

Mathematical Analysis of Antibody by Gamma and PSI Functions

Bin Zhao*, Aibing Li and Lichun Liang

College of Science, Northwest A & F University, China

*Corresponding author: Bin Zhao, College of Science, Northwest A&F University, Yangling, Shaanxi, China, Tel: +86 13028517572; Email: zhaobin835@nwsuaf.edu.cn

Submission: 📅 September 13, 2017; Published: 📅 December 18, 2017

Abstract

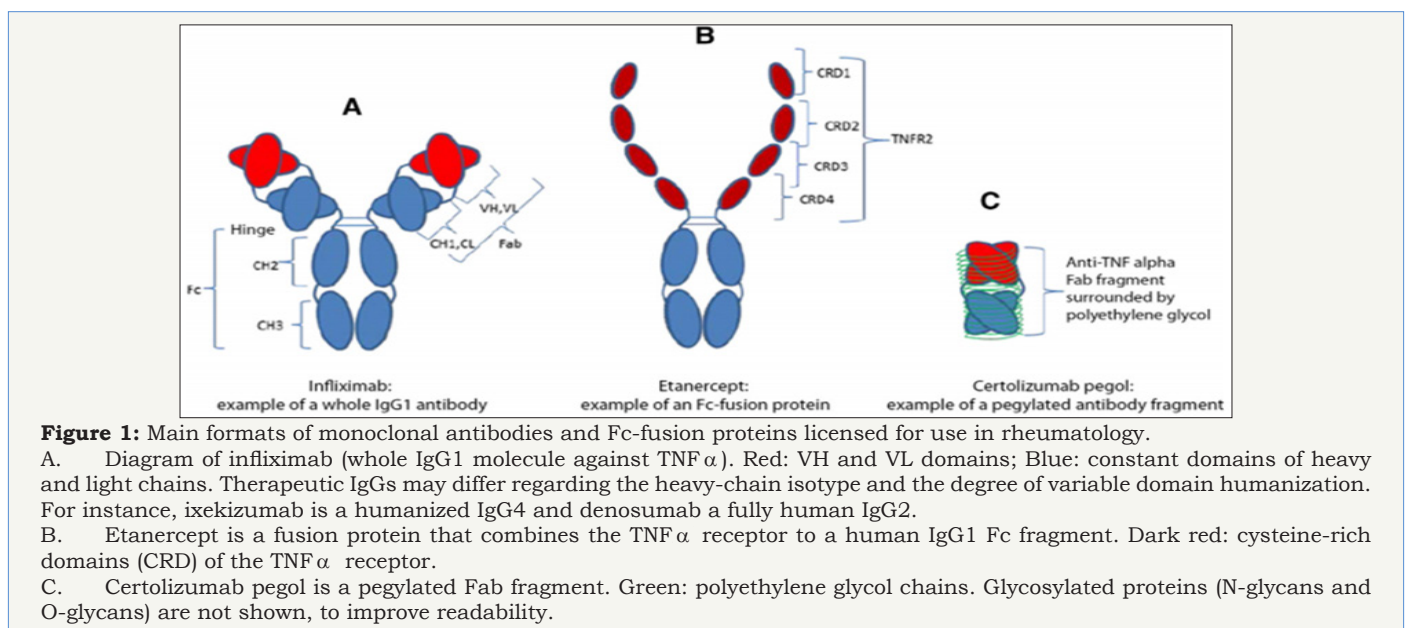
This study aimed to develop mathematical models with important physiological parameters involving immune response for antibody by using Gamma and Psi functions, which could be used in many aspects, for instance to explain the controversies concerning antibody concentration with biological reasons and mathematical calculations, to optimize the antibody yield in large scale production. In this paper, we establish two logarithmically completely monotonic functions involving the gamma function according to two preferred interaction geometries, necessary and sufficient conditions are presented for one of them to be logarithmically completely monotonic. As a consequence, a sharp inequality involving the gamma function is deduced to solve the problems of genetically engineered antibody. This approach would aid to utilize the engineered antibodies and their derivatives for various biomedical applications.

Keywords: Gamma function; Genetically engineered antibody; Logarithmically completely monotonic; Inequality; PSI function

Introduction

Mathematical or computational models have been proposed in different biological subjects in order to explain various complex phenomena. We have designed this study to monoclonal antibodies and Fc-fusion proteins (Figure 1). At present, it is still difficult to predict the optimal structure of antibodies. Topology knowledge can be important in antibody application as well as transformation. Theoretically, we can obtain desired antibodies by using protein/gene engineering technology. For instance, we can transform

the complementarities determining region (CDR) to promote the affinity of the antibody to antigen. Similarly, we could also transform any domain of antibody to make it bind with any desired target. Under this vision, topology is a powerful tool to predict the structure of protein and it will serve to antibody engineering. Our present work tries to explain, and predict, if possible, the change of structure, size and function of antibodies as well as their fragments from a topological perspective.



Main formats of monoclonal antibodies and Fc-fusion proteins licensed for use in rheumatology (Figure 1).

- a. Diagram of infliximab (whole IgG1 molecule against TNF α). Red: VH and VL domains; Blue: constant domains of heavy and light chains. Therapeutic IgGs may differ regarding the heavy-chain iso type and the degree of variable domain humanization. For instance, Ixekizumab is a humanized IgG4 and denosumab a fully human IgG2.
- b. Etanercept is a fusion protein that combines the TNF α receptor to a human IgG1 Fc fragment. Dark red: cysteine-rich domains (CRD) of the TNF α receptor.
- c. Certolizumab pegol is a pegylated Fab fragment. Green: polyethylene glycol chains. Glycosylated proteins (N-glycans and O-glycans) are not shown, to improve readability.

