Longterm existence of solutions of a reaction diffusion system with non-local terms modeling an immune response—An interpretation-orientated proof

Cordula Reisch *, Dirk Langemann

TU Braunschweig, Institute for Partial Differential Equations, Universitätssplatz 2, 38106 Braunschweig, Germany

**A R T I C L E I N F O**

**A B S T R A C T**

This paper shows the global existence and boundedness of solutions of a reaction diffusion system modeling liver infections. The existence proof is presented step by step and the focus lies on the interpretation of intermediate results in the context of liver infections which is modeled. Non-local effects in the dynamics between the virus and the immune system cells coming from the immune response in the lymphs lead to an integro-partial differential equation. While existence theorems for parabolic partial differential equations are textbook examples in the field, the additional integral term requires new approaches to proving the global existence of a solution. This allows to set up an existence proof with a focus on interpretation leading to more insight in the system and in the modeling perspective at the same time.

We show the boundedness of the solution in the $L^1(\Omega)$- and the $L^\infty(\Omega)$-norms, and use these results to prove the global existence and boundedness of the solution. A core element of the proof is the handling of oppositely acting mechanisms in the reaction term, which occur in all population dynamics models and which results in reaction terms with opposite monotonicity behavior. In the context of modeling liver infections, the boundedness in the $L^\infty(\Omega)$-norm has practical relevance: Large immune responses lead to strong inflammations of the liver tissue. Strong inflammations negatively impact the health of an infected person and lead to grave secondary diseases. The gained rough estimates are compared with numerical tests.

1. Introduction

Modeling the coupled dynamics of virus and the immune system during a liver infection caused by a hepatitis virus is challenging because the mechanisms behind persisting infections over month or years are still unknown.¹ An opportunity for overcoming the problem of unknown mechanisms on the cell scale contains two integrative changes. First, the modeling scale is changed from the cell scale towards a mesoscopic scale on the length scale of a few centimeters. Second, the mechanisms, which are unknown in detail, are replaced by integrative mechanisms representing the commonly accepted properties of the unknown mechanisms. This change of view results in a compact model of partial differential equations.

Modeling inflammations with differential equations is a widely used approach. For example in Refs. 2, 3, atherogenesis as a particular inflammation is modeled with reaction diffusion equations. In Ref. 2, instable states are interpreted as persisting infections, whereas in Ref. 3 traveling waves are interpreted as persisting infections. Reaction diffusion systems for modeling the dynamics of liver infections are presented in Refs. 4, 5. In Refs. 6–9 systems of ordinary equations are used for modeling the total amount of immune system cells and virus during a hepatitis C liver infection.

In Refs. 9–13 liver inflammations are modeled by using reaction diffusion equations describing the virus concentration and the T cell population during an infection. As a specific feature, the reaction diffusion equations include a space-dependent and non-local term describing the inflow of T cells in a small part of the modeled region. The amount of inflowing T cells depends on the total virus amount in the regarded part of the liver. The dependency on the total virus amount is represented by an integral term over the whole domain. The non-local term models the T cell dispersal starting in the lymphs.

The description of the inflow region, called portal field, reflects some important parts of the real liver structure. Therefore, the term is desirable and necessary for modeling liver infections even if it makes the mathematical analysis of the model more difficult. One challenging task caused by the non-local and space-depending inflow term is the proof of the longterm existence of a solution. Often used results for parabolic partial differential equations are based on Lipschitz continuous reaction functions with respect to the state variable or require monotonous reaction functions. Due to the integral term and the oppositely acting mechanisms, these results are not directly applicable to the system modeling the dynamics of liver infections, see Section 3.

In this paper, the longterm existence and boundedness of solutions of the model proposed in Ref. 9 is proven and the results are interpreted...
We regard, as a simplification, the boundary $\partial \Omega$ of the bounded domain $\Omega$ to be impermeable. This results in zero flux or homogeneous Neumann boundary conditions.

