

Bricks

Semantic Content Enrichment

Sam Herbert

Overview

Science article abstract:

1. What semantic information can we add?
2. What are the benefits to the content consumer?

Ebola Virus Can Be Effectively Neutralized by Antibody Produced in Natural Human Infection

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant human monoclonal antibodies to Ebola virus antigens was isolated from phage display libraries constructed from RNA from donors who recovered from infection in the 1997 Ebola virus outbreak in Kikwit, Democratic Republic of Congo. Antibodies reactive with nucleoprotein (NP), envelope glycoprotein (GP), and secreted envelope glycoprotein (sGP) were characterized by immunofluorescence and radioimmunoprecipitation assays. Four antibodies reacting strongly with sGP and weakly with GP and two antibodies reacting with NP were not neutralizing. An antibody specific for GP neutralized Ebola virus to 50% at 0.4 µg/ml as the recombinant Fab fragment and to 50% at 0.3 µg/ml (90% at 2.0 µg/ml) as the corresponding whole immunoglobulin G1 molecule. The studies indicate that neutralizing antibodies are produced in infection by Ebola virus although probably at a relatively low frequency. The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Toshiaki Maruyama,¹ Luis L. Rodriguez,^{2,†} Peter B. Jahrling,³ Anthony Sanchez,² Ali S. Khan,² Stuart T. Nichol,² C. J. Peters,² Paul W. H. I. Parren,¹ and Dennis R. Burton^{1,*} Ebola Virus Can Be Effectively Neutralized by Antibody Produced in Natural Human Infection.

Keywords

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant human monoclonal antibodies to Ebola virus antigens was isolated from phage display libraries constructed from RNA from donors who recovered from infection in the 1995 Ebola virus outbreak in Kikwit, Democratic Republic of Congo. Antibodies reactive with nucleoprotein (NP), envelope glycoprotein (GP), and secreted envelope glycoprotein (sGP) were characterized by immunofluorescence and radioimmunoprecipitation assays. Four antibodies reacting strongly with sGP and weakly with GP and two antibodies reacting with NP were not neutralizing. An antibody specific for GP neutralized Ebola virus to 50% at 0.4 $\mu\text{g/ml}$ as the recombinant Fab fragment and to 50% at 0.3 $\mu\text{g/ml}$ (90% at 2.6 $\mu\text{g/ml}$) as the corresponding whole immunoglobulin G1 molecule. The studies indicate that neutralizing antibodies are produced in infection by Ebola virus although probably at a relatively low frequency. The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Keywords

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant monoclonal antibodies to Ebola virus antigens was isolated from mice constructed from RNA from donors who recovered from infection during an Ebola virus outbreak in Kikwit, Democratic Republic of Congo. The antibodies were characterized for reactivity with nucleoprotein (NP), envelope glycoprotein (GP), and secreted phospholipase 2 (SLP2) were characterized by ELISA and radioimmunoassay. Four antibodies reacted with GP and two reacted weakly with NP and two antibodies reacting with NP were specific for GP neutralized Ebola virus to 50% at 0.4 $\mu\text{g/ml}$ Fab fragment and to 50% at 0.3 $\mu\text{g/ml}$ (90% at 2.6 $\mu\text{g/ml}$) including whole immunoglobulin G1 molecule. The studies indicate that neutralizing antibodies are produced in infection by Ebola virus although probably at a relatively low frequency. The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Content
Insights

Related
Articles

Entities (genes and proteins)

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant human monoclonal antibodies to Ebola virus antigens was isolated from phage display libraries constructed from **RNA** from donors who recovered from infection in the 1995 Ebola virus outbreak in Kikwit, Democratic Republic of Congo. Antibodies reactive with **nucleoprotein (NP)**, **envelope glycoprotein (GP)**, and secreted **envelope glycoprotein (sGP)** were characterized by immunofluorescence and radioimmunoprecipitation assays. Four antibodies reactive with sGP and

weakly w **Accession number (ID): Q05320** body
specific f **Alternative names: GP1, GP2, GP2-delta**
fragment **Virus host: Homo sapiens, Franquet's epauleted fruit**
immunog **bat, Little collared fruit bat**
produced in infection by Ebola virus although probably at a relatively low frequency.
The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Entities (medical)