For $\text{Re}(z) > 0$ the classical Euler's gamma function Γ and psi (digamma) function ψ are defined by

$$\Gamma(z) = \int_0^\infty t^{z-1} e^{-t} dt, \quad \psi(z) = \frac{\Gamma'(z)}{\Gamma(z)} \quad (1.1)$$

respectively. The derivatives $\psi^{(n)}(z)$ for $n \in \mathbb{N}$ are known as polygamma functions. For $\psi(z)$ [1], the following series representations are established:

$$\psi(z) = -\gamma - \frac{1}{z} + \sum_{n=1}^\infty \frac{z}{n(z+n)}, \quad (1.2)$$

$$\psi(z) = \ln z + \int_0^\infty \left(\frac{1}{t} - \frac{1}{1-e^{-t}} \right) e^{-zt} dt, \quad (1.3)$$

$$\psi(z) = -\gamma + \int_0^\infty \frac{e^{-t} - e^{-zt}}{1-e^{-t}} dt, \quad (1.4)$$

Where, $\gamma = 0.577215664901\dots$ denotes the Euler's constant. We next recall [2-5] that a function f is said to be completely monotonic on an interval I , if f has derivatives of all orders on I which alternate successively in sign, that is,

$$(-1)^n f^{(n)}(x) \geq 0, \quad (1.5)$$

for all $x \in I$ and for all $n \geq 0$. If inequality (1.5) is strict for all $x \in I$ and all $n \geq 0$, then f is said to be strictly completely monotonic. The classical Bernstein-Widder theorem [6], Theorem

12a states that a function f is completely monotonic on $(0, \infty)$ if and only if it is a Laplace transform of some nonnegative measure μ , that is,

$$f(x) = \int_0^\infty e^{-xt} d\mu(t), \quad (1.6)$$

Where, $\mu(t)$ is non-decreasing and the integral converges for $x > 0$. We recall also [7-9] that a positive function f is said to be logarithmically completely monotonic on an interval I if f has derivatives of all orders on I and

$$(-1)^n [\ln f(x)]^{(n)} \geq 0, \quad (1.7)$$

for all $x \in I$ and for all $n \geq 1$. If inequality (1.7) is strict for all $x \in I$ and all $n \geq 1$, then f is said to be strictly logarithmically completely monotonic. The antibody structure will be changed when it binds certain target (Figure 2), i.e.: antigen, receptor. How to describe the changes in the view of topology? The following cases will explain it in detail.

Diagrammatic representation of VH germ-line diversity (top) leading to domain structure diversity (bottom) (Figure 2). In humans, multiple V, D, J regions are rearranged to produce a VDJ segment that encodes the variable region (in this case the VH). The assembled polypeptide is represented by the classically 'Y' shaped IgG (150kDa) molecule (bottom), where the various regions of the antibody are labeled and the carbohydrates are indicated by pink dots. Unlike humans or mice, chickens use gene conversion to diversify the antibody repertoire. Single genes encode the VH (and VL) and a single VDJ event, which generates minimal diversity, is preceded by insertion of upstream pseudogenes (Ψ VH) in a process that results in donation of diverse sequence fragments into the V region. Chicken antibody (IgY) has a molecular weight of 180 kDa by virtue of an additional constant heavy domain and additional carbohydrate moieties. Cows employ an ultra long CDRH3 repertoire resulting from a VDJ event that utilises a single VH segment (VH Bul), which combines with an elongated D region (DH2) to produce a recombined V-region that undergoes diversification by somatic hyper-mutation. This can introduce or remove cysteine thus, diversifying the disulfide landscape of the repertoire.

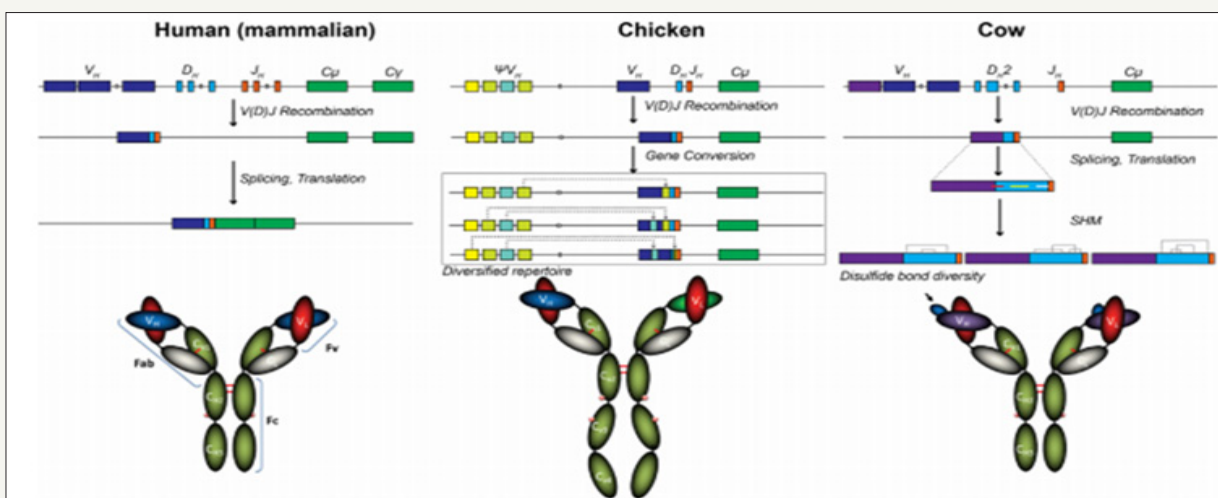


Figure 2: Diagrammatic representation of VH germ-line diversity (top) leading to domain structure diversity (bottom).

It was proved explicitly in [8] and other articles that a logarithmically completely monotonic function must be completely monotonic. In [10], G. D. Anderson et al. proved that the function

$$g(x) = x(\ln x - \psi(x)) \tag{1.8}$$

is strictly decreasing and strictly convex on $(0, \infty)$, with two limits

$$\lim_{x \rightarrow 0} g(x) = 1, \quad \lim_{x \rightarrow \infty} g(x) = \frac{1}{2} \tag{1.9}$$

From (1.9) and the monotonicity of $g(x)$, then the double inequalities

$$\frac{1}{2x} < \ln x - \psi(x) < \frac{1}{x} \tag{1.10}$$

holds for all $x > 0$.

In [11], Theorem [1], by using the well-known Binet's formula, H. Alzer generalized the monotonicity and convexity of $g(x)$, that is, the function

$$g_\alpha(x) = x^\alpha (\ln x - \psi(x)) \tag{1.11}$$

is strictly completely monotonic on $(0, \infty)$ if and only if $\alpha \leq 1$.