Using as few mechanisms as possible, see Ref. 12, we find the predator–prey model

$$
\begin{align*}
  u_t &= u(u) - yu + a \Delta u & \text{for } x \in \Omega, t > 0, \\
  v_t &= j[u] - \eta(1 - u) + \beta \Delta v & \text{for } x \in \Omega, t > 0, \\
  w(0, x) &= u_0(x), & v(0, x) &= v_0(x) & \text{for } x \in \Omega, \\
  0 &= \nabla u \cdot n = \nabla v \cdot n & \text{for } x \in \partial \Omega, t > 0
\end{align*}
$$

with a growth function $u(u)$ describing the non-linear growth of the virus in absence of other mechanisms and the non-local inflow $j = j[u](x)$ of T cells. The constants $a$ and $\beta$ describe the strength of diffusion. The reaction diffusion system in Eq. (1) contains the predator term $\gamma u v$ diminishing the virus in presence of the immune response $v$, and the decay term $\eta(1 - u)v$ describing the fade out of the immune response in absence of any virus.

The growth rate $w$ in Ref. 9 describes a logistic growth of the virus with a strong Allee effect, i.e.

$$
u(u) = (1 - u) \frac{u - u_{\text{min}}}{u + k} \quad \text{with } 0 < u_{\text{min}} < 1 \quad \text{and} \quad k > 0.$$

The minimal density for the survival of the virus is $u_{\text{min}}$. Otherwise, the virus is locally attacked and it decreases without the secondary immune response from the lymphs. The parameter $k$ is a small parameter fitting the growth in Eq. (2) to a pure logistic growth for values $u$ close to 1.

As usual in population dynamics models, the reaction functions in Eq. (1) contain terms with opposite monotonicity behavior. The growth term $u(u)$ and the decay term $-\gamma u v$ act oppositely for $u$ in the equation for $u$. Just like the inflow term $j[u]$ and the decay term $-\eta(1 - u)v$ do for $v$.

Remark 2. The particular choice of the growth rate makes $w(u_{\text{min}}) = 0$ and $w(1) = 0$, and it is positive between the zeros. Furthermore $w$ behaves asymptotically like $1 - u$ for large $u$, and we find that $w$ is increasing in the interval $[0, u_{\text{min}}]$. Thus, the minimal value $w(u)$ for $u \in [0, 1]$ is $w(0) = -u_{\text{min}}/k$.

Opposite to the classical Lotka–Volterra model, the Allee effect allows a population to become extinct. Besides, the Allee effect does not influence qualitatively the system behavior for larger values $u$.

Remark 3. Eq. (2) norms the capacity of the logistic growth to 1 because $w(u) < 0$ for all $u > 1$. There is no loss of generality because the normalization of $u$ is a pure scaling. A possible $u$ with $u(t, x) > 1$ at some $x$ decays in finite time below 1. Due to this realistic property of the model, system (1) is suitable only for $u(t, x) \leq 1$.

The non-local inflow term is

$$
\begin{align*}
  j[u](x) &= \delta \chi_\theta(x) \int_\Omega u(t, x) \, dx = \delta \chi_\theta(x) U(t) \quad \text{where} \quad U(t) = \int_\Omega u(t, x) \, dx
\end{align*}
$$

is the total amount of virus, and $\chi_\theta(x)$ is a non-negative function with support $\supp \chi_\theta(x) = \theta \subset \Omega$ and

$$
\int_\Omega \chi_\theta(x) \, dx = \int_\Omega \chi_\theta(x) \, dx = 1.
$$

As a realistic inflow, we consider $\chi_\theta$ to be at least a bounded and piecewise continuous function.

A non-smooth example for $\chi_\theta$ is the characteristic function on the subdomain $\theta \subset \Omega$ providing $1/|\theta|$ for $x \in \theta$ and 0 elsewhere.

Analogously, to Eq. (3), we define the integral of the non-negative $v$ over $\Omega$ as

$$
V(t) = \|v(t, \cdot)\|_{L^1(\Omega)} = \int_\Omega v(t, x) \, dx.
$$

This expression gives the total amount of T cells in $\Omega$ and is important for the harm of an infected organism.