The activity of antibodies against **filoviruses** is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant **human** monoclonal antibodies to **Ebola virus** antigens was isolated from **phage** display libraries constructed from **RNA** from donors who recovered from **infection** in the 1995 **Ebola virus** outbreak in Kikwit, Democratic Republic of Congo. Antibodies reactive with nucleoprotein (NP), envelope glycoprotein (GP), and secreted envelope glycoprotein (sGP) were characterized by immunofluorescence and

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MeSHID=D029043

Tree=Viruses/Vertebrate Viruses/RNA

Viruses/Mononegavirales/Filoviridae

TreeNumber=B04.820.455.300.200

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produced in **infection** by **Ebola virus** although probably at a relatively low frequency. The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against **Ebola virus infection**.

Entities (medical)

The activity of a ... most **filoviruses** is poorly understood. ... and passive prophylaxis ... monoclonal antibodies ...
consequences ... and passive prophylaxis ... monoclonal antibodies ...
panels ... monoclonal antibodies ...
isolated ...
from ...
Concomitant ... active with nucleoprotein (NP) ...
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Entity Pages

Taxonomic Browsing

Faceted Search

MeSHID=D029043
Tree=Viruses/Vertebrate Viruses/RNA
Viruses/Mononegavirales/Filoviridae
TreeNumber=B04.820.455.300.200

produced in **infection** by **Ebola virus** although probably at ...
The neutralizing antibody may be useful in vaccine design ...
against **Ebola virus infection**.

Entities (places)

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant human monoclonal antibodies to Ebola virus antigens was isolated from phage display libraries constructed from RNA from donors who recovered from infection in the 1995 Ebola virus outbreak in **Kikwit**, **Democratic Republic of Congo**. Antibodies reactive with nucleoprotein (NP), envelope glycoprotein (GP), and secreted envelope glycoprotein (sGP) were characterized by immunofluorescence and radioimmunoprecipitation assays. Four antibodies strongly with sGP and

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Geoid=2314705 (Identifier in Geonames DB)
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The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Entities (places)

Map Visualisation

The activity of antibodies against Ebola virus antigen has important consequences for vaccine development. To investigate this activity, a panel of recombinant human monoclonal antibodies against Ebola virus antigens was isolated from phage display libraries derived from donors who recovered from infection in the Itombwe region of the Democratic Republic of Congo. Antibodies recognizing the Ebola virus glycoprotein (GP) secreted envelope glycoprotein (sGP) were characterized by immunofluorescence and radioimmunoprecipitation assays. Four antibodies strongly with sGP and we an antibody spc nt Fab fra whole im es are pro quency. The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Population=186991

Geoid=2314705 (Identifier in Geonames DB)

Admindiv1=Bandundu

Longitude=18.818

Latitude=-5.039

Relationships

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant human monoclonal antibodies to Ebola virus antigens was

isolated from individuals in Congo. secreted

Entity A: Envelope glycoprotein

Entity B: Ebola virus

Reaction type: Inhibition:Neutralise

radioimmunoprecipitation. Four antibodies reacting strongly with sGP and weakly with GP and two antibodies reacting with NP were not neutralizing. An antibody specific for GP neutralized Ebola virus to 50% at 0.4 µg/ml as the recombinant Fab fragment and to 50% at 0.3 µg/ml (90% at 2.6 µg/ml) as the corresponding whole immunoglobulin G1 molecule. The studies indicate that neutralizing antibodies are produced in infection by Ebola virus although probably at a relatively low frequency. The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Relationships

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To address this, a panel of recombinant human monoclonal antibodies to the GP of Ebola virus was isolated from immunized mice. Congo. secretes radioimmunoprecipitation assays showed four antibodies reacting strongly with sGP and weakly with GP and two antibodies reacting with NP were not neutralizing. An antibody specific for GP neutralized Ebola virus to 50% at 0.4 µg/ml as the recombinant GP at 0.3 µg/ml (90% at 2.6 µg/ml) as the control. The studies indicate that neutralizing antibody may be useful in vaccine design and as a treatment for Ebola virus infection.