In [12], D. Kershaw and A. Laforgia proved that the function $[\Gamma(1+1/x)]^x$ is decreasing on $(0, \infty)$ and $x[\Gamma(1+1/x)]^x$ is increasing on $(0, \infty)$. These are equivalent to the function $[\Gamma(1+x)]^{1/x}$ being increasing and $[\Gamma(1+x)]^{1/x}/x$ being decreasing on $(0, \infty)$, respectively. In Theorem 5, F Qi & Ch-p Chen [13] generalized these functions. They obtained the fact that for all $x > 0$ the function $x^r [\Gamma(1+x)]^{1/x}$ is strictly increasing for $r \geq 0$ and strictly decreasing for $r \leq -1$, respectively. After the papain digestion, the remained antibody functional part (usually the Fab domain), will be smaller and the structure is also changed. These changes can be revealed vividly using topology. Recently [14,15], Theorem 1, F Qi, et al. [14] established another excellent result, which states that for given $y \in (-1, \infty)$ and $\alpha \in (-\infty, \infty)$, let

$$g_{\alpha,y}(x) = \begin{cases} \frac{1}{(x+y+1)^\alpha} \left[\frac{\Gamma(x+y+1)}{\Gamma(y+1)} \right]^{1/x}, & x \in (-y-1, \infty) \setminus \{0\}; \\ \frac{1}{(y+1)^\alpha} \exp[\psi(y+1)], & x = 0. \end{cases} \tag{1.12}$$

The function (1.12) is logarithmically completely monotonic with respect to $x \in (-y-1, \infty)$ if and only if $\alpha \geq \max\{1, 1/(y+1)\}$; and if $\alpha \leq \min\{1, 1/2(y+1)\}$, the reciprocal of the function (1.12) is logarithmically completely monotonic with respect to $x \in (-y-1, \infty)$. Antibodies occur spontaneously gathering and forming dimer, polymer, which will influence their functions (Figure 3). In antibody engineering practice, it urgently needs some measures to overcome this difficulty. From topology perspective, we could understand this issue as follow. Stimulated by the above results, we put forward the function as follows: for given $y \in (0, \infty)$ and real number α , let the function $f_{\alpha,y}(x)$ be defined by

$$f_{\alpha,y}(x) = \frac{1}{x^\alpha} \left[\frac{\Gamma(x+y)}{\Gamma(y)} \right]^{1/x}, \quad x \in (-y, \infty) \setminus \{0\}. \tag{1.13}$$

Our First Result is Contained in the Following Theorem

Theorem 1: For the function (1.13), then the following statements are true:

- i. For any given $y \geq 1$, the function (1.13) is strictly logarithmically completely monotonic with respect to $x \in (-y, \infty) \setminus \{0\}$ if and only if $\alpha \geq 1$;
- ii. For any given $0 < y < 1$, if $\alpha \geq e^{-(1-y)}/y$, then the function (1.13) is strictly logarithmically completely monotonic with respect to $x \in (-y, \infty) \setminus \{0\}$;
- iii. For any given $y > 0$, the reciprocal of the function (1.13) is strictly logarithmically completely monotonic with respect to $x \in (-y, \infty) \setminus \{0\}$ if and only if $\alpha \leq 0$.

Our second result is presented in the following theorem.

Theorem 2: For any given $y \in [1, \infty)$, let the function $h_y(x)$ be defined on $(0, \infty)$ by

$$h_y(x) = \frac{(x+y)^x}{\Gamma(x+y)x^{2x}} e^{\left(\frac{1}{2}y\right)^{x^2-2x}} \exp\left(-\sum_{n=1}^{\infty} \left(\frac{x^2}{n} + \ln\left(1 + \frac{x}{n}\right) - 2x\right)\right), \tag{1.14}$$

Where, γ denotes the Euler's constant, then the function (1.14) is strictly logarithmically completely monotonic with respect to x on $(0, \infty)$. The following corollary can be derived from Theorems 2 immediately.

Corollary 1: For any given $y \geq 1$, the inequality

$$\frac{(x+y)^x}{\Gamma(x+y)x^{2x}} e^{\left(\frac{1}{2}y\right)^{x^2-2x}} < \frac{1}{\Gamma(y)} \exp\left(-\sum_{n=1}^{\infty} \left(\frac{x^2}{n} + \ln\left(1 + \frac{x}{n}\right) - 2x\right)\right) \tag{1.15}$$

holds for all $x > 0$.

Lemma

In order to prove our main results, we need the following lemmas. It is well known that Bernoulli polynomials $B_k(x)$ and Euler polynomials $E_k(x)$ are defined by

$$\frac{te^{tx}}{e^t - 1} = \sum_{n=0}^{\infty} \frac{B_n(x)}{n!} t^n, \quad |t| < 2\pi \tag{2.1}$$

$$\frac{2e^{tx}}{e^t + 1} = \sum_{n=0}^{\infty} \frac{E_n(x)}{n!} t^n, \quad |t| < \pi \tag{2.2}$$

respectively. The Bernoulli numbers B_n are denoted by $B_n = B_n(0)$, while the Euler numbers E_n are defined by. In [16], the following summation formula is given:

$$\sum_{n=0}^{\infty} \frac{(-1)^n}{(2n+1)^{2k+1}} = \frac{\pi^{2k+1} E_k}{2^{2k+1} (2k)!} \tag{2.3}$$

for any nonnegative integer k , which implies

$$B_{2n} = (-1)^{n-1} \frac{2(2n)!}{(2\pi)^{2n}} \sum_{m=1}^{\infty} \frac{1}{m^{2n}}, \quad n \in \mathbb{N} \tag{2.4}$$

In particular, it is known that for all $n \in \mathbb{N}$ we have

$$B_{2n+1} = 0, \quad B_{2n} = (-1)^{n+1} |B_{2n}|, \tag{2.5}$$

$$E_{2n+1} = 0, \quad E_{2n} = (-1)^n |E_{2n}|, \tag{2.6}$$

And the first few nonzero values are

$$B_0 = 0, \quad B_1 = -\frac{1}{2}, \quad B_2 = \frac{1}{6}, \quad B_4 = -\frac{1}{30},$$

$$E_0 = 1, \quad E_2 = -1, \quad E_4 = 5,$$

[17].

