG protein expression and antibody expression (Table 1).

The concentrations of the heavy and light chain proteins were quantified by densitometry in comparison to IgG standards (Bergiglobin; pooled human immunoglobulin; Behringwerke). Maximum levels of 0.5 and 2.9 mg/ml, respectively, of heavy and light chain proteins were obtained in transgenic line HEP50. The heavy chain, which comprises 69% of the total molecular weight of an IgG molecule, can only be secreted from cells as part of a complete antibody. In contrast, the light chain protein can be secreted individually. Therefore the 0.5 mg of heavy chain protein per ml measured in the milk of transgenic line HEP50 corresponds to a total recombinant IgG concentration of 0.7 mg/ml. The heavy chain, which comprises 69% of the total molecular weight of an IgG molecule, can only be secreted from cells as part of a complete antibody.

The defatted milk of transgenic line HEP50 was subsequently analyzed in a virus neutralization assay. A 10^{-4} dilution of defatted milk reduced MHV-JHM infectivity by 90% (Fig. 3A), which corresponds to 125 pg of IgG being required to neutralize 1 PFU of MHV-JHM. This is consistent with results we have obtained in cell culture, where 100 pg of IgG was required to neutralize 1 PFU of MHV-JHM (21). Milk isolated from nontransgenic mice did not show any neutralizing effect against the MHV-JHM infection (Fig. 3A). This confirms the fact that the breeding colony of mice was seronegative for MHV-JHM and also that the recombinant antibody is the decisive virus-neutralizing factor in milk. Defatted milk samples from four other transgenic lines were also analyzed by neutralization assays. All of the samples neutralized MHV infectivity, albeit with different levels of efficacy (Fig. 3B). The dilutions at which MHV-JHM infectivity was reduced by 90 and 50% were correlated with expression of the heavy chain (which is the limiting factor for antibody formation) (Fig. 3C). The linear correlation observed confirms that the concentration of the recombinant antibody in the milk of these transgenic animals is the critical factor determining neutralization activity.

Finally, the ability of recombinant antibody secreted in breast milk to protect suckling mice in vivo was determined. Transgenic mice and their nontransgenic littermates were infected intranasally with virulent MHV-JHM. Under these conditions (31), 100% of naive mice succumb to acute, fatal encephalitis by 5 to 7 days postinoculation. However, as shown in Table 2, 23 of 23 suckling mice nursed by transgenic dams did not develop acute encephalitis, whereas 6 of 6 suckling mice nursed by transgene-negative animals died by 7 days postinfection (Table 2). Infected mice nursed by transgenic dams also grew at the same rate as uninfected mice nursed by transgenic
We have analyzed whether expressing a neutralizing antibody as a recombinant protein in the milk of transgenic animals may provide an alternative to these procedures. This transgenic approach is particularly useful if (i) no useful vaccines are available, (ii) neutralizing antibodies are a major component of the host’s defense against the pathogen in question, and (iii) the period during which the infection is life-threatening is limited. The use of transgenic animals offers the additional advantage that the most potent antibodies can be selected by in vitro analyses before transgenes are established from the respective hybridoma cell lines. Moreover, no live pathogens have to be introduced into an animal colony. Coronavirus, which cause diseases of economic importance in animals and humans, fulfill all of the above-mentioned criteria. Therefore, transgenic animals that express neutralizing antibodies in the milk may, in the long term, provide a strategy to protect animals during the suckling period. The experiments described here demonstrate that this approach is practicable. However, antibody isotypes mediating lactogenic immunity vary between different species (27). There are also profound differences in the intestinal uptake of immunoglobulins (27). Transgene-mediated strategies manipulating lactogenic immunity in livestock will therefore have to be adapted to the requirements of individual species in terms of immunoglobulin isotype and expression strategy. In that respect, the transgenic mouse-MHV model provides an excellent opportunity to determine the critical factors for successful immune protection through in vivo challenge experiments.