Entity A: Envelope glycoprotein

Entity B: Ebola virus

Reaction type: Inhibition:Neutralise

Workflow Integration

Advanced Search

Classification - what is this content about?

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant human monoclonal antibodies to Ebola virus antigens was isolated from phage display libraries constructed from RNA from donors who recovered from infection in the 1995 Ebola virus outbreak in Kikwit, Democratic Republic of Congo. Antibodies reactive with nucleoprotein (NP), envelope glycoprotein (GP), and secreted envelope glycoprotein (sGP) were characterized by immunofluorescence and radioimmunoprecipitation assays. Four antibodies reacting strongly with sGP and weakly with GP and two antibodies reacting with NP were not neutralizing. An antibody specific for GP neutralized Ebola virus to 50% at 0.4 $\mu\text{g/ml}$ as the recombinant Fab fragment and to 50% at 0.3 $\mu\text{g/ml}$ (90% at 2.6 $\mu\text{g/ml}$) as the corresponding whole immunoglobulin G1 molecule. The studies indicate that neutralizing antibodies are produced in infection by Ebola virus although probably at a relatively low frequency. The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Virology

Classification - what is this content about?

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant human monoclonal antibodies to Ebola virus antigens was isolated from phage display libraries constructed from RNA from donors who recovered from infection in the 1995 Ebola virus outbreak in Kikwit, Democratic Republic of Congo. Antibodies reactive with nucleoprotein (NP), envelope glycoprotein (GP) and secreted envelope glycoprotein (sGP) were characterized by immunoblotting and radioimmunoprecipitation assays. Four antibodies reacting strongly with GP reacted weakly with NP and two antibodies reacting with NP were not reactive with GP. A GP-specific antibody neutralized virus to 50% at 0.4 µg/ml as the complement of NP and GP fragments (50% at 2.6 µg/ml) as the complement of GP and NP fragments. These results indicate that neutralizing activity is probably at a relative concentration of 1:1. The results have implications for vaccine design and as a prophylactic agent.

Virology

Content Slices

Virtual Issues

Article semantic fingerprint

The activity of antibodies against filoviruses is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant human monoclonal antibodies to Ebola virus antigens was isolated from phage display libraries constructed from RNA from donors who recovered from infection in the 1995 Ebola virus outbreak in Kikwit, Democratic Republic of Congo. Antibodies reactive with nucleoprotein (NP), envelope glycoprotein (GP), and secreted envelope glycoprotein (sGP) were characterized by immunofluorescence and radioimmunoprecipitation assays. Four antibodies reacting strongly with sGP and weakly with GP and two antibodies reacting with NP were not neutralizing. An antibody specific for GP neutralized Ebola virus to 50% at 0.4 $\mu\text{g/ml}$ as the recombinant Fab fragment and to 50% at 0.3 $\mu\text{g/ml}$ (90% at 2.6 $\mu\text{g/ml}$) as the corresponding whole immunoglobulin G1 molecule. The studies indicate that neutralizing antibodies are produced in infection by Ebola virus although probably at a relatively low frequency. The neutralizing antibody may be useful in vaccine design and as a prophylactic agent against Ebola virus infection.