The Bernoulli and Euler numbers and polynomials are generalized [18-21].

Lemma 1: [22,23]

For real number $x > 0$ and natural number m , then

$$\ln \Gamma(x) = \ln \sqrt{2\pi} + \left(x - \frac{1}{2}\right) \ln x - x + \sum_{n=1}^m \frac{B_{2n}}{2(2n-1)n} \cdot \frac{1}{x^{2n-1}} + \theta_1 \frac{B_{2(m+1)}}{(2m+1)(2m+2)} \cdot \frac{1}{x^{2m+1}}; \quad 0 < \theta_1 < 1 \tag{2.7}$$

$$\psi(x) = \ln x - \frac{1}{2x} - \sum_{n=1}^m \frac{B_{2n}}{2n} \cdot \frac{1}{x^{2n}} - \theta_2 \frac{B_{2(m+1)}}{2m+2} \cdot \frac{1}{x^{2m+2}}; \quad 0 < \theta_2 < 1 \tag{2.8}$$

$$\psi'(x) = \frac{1}{x} + \frac{1}{2x^2} + \sum_{n=1}^m \frac{B_{2n}}{x^{2n+1}} + \theta_3 \frac{B_{2(m+1)}}{x^{2m+3}}; \quad 0 < \theta_3 < 1 \tag{2.9}$$

$$\psi''(x) = -\frac{1}{x^2} - \frac{1}{x^3} - \sum_{n=1}^m \frac{B_{2n}}{x^{2n+2}} - (2m+3)\theta_4 \frac{B_{2(m+1)}}{x^{2m+4}}; \quad 0 < \theta_4 < 1 \tag{2.10}$$

Remark 1: $\theta_1, \theta_2, \theta_3, \theta_4$ only depend on natural number m .

Lemma 2: [24]

For real number $x > 0$ and natural number n ,

We have

$$\frac{(n-1)!}{x^n} + \frac{n!}{2x^{n+1}} < (-1)^{n+1} \psi^{(n)}(x) < \frac{(n-1)!}{x^n} + \frac{n!}{x^{n+1}}. \tag{2.11}$$

Lemma 3: [1,17]

For real number $x > 0$ and natural number n , we have

$$\ln x = \int_0^\infty \frac{e^{-t} - e^{-xt}}{t} dt, \tag{2.12}$$

$$\frac{\Gamma(n)}{x^n} = \int_0^\infty t^{n-1} e^{-xt} dt, \tag{2.13}$$

$$\psi^{(n)}(x) = (-1)^{n+1} \int_0^\infty \frac{t^n e^{-xt}}{1-e^{-t}} dt. \tag{2.14}$$

Lemma 4:

Let the sequence of functions $u_n(x)$ for $n \in \mathbb{N}$ be defined on $(0, \infty)$ by

$$u_n(x) = \frac{x^2}{n} + \ln \left(1 + \frac{x}{n}\right) - 2x, \tag{2.15}$$

The series $\sum_{n=1}^\infty u_n(x)$ is differentiable on $(0, \infty)$, that is,

$$\left(\sum_{n=1}^\infty u_n(x)\right)' = \sum_{n=1}^\infty \left(\frac{2x}{n} - \frac{2x}{n+x}\right). \tag{2.16}$$

Proof: It is obvious that $\sum_{n=1}^\infty u_n(0) = 0$, therefore $\sum_{n=1}^\infty u_n(x)$ converges at $x = 0$. In order to prove (2.16), we need only to show that the inner closed uniform convergence of the series $\sum_{n=1}^\infty u'_n(x)$ on $(0, \infty)$. From (2.15), We have

$$\sum_{n=1}^\infty u'_n(x) = \sum_{n=1}^\infty \left(\frac{2x}{n} - \frac{2x}{n+x}\right). \tag{2.17}$$

For any interval $[a, b] \subset [0, \infty)$, we have

$$\left|\sum_{n=1}^\infty \left(\frac{2x}{n} - \frac{2x}{n+x}\right)\right| \leq \sum_{n=1}^\infty \frac{2b^2}{n^2} \tag{2.18}$$

for all $x \in [a, b]$. It is easy to check that the series $\sum_{n=1}^\infty \frac{2b^2}{n^2}$ converges, which and

Weierstrass M-test implies that the series $\sum_{n=1}^\infty u'_n(x)$ is inner closed uniformly convergent on $(0, \infty)$. Hence, the series $\sum_{n=1}^\infty u_n(x)$ is differentiable on $(0, \infty)$ and the identity (2.17) holds for $x \geq 0$.

The lemma is proved.

Lemma 5:

For $0 < a \leq 1$ and real number b , let the function $Q_{a,b}(x)$ be defined by

$$Q_{a,b}(x) = \frac{(ax+b)^x}{\Gamma(ax+b)}, \quad x \in \left(-\frac{b}{a}, \infty\right). \tag{2.19}$$

If $b \geq a$, then the function (2.19) satisfies

$$(-1)^n \left[\ln Q_{a,b}(x)\right]^{(n)} > 0 \tag{2.20}$$

for all $x \in \left(-\frac{b}{a}, \infty\right)$ and $n = 2, 3, \dots$.

Proof: Taking the logarithm of $Q_{a,b}(x)$ yields

$$\ln Q_{a,b}(x) = x \ln(ax+b) - \ln \Gamma(ax+b), \tag{2.21}$$

and differentiating $\ln Q_{a,b}(x)$, then

$$\left(\ln Q_{a,b}(x)\right)' = \ln(ax+b) - a\psi(ax+b) - \frac{b}{ax+b} + 1. \tag{2.22}$$

For given integer $n \geq 2$, we get

$$\left(\ln Q_{a,b}(x)\right)^{(n)} = \frac{(-1)^n (n-2)! a^{n-1}}{(ax+b)^{n-1}} - a^n \psi^{(n-1)}(ax+b) - \frac{(-1)^{n-1} (n-1)! b a^{n-1}}{(ax+b)^n}, \tag{2.23}$$

and, by the identities (2.13) and (2.14), (2.23) can be written as

$$\left(\ln Q_{a,b}(x)\right)^{(n)} = (-1)^n a^{n-1} \int_0^\infty \frac{e^{-(ax+bt)t^{n-2}}}{1-e^{-t}} \left((1+bt)(1-e^{-t}) - at\right) dt. \tag{2.24}$$

Let $p(t) = (1+bt)(1-e^{-t}) - at$ and $q(t) = (1+t)(1-e^{-t}) - t$. It is easy to check that

$$q'(t) = te^{-t} > 0, \quad t \in (0, \infty), \tag{2.25}$$

Therefore, $q(t)$ is strictly increasing on $(0, \infty)$, and then $q(t) > q(0) = 0$.