Two features distinguish transgene-mediated lactogenic immunity from the natural process of passive immunization by

![Image](image_url)

**FIG. 4.** Weight gain in infected and control animals. MHV-JHM-infected mice (n = 23, 12 females and 11 males) nursed by HEP50-positive transgenic dams and noninfected control mice (n = 20, 11 females and 9 males) were weighed at 44 days of age (34 days postinfection). Mean values and standard deviations are shown.
maternal antibodies. In the natural situation, newborn animals are provided with a polyclonal mixture of antibodies against a particular pathogen. The transgenic dams described here only provide a single, albeit highly neutralizing, antibody to their offspring. As has been shown previously, the β-LG-based expression system is able to supply immunoglobulin at a high concentration throughout the entire lactation period (39). In mammals, the total immunoglobulin concentration in colostrum ranges between 5 mg/ml (rat) and 250 mg/ml (cow). Lower immunoglobulin levels, ranging from 1 mg/ml (human) up to 10 mg/ml (pig), are found in milk (27, 41). Therefore, the concentration of recombinantly expressed neutralizing antibody of 0.7 mg/ml exceeds the concentration of any single monospecific antibody present in milk by some orders of magnitude. Nevertheless, there appeared to be no selection of virus escape mutants, which could overcome the immune protection and lead to overt disease within the time frame of the experiment. This indicates that production of a single MHV-JHM neutralizing antibody in milk is sufficient to provide full protection during the suckling period.

Milk proteins and their derivatives have been shown to have antibacterial and antiviral effects (29, 34, 44). In our experimental model, the neutralizing effect of the mouse milk samples was dependent on the concentration of the recombinant antibody. This suggests that no other natural components of milk provide a protective effect against MHV-JHM infections.

The levels of antibody production and secretion into milk are critical factors for the establishment of transgene-mediated virus resistance. The concentration of 0.7 mg of mature IgG per ml is at the lower end of expression levels that have been previously reported for other recombinant antibodies expressed in the milk of transgenic mice (0.4 mg/ml [28], 0.8 mg/ml [30], 4 mg/ml [13], 5 mg/ml [3], and 6 mg/ml [39]). Nevertheless, complete protection of litters suckling transgenic dams could be obtained at these expression levels. The comparatively low expression level is at least in part due to the failure to produce equimolar amounts of heavy and light chain protein (compare Table 1 and Fig. 2). This confirms our previous experiences with cell culture (21) and data published by others (33, 44). The consistent excess of light chain protein over heavy chain protein is due to the cytotoxicity of unpaired heavy chains (19), which leads to the death of cells in which the heavy chain is overexpressed. Additionally, we found that the MAb A1 heavy chain transgene was present at a lower copy number than the MAb A1 light chain construct in all strains of mice analyzed. This is consistent with the findings of Sola and colleagues (39), who used a β-LG-based expression system to express a recombinant antibody that neutralizes the porcine coronavirus transmissible gastroenteritis virus in the milk of transgenic mice. In contrast, when an expression system based on the whey acidic protein (WAP) promoter was used to express IgG-encoding genes, most of the transgenic lines generated carried an excess of heavy chain expression constructs (3). One possible explanation is that the β-LG promoter is active at an early stage of development and causes the loss of embryos in which the (cytotoxic) heavy chain is overexpressed. Transgenic animals in which the two antibody chains are expressed in equimolar amounts can be generated by microinjection (33) but may require the screening of a larger number of transgenic lines.

In conclusion, we have demonstrated that high levels of a virus-neutralizing antibody can be generated in the milk of transgenic mice and that full protection against virus-induced disease can be accomplished in newborn animals via this route. Thus, in the murine system, we have established a proof of principle, and the application of this technology to generate virus-resistant animals can now be pursued. Additionally, it now seems feasible that ruminants could be modified by transgenic methodology to produce milk containing neutralizing antibodies directed against pathogens responsible for major human infant diseases.

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