C. Reisch and D. Langemann

Partial Differential Equations in Applied Mathematics 6 (2022) 100446
The second mechanism is a decrease in growth by diminishing the prey population in the way that an increasing predator population slows down its own growth in the classical Lotka-Volterra system because the growth of the predator population is negative. Additional to the dynamics of the classical predator-prey model, there is a positive influence on $v$ just depending on $u$, compare the thicker line. This might lead to an unbounded growth of $v$, what is part of our discussion.

As a third mechanism, the term $\eta uv$ spatially inhomogeneous. The used parameters and the shape and size of the domain $\Omega$ correspond to $\eta uv$ in the model. The reaction terms in system (1) contain oppositely acting monotonous terms, compare the classical Lotka-Volterra system. Consequently, the model equations in (1) are not globally Lipschitz continuous with respect to $u$ and $v$ nor monotonous. Even if $u$ is bounded by construction by an upper limit $1$, an a-priori upper bound is not obvious. We show the existence of a global upper bound for $v$ by the inflow term $j[u]$.

Fig. 1 shows a state chart of the local reaction mechanisms. It is simplified and abstracts from the space dependency of the increase of $v$ by the inflow term $j[u]$. The non-local inflow term $j[u]$ is a considerate expansion of the classical Lotka-Volterra system because the growth of the predator depends directly on the prey in Eq. (1). That enforces the feedback loop in the way, that an increasing predator population slows down its own growth by diminishing the prey population in $u$, compare (−) in Fig. 1.

The interplay of oppositely acting mechanisms leads to interesting solutions. We observe in Ref. 9 that the system (1) has solutions which can be divided into two qualitative different types. On the one hand, there are solutions tending towards zero. On the other hand, we find solutions with a tendency towards a stationary state which is spatially inhomogeneous. The used parameters and the shape and size of the domain $\Omega$ control towards which stationary state the solution is tending. See Refs. 9, 10, 12, 13 for further details on the analytical results.

As the model was found in the context of liver infections, we interpret the two qualitative different solution types as different infection courses. Solutions with a tendency towards zero are associated with healing courses, see Fig. 2. The immune system is able to kill all infected cells during an active phase and therefore, the virus vanishes. Afterwards, the immune reaction fades out and the T cell amount tends towards zero as well, see Fig. 2(b).

In Ref. 19, p. 188, an existence theorem for a reaction diffusion term is shown. The proof is based on the Schauder fixed point theorem and uses a non-local and space-dependent term for modeling the biological structure of the application. The analysis of this model leads to an interesting new problem which cannot be handled easily by standard approaches. Besides, we are interested in a proof using interpretable intermediate steps for gaining a deeper understanding of the systems dynamics.

Of course, there are many theorems for the existence of a solution of a reaction diffusion system or more general a parabolic partial differential equation. In this section, we mention some important results on the existence of solutions for reaction diffusion equations and explain, why they cannot be applied directly to the system (1). There are at least two main approaches often used in proofs of existence theorems for parabolic partial differential equations. One approach uses fixed point theorems, like the Banach fixed point theorem, the Brouwer fixed point theorem and from this following the Schauder and the Leray–Schauder fixed point theorems. The second approach uses semigroup theory, see Refs. 17, 18.

The first approach using fixed point theorems can be found for example in Ref. 16, p. 536. There, the existence and uniqueness of solutions is shown under the requirement, that the local reaction function $f$ is Lipschitz continuous with respect to $q = (u,v)^T$. This requirement is used for showing the contraction of the operator for the fixed point theorem. Additionally, the theorem in Ref. 16 requires Dirichlet boundary conditions.

In Ref. 19, p. 188, an existence theorem for a reaction diffusion system with Lotka-Volterra reaction terms is shown. The proof is based on the Schauder fixed point theorem and uses a-priori bounds for the state variables. There are several proofs for monotonous reaction functions as well, see Ref. 20, p. 120.

Unfortunately, the reaction functions in Eq. (1) are neither globally Lipschitz continuous with respect to $u$ and $v$, nor monotonous. Even if $u$ is bounded by construction by an upper limit $1$, an a-priori upper bound for $v$ is not obvious. We show the existence of a global upper bound in Section 3.3.