The following two cases will complete the proof of Lemma 5.

a. Case 1: If $0 < a \leq 1 \leq b$, then since $q(t) > 0$ for $t > 0$, we have

$$at \leq t < (1+t)(1-e^{-t}) \leq (1+bt)(1-e^{-t}), \tag{2.26}$$

Which, implies $at < (1+bt)(1-e^{-t})$, and then $p(t) > 0$ for all $t > 0$.

From (2.24), we know that the inequality (2.20) holds for $x \in (-b/a, \infty)$ and integer $n \geq 2$.

b. Case 2: If $0 < a \leq b \leq 1$, then we get

$$p'(t) = b - a + e^{-t}(bt+1-b) \geq bte^{-t} > 0, \quad t \in (0, \infty), \tag{2.27}$$

Therefore, $p(t)$ is strictly increasing on $(0, \infty)$, and then $p(t) > p(0) = 0$.

From (2.24), we know that the inequality (2.20) holds for $x \in (-b/a, \infty)$ and integer $n \geq 2$.

The lemma is proved.

Proof of Theorems

Proof of Theorem 1

For $x \neq 0$ and natural number n , taking the logarithmically differential into consideration yields

$$\left[f_{a,y}(x)\right]^{(n)} = \frac{1}{x^{n+1}} \left[\sum_{k=0}^n (-1)^k \frac{n! x^{n-k} \psi^{(n-k-1)}(x+y)}{(n-k)!}\right] \tag{3.1}$$

$$-(-1)^n n! \ln \Gamma(y) - \alpha(-1)^{n-1} (n-1)! x$$

Where, $\psi^{(-1)}(x+y)$ and $\psi^{(0)}(x+y)$ stand for $\ln \Gamma(x+y)$ and $\psi(x+y)$ respectively.

Furthermore, differentiating $x^{n+1} [f_{\alpha,y}(x)]^{(n)}$ directly gives

$$\left\{ x^{n+1} [f_{\alpha,y}(x)]^{(n)} \right\}' = (-1)^{n-1} x^n \left[(-1)^{n+1} \psi^{(n)}(x+y) - \frac{\alpha(n-1)!}{x^n} \right]. \quad (3.2)$$

Making use of (2.11) and (2.13) shows that for all $n \in \mathbb{N}$ and any fixed $y > 0$, the double inequality

$$\int_0^\infty t^{n-1} e^{-xt} \left[e^{-yt} \left(1 + \frac{1}{2}t \right) - \alpha \right] dt = \frac{(n-1)!}{(x+y)^n} + \frac{1}{2} \frac{n!}{(x+y)^{n+1}} - \frac{\alpha(n-1)!}{x^n} \leq \frac{(-1)^{n-1}}{x^n} \left\{ x^{n+1} [f_{\alpha,y}(x)]^{(n)} \right\} \leq (3.3)$$

$$\frac{(n-1)!}{(x+y)^n} + \frac{n!}{(x+y)^{n+1}} - \frac{\alpha(n-1)!}{x^n} = \int_0^\infty t^{n-1} e^{-xt} [e^{-yt}(1+t) - \alpha] dt$$

holds for all $x \in (-y, \infty) \setminus \{0\}$ and $\alpha \in (-\infty, \infty)$.

For any fixed $y \in (0, \infty)$, let $u(t)$ and $v(t)$ be defined on $(-\infty, \infty)$ by

$$u(t) = e^{-yt} \left(1 + \frac{1}{2}t \right) \text{ and } v(t) = e^{-yt} (1+t)$$

respectively.

Differentiating $u(t)$ and $v(t)$ directly, we obtain

$$u'(t) = e^{-yt} \left(\frac{1}{2} - y - \frac{1}{2}yt \right), \quad (3.4)$$

$$v'(t) = e^{-yt} (1 - y - yt). \quad (3.5)$$

Therefore, for given $y \in (0, \infty)$ we have

$$u'(t) \begin{cases} > 0, & t < 1/y - 2 \\ < 0, & t > 1/y - 2 \end{cases}, \quad (3.6)$$

And

$$v'(t) \begin{cases} > 0, & t < 1/y - 1 \\ < 0, & t > 1/y - 1 \end{cases}. \quad (3.7)$$

From (3.6) and (3.7), we conclude that for all $t > 0$ we obtain

$$u(t) > 0, \quad (3.8)$$

And

$$v(t) \begin{cases} < 1, & y \geq 1 \\ \leq e^{-(1-y)/y}, & 0 < y < 1 \end{cases}. \quad (3.9)$$

From (3.3) and (3.8)-(3.9), it is easy to see that

$$\frac{(-1)^{n-1}}{x^n} \left\{ x^{n+1} [f_{\alpha,y}(x)]^{(n)} \right\} \begin{cases} > 0, & \text{if } \alpha \leq 0 \text{ for fixed } y > 0 \\ < 0, & \text{if } \alpha \geq 1 \text{ for fixed } y \geq 1 \\ < 0, & \text{if } \alpha \geq e^{-(1-y)/y} \text{ for fixed } 0 < y < 1 \end{cases} \quad (3.10)$$

for all $n \in \mathbb{N}$ and all $x \in (-y, \infty) \setminus \{0\}$.