Virology

Article semantic fingerprint

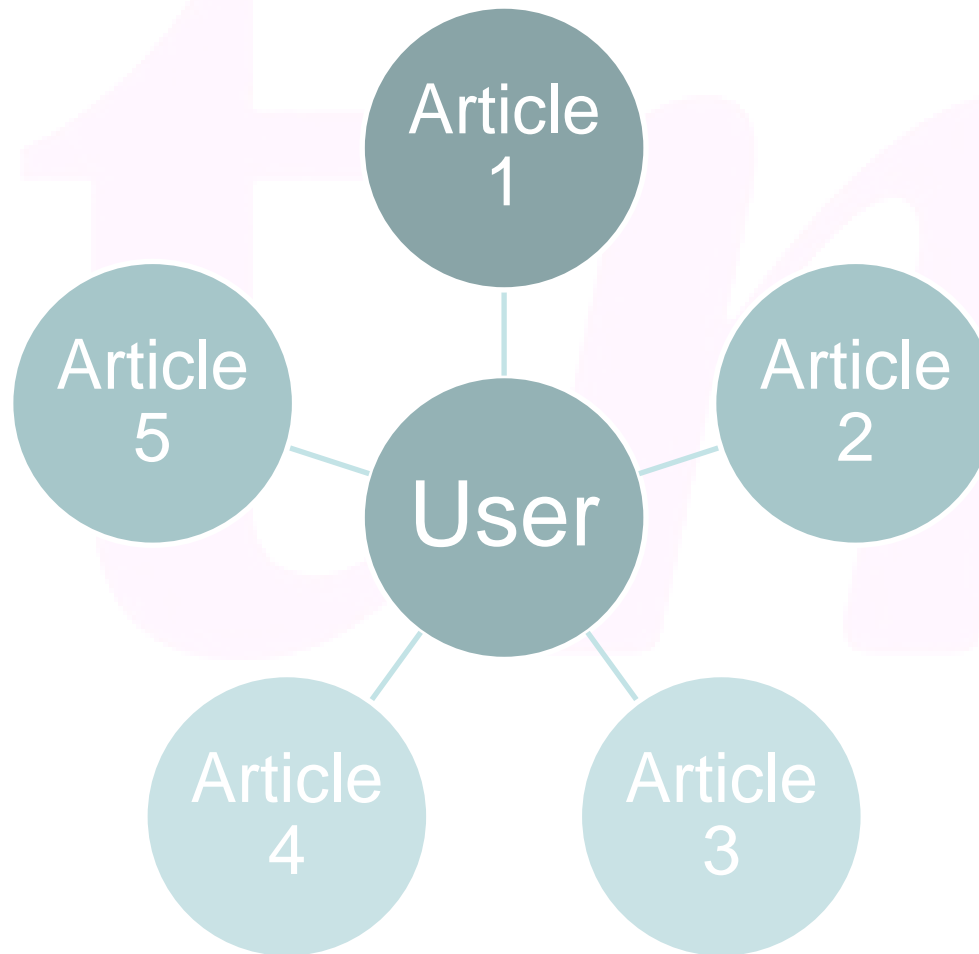
The activity of antibodies against **filoviruses** is poorly understood but has important consequences for vaccine design and passive prophylaxis. To investigate this activity, a panel of recombinant monoclonal antibodies to Ebola virus antigen was isolated from mice. These antibodies were constructed from **RNA** isolated from mice during the **Ebola virus** outbreak in **Kikwit, Congo**. Antigenic sites on the glycoprotein (GP), envelope glycoprotein (GP2), and secreted glycoprotein (GP1) were characterized by radioimmunoassay. Four antibodies reacting strongly with GP1 reacted weakly with GP2 and two antibodies reacting with GP2 were not neutralizing. An antibody specific for GP2 neutralized **Ebola virus** to 50% at 0.4 $\mu\text{g/ml}$ as the recombinant Fab fragment and to 50% at 0.3 $\mu\text{g/ml}$ (90% at 2.6 $\mu\text{g/ml}$) as the corresponding whole immunoglobulin G1 molecule. The studies indicate that **neutralizing antibodies** are produced in **infection** by **Ebola virus** although probably at a relatively low frequency. The **neutralizing antibody** may be useful in **vaccine design** and as a prophylactic agent against **Ebola virus infection**.