Existence results using a semigroup approach are based on limited growth conditions, for example Ref. 21, p. 276 or Ref. 22, p. 75. Due to the non-local integral term, the nonlinear terms and the unavailable a-priori bound for $v$, the system in Eq. (1) does not fulfill the requirements for these existence results. As already mentioned, the existence of a finite a-priori bound for $v$ and therefore the boundedness of $v$ in the $L^\infty$-norm is a relevant question concerning the application in modeling liver infections.
Fig. 2. Numerical simulation with a solution interpreted as healing infection course. Used parameter values are $u_{\min} = 0.05$, $\kappa = 0.01$, $\gamma = 0.9$, $\delta = 3.7$, $\eta = 0.9$, $\alpha = 0.6$, and $\beta = 0.3$. The initial conditions for $t_0 = 0$ (bright mesh) show the amount of virus and $T$ cells right after the activation of the immune response. In (b) $T$ cells enter the domain through an area $\Theta$ around $(x_1, x_2) = (1, 1)$. The virus is killed by the $T$ cells and decays for $t_1 = 0.75$ and $t_2 = 3$ (dark mesh). The amount of $T$ cells reduces due to the very low virus concentration. Both population vanish after an active phase.

Fig. 3. Numerical simulation with a solution interpreted as chronic or persisting infection course. Used parameter values are the same as in Fig. 2 but $\delta = 0.7$ and $\eta = 0.9$. The time steps are $t_0 = 0$ (bright), $t_1 = 10.5$, $t_2 = 30$ (dark). Starting with the same initial conditions as in Fig. 2, the virus and the $T$ cells persist in the whole domain. The $T$ cell amount is higher around the portal field $\Theta$. There is nearly no difference between the spread at $t_1$ and at $t_2$.

Fig. 4. Numerical simulations according to those in Figs. 2 and 3. (a) Total virus $U$ and $T$ cell amount $V$ during a chronic or healing infection course over the time. (b) Summed dynamics of a healing or chronic infection course in phase space.

Results for reaction diffusion systems with non-local effects can be divided into results for nonlinear diffusion and nonlinear reaction terms. The global existence of solutions for systems with nonlinear diffusion,

$$u_t = f(u_1, \ldots, u_m) + \Delta \phi(u_i)$$

with homogeneous Dirichlet boundary conditions is shown in Ref. 22. The results yield if the solutions are non-negative and the total mass is controlled. Additionally, an a-priori estimate in the $L^1(\Omega)$-norm for the nonlinear reaction functions is required.

In Ref. 23, the reaction diffusion equation

$$u_t = \Delta u + \int_{\Omega} \psi(y)$$

with a non-local term and with homogeneous Dirichlet boundary conditions is analyzed. The global existence of non-negative solutions is shown for any $p > 1$.

As a third example, the global existence of solutions of the general formulation

$$u_t + A(u) = F(u),$$

where $A$ is a parabolic operator and $F$ is bounded in the $L^2(\Omega)$-norm is shown in Ref. 24.

The results are mainly for single equations instead of systems, and the requirements are not fulfilled for system (1). Again, the system with coupled equations and an integral term require new approaches for proving the existence of globally bounded solutions.

The adaption of the named existence theorems on our system (1) requires – if possible at all – severe modifications on a technical mathematical level. However, by proving the longterm existence, we aim to develop a deeper understanding of the infection application. Therefore, we present a step by step proof and accompany it by biological and medical applications.

Now, we show the existence of solutions and their boundedness in $L^\infty(\Omega)$, which allows a point-wise estimation of the maximal virus concentration.
and T cell amounts. The section has the following structure. First, the existence of a weak solution for a small time span \([0, T]\) is shown. We discuss some basic properties of such solutions like non-negativity of \(u\) and \(v\) and boundedness of \(u\). These properties are important for modeling purposes as negative values are not interpretable in the context of densities of virus and T cells.