On the one hand, if $x \in (0, \infty)$, then the inequalities (3.10) can be equivalently

changed into

$$\left\{ x^{2k+1} [f_{\alpha,y}(x)]^{(2k)} \right\} \begin{cases} < 0, & \text{if } \alpha \leq 0 \text{ for fixed } y > 0 \\ > 0, & \text{if } \alpha \geq 1 \text{ for fixed } y \geq 1 \\ > 0, & \text{if } \alpha \geq e^{-(1-y)/y} \text{ for fixed } 0 < y < 1 \end{cases} \quad (3.11)$$

And

$$\left\{ x^{2k} [f_{\alpha,y}(x)]^{(2k-1)} \right\} \begin{cases} > 0, & \text{if } \alpha \leq 0 \text{ for fixed } y > 0 \\ < 0, & \text{if } \alpha \geq 1 \text{ for fixed } y \geq 1 \\ < 0, & \text{if } \alpha \geq e^{-(1-y)/y} \text{ for fixed } 0 < y < 1 \end{cases} \quad (3.12)$$

for $k \in \mathbb{N}$.

From (3.1), then simple computation shows that

$$\lim_{x \rightarrow 0} x^{n+1} [f_{\alpha,y}(x)]^{(n)} = 0 \quad (3.13)$$

for all $n \in \mathbb{N}$ and any given $y \in (0, \infty)$. As a result,

$$x^{2k+1} [f_{\alpha,y}(x)]^{(2k)} \begin{cases} < 0, & \text{if } \alpha \leq 0 \text{ for fixed } y > 0 \\ > 0, & \text{if } \alpha \geq 1 \text{ for fixed } y \geq 1 \\ > 0, & \text{if } \alpha \geq e^{-(1-y)/y} \text{ for fixed } 0 < y < 1 \end{cases} \quad (3.14)$$

and

$$x^{2k} [f_{\alpha,y}(x)]^{(2k-1)} \begin{cases} > 0, & \text{if } \alpha \leq 0 \text{ for fixed } y > 0 \\ < 0, & \text{if } \alpha \geq 1 \text{ for fixed } y \geq 1 \\ < 0, & \text{if } \alpha \geq e^{-(1-y)/y} \text{ for fixed } 0 < y < 1 \end{cases} \quad (3.15)$$

for all $k \in \mathbb{N}$ and all $x > 0$.

Therefore, (3.14) and (3.15) imply

$$(-1)^n [f_{\alpha,y}(x)]^{(n)} \begin{cases} < 0, & \text{if } \alpha \leq 0 \text{ for fixed } y > 0 \\ > 0, & \text{if } \alpha \geq 1 \text{ for fixed } y \geq 1 \\ > 0, & \text{if } \alpha \geq e^{-(1-y)/y} \text{ for fixed } 0 < y < 1 \end{cases} \quad (3.16)$$

for all $n \in \mathbb{N}$ and all $x > 0$.

Hence, if either $\alpha \geq e^{-(1-y)/y}$ for given $0 < y < 1$ or $\alpha \geq 1$ for given $y \geq 1$, the function (1.13) is strictly logarithmically completely monotonic with respect to x on $(0, \infty)$, and if $\alpha \leq 0$ for given $y > 0$, so is the reciprocal of the function (1.13).

On the other hand, if $x \in (-y, 0)$ for any given $y > 0$, then (3.10) implies

$$\left\{ x^{n+1} [f_{\alpha,y}(x)]^{(n)} \right\} \begin{cases} < 0, & \text{if } \alpha \leq 0 \text{ for fixed } y > 0 \\ > 0, & \text{if } \alpha \geq 1 \text{ for fixed } y \geq 1 \\ > 0, & \text{if } \alpha \geq e^{-(1-y)/y} \text{ for fixed } 0 < y < 1 \end{cases} \quad (3.17)$$

for $n \in \mathbb{N}$.

In view of (3.13), we can conclude that

$$x^{n+1} [f_{\alpha,y}(x)]^{(n)} \begin{cases} > 0, & \text{if } \alpha \leq 0 \text{ for fixed } y > 0 \\ < 0, & \text{if } \alpha \geq 1 \text{ for fixed } y \geq 1 \\ < 0, & \text{if } \alpha \geq e^{-(1-y)/y} \text{ for fixed } 0 < y < 1 \end{cases} \quad (3.18)$$

for $n \in \mathbb{N}$. It is obvious that (3.18) is equivalent to that (3.14) and (3.15) hold for any given $y > 0$ and $x \in (-y, 0)$. Therefore, it is easy to prove similarly that (3.16) is also valid on $x \in (-y, 0)$ for any given $y > 0$ and all $n \in \mathbb{N}$.

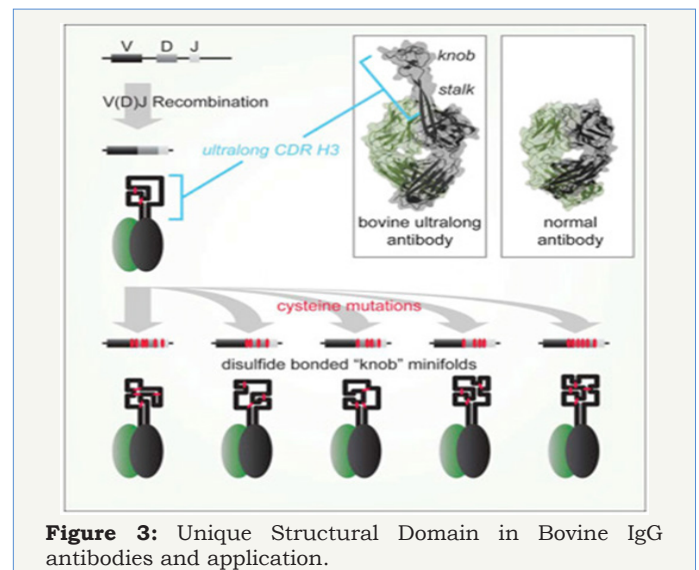


Figure 3: Unique Structural Domain in Bovine IgG antibodies and application.

The amino acid of antibody/protein possesses different preferences. Thus we can conduct site-directed mutation to promote the affinity and/or hydrophilic with the prediction of topology. For example, bovine antibodies have an unusual structure comprising a β -strand ‘stalk’ domain and a disulphide-bonded ‘knob’ domain in CDR3 (Figure 3). Attempts have been made to utilize such amino acid preference for antibody drug development. Unique Structural Domain in Bovine IgG antibodies and application (Figure 3).