Smart
Content
Analysis

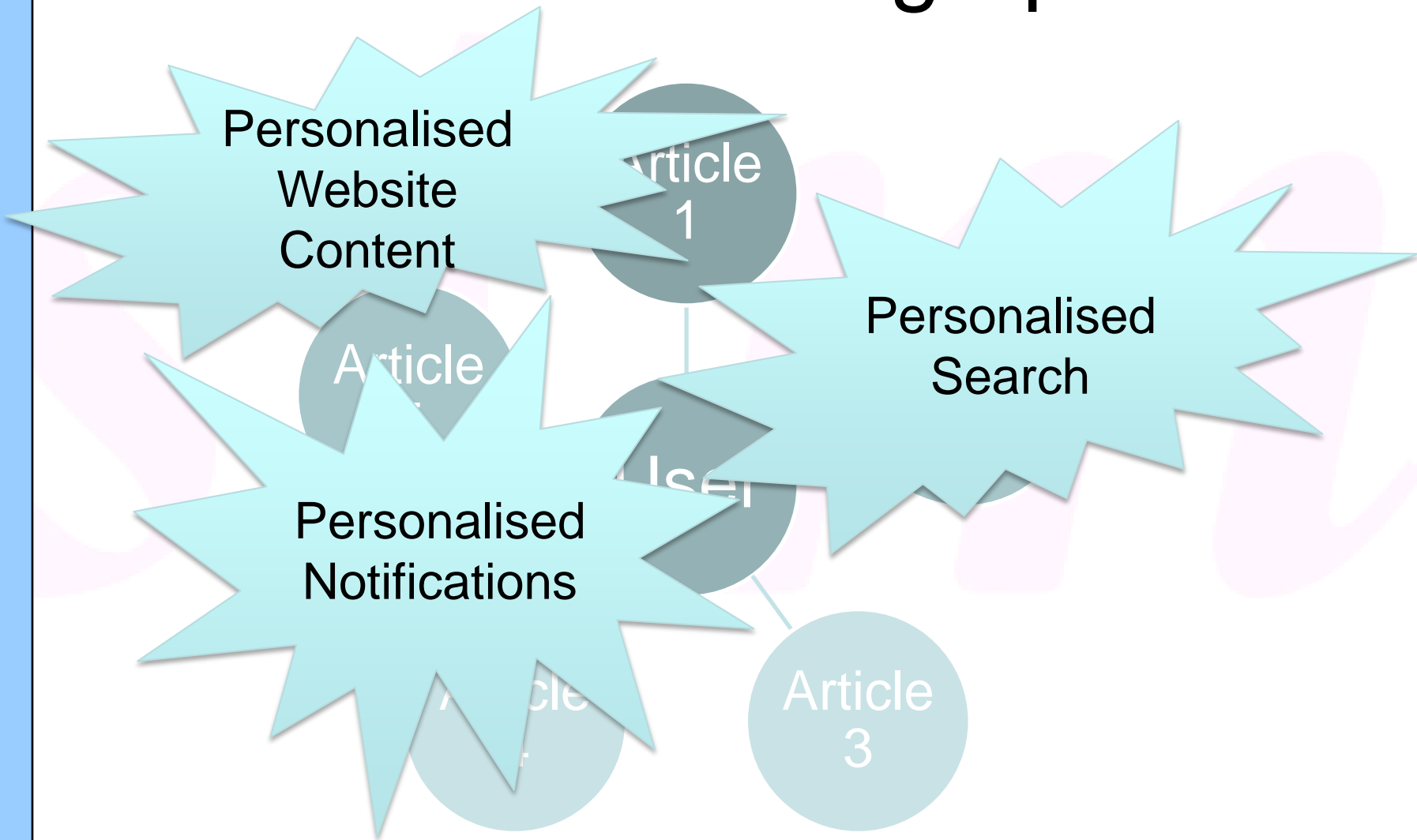
Smart
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Prology

User semantic fingerprint




User semantic fingerprint



Semantic Content Enrichment

Core competency for
information providers and publishers

Time's Up!



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