In Section 3.2, the boundedness of \(v\) in \(L^1(\Omega)\) is shown. This result shows a boundedness of the total amount of virus and T cells at a certain time. Afterwards and building up on this result, the boundedness of the norm \(\|v\|_{L^\infty(\Omega)}\) is proven.

Finally, in Section 3.3 the boundedness of the norm \(\|v\|_{L^\infty(\Omega)}\) is shown and using it, the global longterm existence of weak solutions of Eq. (1) is shown.

3.1. Properties of the weak solution

Starting with the definition of a weak solution, the existence of a weak solution of Eq. (1) for a small time span \([0, T]\) is shown.

**Definition 5.** A weak solution of (1) on the time-interval \([0, T]\) is a pair of functions \((u, v)\) with \(u, v \in L^1([0,T]; H^1(\Omega))\) and \(u, v \in L^2([0,T]; \Omega)\) for which

\[
\int_\Omega u_0 \varphi \, dx = \int_\Omega u_0(\varphi - \gamma uv\varphi - \alpha Vu - \nabla \varphi \cdot \nabla v) \, dx,
\]

\[
\int_\Omega v_0 \varphi \, dx = \int_\Omega v_0(\varphi - \eta(1-u)\varphi - \beta \nabla \varphi \cdot \nabla \varphi) \, dx
\]

is fulfilled for all \(\varphi \in H^1(\Omega)\) with \(\varphi = \varphi(x)\) and almost every time \(t \in [0,T]\).

In Ref. 16, Theorem 9.2.2, p. 536 the existence of a unique weak solution of a reaction diffusion system

\[ q = f(q) + \Delta q \quad \text{with} \quad q = \begin{pmatrix} u \\ v \end{pmatrix}, \quad D = \begin{pmatrix} a & 0 \\ 0 & \beta \end{pmatrix} \]

with Dirichlet boundary conditions and a Lipschitz continuous reaction function \(f(q)\) with respect to \(q\) is proven. The first step of the proof shows the existence of a weak solution in case of an externally given function \(h(t) = f(q(t))\) replacing the reaction terms. Additionally, it is shown in this step, that the time derivative of the solution is a \(L^1(\Omega)\)-function as well. Even if the reaction diffusion system (1) has Neumann boundary conditions and the reaction function does not fulfill global Lipschitz conditions with respect to the state variables, this step is adaptable by the following considerations.

Regarding a solution \(q \in C([0,T]; L^1(\Omega, \mathbb{R}^2))\), which is bounded in a suitable chosen time interval \(t \in [0,T]\). Define \(h(t) = f(q(t))\) as a right-hand side for the general parabolic system

\[ q_t - \Delta q = h(t) \quad \text{for} \quad x \in \Omega, 0 < t < T, \]

\[ \nabla q \cdot n = 0 \quad \text{for} \quad x \in \partial \Omega, 0 < t < T, \]

\[ q(0, x) = q_0(x) \quad \text{for} \quad x \in \Omega. \]

Due to the boundedness of \(q\) in the limited time interval and the smoothness of \(f\), the function \(h\) is regular in the sense, that \(h \in L^1((0,T); \Omega)\). In Ref. 16, Theorem 3, p. 378, the existence of a weak solution for systems like in Eq. (7) but with Dirichlet boundary conditions is shown. By replacing the Sobolev space \(H^1(\Omega)\) by \(H^1(\Omega)\) and changing some of the constants, a completely analogous proof assures the existence of a weak solution in case of homogeneous Neumann boundary conditions.

**Theorem 6.** Let \(u_0, v_0 \in L^\infty(\Omega)\) and \(T > 0\) such that \(u(t), v(t) \in L^\infty(\Omega)\) for \(t \in [0,T]\). Then, there exists a weak solution \((u(t), v(t))\) in \(L^1((0,T); \Omega)\) for \(u_0(x), v_0(x) \in L^1((0,T); \Omega)\) with \(u(t), v(t) \in L^1((0,T); \Omega)\) of system (1).

The proof follows Ref. 16, Theorem 9.2.2, p. 536 and Ref. 16, Theorem 3, p. 378 with the mentioned adaptions of the boundary conditions.