Consequently, the function (1.13) is the same logarithmically completely monotonic on $(-y, 0)$ as on $(0, \infty)$, that is, if either $\alpha \geq e^{-(y)} / y$ for given $0 < y < 1$ or $\alpha \geq 1$ for given $y \geq 1$, the function (1.13) is strictly logarithmically completely monotonic with respect to x on $(-y, 0)$, and if $\alpha \leq 0$ for given $y > 0$, so is the reciprocal of the function (1.13).

Conversely, we assume that the reciprocal of the function (1.13) is strictly logarithmically completely monotonic on $(-y, \infty) \setminus \{0\}$ for any given $y > 0$. Then we have for any given $y > 0$ and all $x > 0$

$$f'_{\alpha, y}(x) = \frac{\ln \Gamma(x+y) - x\psi(x+y) - \ln \Gamma(y)}{x^2} + \frac{\alpha}{x} < 0, \tag{3.19}$$

Which implies

$$\alpha < - \frac{\ln \Gamma(x+y) - x\psi(x+y) - \ln \Gamma(y)}{x}. \tag{3.20}$$

By L'Hospital's rule, we have

$$\lim_{x \rightarrow 0} \frac{-\ln \Gamma(x+y) + x\psi(x+y) + \ln \Gamma(y)}{x} = 0 \tag{3.21}$$

for any given $y > 0$. By virtue of (3.20) and (3.21), we conclude that the necessary condition for the reciprocal of the function (1.13) to be strictly logarithmically completely monotonic is $\alpha \leq 0$.

If the function (1.13) is logarithmically completely monotonic on $(-y, \infty) \setminus \{0\}$ for any given $y > 0$, then the inequality (3.19) and (3.20) are reversed for any given $y > 0$ and all $x > 0$.

By utilizing (2.7) and (2.8), it is easy to see that

$$\lim_{x \rightarrow \infty} \frac{-\ln \Gamma(x+y) + x\psi(x+y) + \ln \Gamma(y)}{x} = 1 \tag{3.22}$$

for any given $y > 0$. In fact, it is not difficult to show that the necessary condition for the function (1.13) to be strictly logarithmically completely monotonic is $\alpha \geq 1$. The proof of Theorem 1 is completed.

Proof of Theorem 2

Taking the logarithm of $h_y(x)$ gives

$$\ln h_y(x) = \ln \frac{(x+y)^x}{\Gamma(x+y)} - \ln x^x + \left(\frac{1}{2} - \gamma\right)x^2 - 2x + \sum_{n=1}^{\infty} \left(\frac{x^2}{n} + \ln \left(1 + \frac{x}{n}\right)^{2n} - 2x\right). \tag{3.23}$$

Let,

$$\mu(x) = \ln \frac{(x+y)^x}{\Gamma(x+y)}, \tag{3.24}$$

$$\omega(x) = -\ln x^x + \left(\frac{1}{2} - \gamma\right)x^2 - 2x + \sum_{n=1}^{\infty} \left(\frac{x^2}{n} + \ln \left(1 + \frac{x}{n}\right)^{2n} - 2x\right), \tag{3.25}$$

Then,

$$\ln h_y(x) = \mu(x) + \omega(x). \tag{3.26}$$

In view of Lemma 4, straightforward calculation gives

$$\begin{aligned} (\ln h_y(x))' &= \ln(x+y) + \frac{x}{x+y} - \psi(x+y) - 2x \ln x + 2\gamma x + 2 - \sum_{n=1}^{\infty} \left(\frac{2x}{n} - \frac{2x}{n+x}\right) \\ &= \mu'(x) + \omega'(x). \end{aligned} \tag{3.27}$$

By virtue of (1.2), the identity (3.27) is equivalent to

$$\begin{aligned} (\ln h_y(x))' &= \ln(x+y) + \frac{x}{x+y} - \psi(x+y) - 2x(\ln x - \psi(x)) \\ &= \mu'(x) + \omega'(x). \end{aligned} \tag{3.28}$$

By Lemma 5, we know that $\mu'(x)$ is strictly increasing on $(0, \infty)$, which and (1.10) imply the limit of $\mu'(x)$ equals 1 as $x \rightarrow \infty$, therefore

$$\mu'(x) < 1 \tag{3.29}$$

holds for all $x > 0$.

We know that $g(x)$ is strictly completely monotonic on $(0, \infty)$, where $g(x)$ is defined by (1.8), hence for given integer $n \geq 0$, the inequality

$$(-1)^{n+1} (\omega'(x))^{(n)} > 0 \tag{3.30}$$

holds for all $x > 0$.

And then by using inequality (1.9) and (1.10), we get

$$-2 < \omega'(x) < -1 \tag{3.31}$$

for all $x > 0$.

From (3.29) and (3.31), we conclude that

$$(\ln h_y(x))' = \mu'(x) + \omega'(x) < 0 \tag{3.32}$$

for all $x > 0$. Utilizing Lemma 5 and (3.30), for given integer $n \geq 2$, it is easy to see that

$$(-1)^n (\ln h_y(x))^{(n)} = (-1)^n (\mu(x))^{(n)} + (-1)^n (\omega(x))^{(n)} > 0 \tag{3.33}$$

for all $x > 0$.

Theorem 2 follows from (3.32) and (3.33). Thus the proof of Theorem 2 is completed [25-31].

Conclusion

In conclusion we establish two new logarithmically completely monotonic functions involving the gamma function according to two preferred interaction geometries, and a sharp inequality involving the gamma function is deduced to solve the problems of genetically engineering antibody. It is necessary to address, many other aspects (such as thermal condition, alkalinity or acidity, adhesion of antibody) are also playing key roles in antibody functioning, which could be also understood from bio-mathematical perspective, and such knowledge will be in return useful for biomedical application of antibodies as well as proteins.

Acknowledgement

This work was supported by the Fundamental Research Funds for the Central Universities (2014YB030), Ministry of Education and State Administration of Foreign Experts Affairs ‘Overseas Teacher’ project (MS2011XBNL057), the Key Construction Pro-



gram (2015SD0018) of International Cooperation Base in S&T, Shaanxi Province, China.

References

- Anon (2016) US Broiler Performance.
- Petracci M, Mudalal S, Soglia F, Cavani C (2015) Meat quality in fast-growing broiler chickens. *World's Poultry Science Journal* 71(2): 363-374.
- Song DJ, King AJ (2015) Effects of heat stress on broiler meat quality. *World's Poultry Science Journal* 71(4): 701-709.
- Sahin K, Smith MO (2016) Regulation of transcription factors by the epigallocatechin-3-gallate in poultry reared under heat stress. *World's Poultry Science Journal* 72(2): 299-306.
- Kapell DNRG, Hill WG, Neeteson AM, McAdam J, Koerhuis ANM, et al. (2012) Twenty-five years of selection for improved leg health in purebred broiler lines and underlying genetic parameters. *Poultry Science* 91(12): 3032-3043.
- Honkatukia M, Tuiskula-HM, Ahola V, Uimari P, Schmutz M, et al. (2011) Mapping of QTL affecting incidence of blood and meat inclusions in egg layers *BMC Genetics* 12: 55.
- Dunn IC (2004) Breeding strategies to improve the egg's natural defense. *World's Poultry Science Journal* 60(4): 458-468.
- Bain MM, MacDade K, Schmutz M, Preisinger R, Waddington D, et al. (2009) The deposition of the cuticle on laying hens eggs is a moderately heritable trait. *British Poultry Abstracts* 5(1): 17-18.
- Kemps BJ, De Ketelaere B, Bamelis FR, Mertens K, Decuyper EM, et al. (2007) Albumen freshness assessment by combining visible near-infrared transmission and low-resolution proton nuclear magnetic resonance spectroscopy. *Poultry Science* 86(4): 752-759.
- Lerner IM, Taylor LW, Lowry DC (1951) Selection for increased incidence of blood spots in white leghorns. *Poultry Science* 30: 748-757.
- Noda K, Nakamura A, Kino K, Kato Y (2007) Genetic improvement by selection for decrease of the meat spots in Nagoya breed. *Japanese Journal of Poultry Science* 44(1): 17-22.
- Albers G (2010) Genomic selection in poultry and pig breeding - a breakthrough technology? *British Poultry Abstracts* 6: 13.
- Tuiskula-HM, Honkatukia M, Preisinger R, Schmutz M, de Koning DJ, et al. (2011) Quantitative trait loci affecting eggshell traits in an F(2) population. *Animal Genetics* 42(3): 293-299.
- Ministry of Agriculture and Cooperatives (2005) Socio-economic impact assessment for the avian influenza crisis: gaps and links between poultry and poverty in smallholders. Department of Livestock Development, Ministry of Agriculture and Cooperatives, The Kingdom of Thailand. (FAO/TCP/RAS/3010e).
- Connolly A (2012) A vision for 2020, the future of poultry. XXIV World's Poultry Congress 5 - 9 August - 2012, Salvador, Bahia, Brazil.
- Leinonen I, Williams AG, Waller AH, Kyriazakis I (2013) Comparing the environmental impact of alternative protein crops in poultry diets: the consequences of uncertainty. *Agricultural systems* 121: 33-42.
- Leinonen I, Williams AG, Wiseman J, Guy J, Kyriazakis I (2012) Predicting the environmental impact of chicken systems in the UK through a life cycle assessment: broiler production systems. *Poultry Science* 91(1): 8-25.
- Leinonen I, Williams AG, Wiseman J, Guy J, Kyriazakis I (2012) Predicting the environmental impact of chicken systems in the UK through a life cycle assessment: egg production systems. *Poultry Science* 91(1): 26-40.
- Vaarst M, Steinfeldt S, Horsted K (2015) Sustainable development perspectives of poultry production. *World's Poultry Science Journal* 71(4): 609-620.
- Hermansen JE, Horsted K, Kongsted AG (2014) Meat production in organic farming. In: Dikeman M and Devine C (Eds) *Encyclopedia of Meat Sciences* 2e, 2: 199-203.
- Berardinelli A, Ragni L, Giunchi A, Gradari P, Guarnieri A (2008) Physical-mechanical modifications of eggs for food-processing during storage. *Poultry Science* 87(10): 2117-2125.
- Cheema MA, Qureshi MA, Havenstein GB (2003) A comprehension of the immune response of a 2001 commercial broiler with a 1957 randombred broiler strain when fed representative 1957 and 2001 broiler diets. *Poultry Science* 82(10): 1519-1529.
- Dekkers JC (2004) Commercial application of marker - and gene-assisted selection in livestock: Strategies and lessons. *Journal of Animal Science* 82(E-Suppl): E313-E328.
- Elshire RJ, Glaubitz JC, Sun Q, Poland JA, Kawamoto K, et al. (2011) A Robust, Simple Genotyping-by-Sequencing (GBS) Approach for High Diversity Species. *PLoS One* 6(5): e19379.
- Elson HA (2015) Poultry welfare in intensive and extensive production systems. *World's Poultry Science Journal* 71(3): 449-460.
- Havenstein GB, Ferket PR, Qureshi MA (2003) Growth, liveability, and feed conversion of 1957 versus 2001 broilers when fed representative 1957 and 2001 broiler diets. *Poultry Science* 82(10): 1500-1508.
- Laughlin K (2009) Temperton Fellowship Report No.15, Harper Adams University College, Newport, Shropshire, UK, England.
- Li Y, Willer CJ, Ding J, Scheet P, Abecasis GR (2010) MaCH: using sequence and genotype data to estimate haplotypes and unobserved genotypes. *Genetic Epidemiology* 34(8): 816-834.
- Nys Y, Bain M, Immerseel, Van F (2011) Improving the safety and quality of eggs and egg products, Cambridge: Woodhead Publishing, Cambridge, UK, pp. 581.
- Van Der Laan MHP (2012) Breeding for genetic resistance to disease. *World's Poultry Congress* 5 - 9, Salvador, Bahia, Brazil.
- Williams AG, Audsley E, Sandars DL (2009) A Lifecycle Approach to Reducing the Environmental Impacts of Poultry Production. *Proceedings and Abstracts, 17th European Symposium on Poultry Nutrition S4.1, p. 7.*