The interpretation of the theorem fits with the observation in nature: In a finite time, there cannot be an infinite amount of virus or T cells at a single point. The maximal amount of virus and T cells is bounded for a time interval \([0, t]\).

The weak solution \((u, v)\) of Eq. (1) fulfills some basic properties.

**Lemma 7 (Non-negativity).** If \(u_0(x) \geq 0\) and \(v_0(x) \geq 0\) for all \(x \in \Omega\), then \(u(t, x) \geq 0\) and \(v(t, x) \geq 0\) yield for all \(t \in (0,T)\) and all \(x \in \Omega\).

**Proof.** Regard the point \(x_{\min} \in \Omega\), where one state variable has its minimal value. If at one time \(t\) the minimum \(\min_{x \in \Omega}(u(t, x), v(t, x)) = 0\) touches the lower bound of the positive domain, then Eq. (1) provides that the reaction term

\[ u(t, x_{\min})u(t, x_{\min}) - \gamma(t, x_{\min})v(t, x_{\min}) = 0 \]

vanishes at the point \(x_{\min}\) of the minimum. At the same time, \(u(t, x_{\min}) \geq 0\) at this point. Thus, \(u\) cannot pass zeros, and stays non-negative.

Similarly, if \(\min_{x \in \Omega}(v(t, x)) = v(t, x_{\min}) = 0\) then

\[ f(u)(x_{\min}) \geq 0, \quad \eta(1 - u(t, x_{\min}))v(t, x_{\min}) = 0 \quad \text{and} \quad u(t, x_{\min}) \geq 0, \quad \text{and} \quad v(t, x_{\min}) \text{ stays non-negative as long as it exists, too}. \]

Consequently, the model reflects the fact that the amount of virus \(u\) and of T cells \(v\) are non-negative.

Next, we prove that the amount of virus is bounded point-wise.

**Lemma 8 (Boundedness of u).** If \(u_0(x) \leq 1\) for all \(x \in \Omega\), then the (weak) solution \(u(t, x)\) is bounded by \(u(t, x) \leq 1\) for all \(t \in (0,T)\) and \(x \in \Omega\).

**Proof.** Regard the maximum value \(\max_{x \in \Omega} u(t, x) = u(t, x_{\max})\). If this maximum is equal to 1, then the growth term \(u(t, x)\) vanishes at \(x_{\max}\), see Remark 2, the predator term \(-uv\) is not positive, and the diffusion term \(u\Delta u\) is not positive, too, cf. proof of Lemma 7. Consequently, the maximum \(u(t, x)\) cannot grow above 1.

According to Remark 3, the model in Eq. (1) is not suitable for values of \(u\) larger than 1. Initial conditions with \(u_0 > 1\) does not affect the boundedness of \(u\) by 1.

**Corollary 9.** If \(u_0(x) > 1\) for some \(x \in \Omega\), there exists a time \(t_1\) with \(u(t, x) \leq 1\) for all \(t > t_1\).

**Proof.** Again, we regard the maximum value \(\max_{x \in \Omega} u(t, x) = u(t, x_{\max})\). If it is larger than 1, the logistic growth \(u(t, x)\) is strictly negative at the point \(x_{\max}\). Since \(u\) has its maximum at \(x_{\max}\), the diffusion term fulfills \(\alpha \Delta u \leq 0\). At the same time, \(v\) is increasing, so that the predator term \(-uv\) is larger than 0, and the maximum value \(u(t, x)\) passes the value 1 with a non-zero time derivative at a finite time instant \(t_1\).

The proof shows that \(u\) does not tend to 1, but rather passes 1. That means that the virus decays under its capacity, whenever an active immune response exists. We formulate this observation in a next corollary saying that \(u\) becomes smaller than 1 together with a non-vanishing \(v\) on some sub-domain of \(\Omega\).

**Corollary 10.** All bounded and non-vanishing initial values allow to find a time instant \(t_2\) for which \(u(t_2, x) \leq 1\) holds true for all \(x \in \Omega\) and \(U(t_2) < (1 - \epsilon)|\Omega|\) yields for \(\epsilon > 0\). At the same time, \(v\) is not vanishing at points \(x \in \Omega\) where \(u\) is smaller than 1.

**Proof.** If the function \(u(t, \cdot)\) is not identical to \(\max_{x \in \Omega} u(t, x)\), then Corollary 9 provides \(t_1\) with \(u(t_1, x) \leq 1\) for all \(t > t_1\). Due to the position-dependency of \(u\), there exists \(\epsilon > 0\) with \(U(t_1) < (1 - \epsilon)|\Omega|\).
For space-independent functions $u(t_1, x) = \max_{x \in \Omega} u(t_1, x)$, only the situation

$$u(t_1, x) = \max_{x \in \Omega} u(t_1, x) = 1$$

is interesting. If $u(t_1, x) > 1$ for all $x \in \Omega$, the reaction terms lead to a decay with $u(t_2, x) \leq 1$ due to Corollary 9. Let $(t_1, x) = \text{argmax}_{x \in \Omega} u(t_1, x) = 1$.

Since the inflow $j[u]$ is positive in both cases, $v$ increases, and the predator term $-\mu w$ is strictly negative in $\theta$ for all $t > t_1$. Therefore, the assertions are fulfilled for every $t_2 > t_1$ with sufficiently small $t_2 - t_1$.

In the following, we assume initial conditions $u(0, x) \leq 1$ for all $x \in \Omega$. As shown in Corollaries 9 and 10 and according to the formulation of system (1), this is not a restriction.

We interpret Corollary 10 in the light of application. Even if there would be a higher amount of virus than the upper limit allows, the additional virus vanish by a negative growth term and spread out by diffusion. The negative growth can be interpreted as a decay due to a limited number of free liver cells where the virus can attach.

**Corollary 11.** Let $(u, v)$ be a solution of system (1) for $t \in (0, T)$ with initial conditions $0 \leq u(0, x) \leq 1$ and $0 \leq v(0, x) \leq v_{\text{max}} < \infty$. Then, the $L^1(\Omega)$-norm $U(t) = \|u(t, \cdot)\|_{L^1(\Omega)}$ is bounded by $U(t) \leq [\Omega]$ for all times $t \in [0, T]$.

**Proof.** Due to Lemma 8, the solution $u(t, x)$ is bounded by 1. Integration of both sides of $u \leq 1$ gives $U(t) \leq [\Omega]$.

With these results, we found a (weak) solution for a time interval $[0, T)$, which is non-negative and at least one component of the solution, namely $u$, is bounded. The increase of the second component $v$ depends on the $L^1(\Omega)$-norm $U$ of $u$. Hence, until now, $v$ could still grow over all bounds.

Consequently, we have to show that the increase of $v$ happens simultaneously to a decrease of $U$, cf. Fig. 1, and that this simultaneity makes $v$ to be bounded in the different norms.

Since we will need it in the next section for showing that blow-ups of the solution of system (1) do not occur, we prove that $u$ is not only bounded by 1 but it is sufficiently remote from 1 after some time. The medical background suggests that a virus density close to 1 provokes an increase of the immune response. Hence, the virus density decreases. This slows down the influx of T cells again, compare the opposite directions of the mechanisms in Fig. 1. The following Lemma 12 will give a very rough estimate for this observation.

But first, we consider the solution $v_{\text{aux}} = v_{\text{aux}}(x)$ of the auxiliary stationary problem

$$-\beta du + \eta v = \chi_0(x) \quad \text{for} \ x \in \Omega,$$

$$u v \cdot n = 0 \quad \text{for} \ x \in \partial \Omega.$$

The function $v_{\text{aux}}(x)$ $\geq 0$ is at least piecewise continuous and not vanishing in the whole domain $\Omega$. Consequently $v_{\text{aux}}$ is continuous, bounded and positive. Since $v_{\text{aux}}(x)$ is positive only in the influx region $\theta$, there is some value $v_{\text{aux}} > 0$ with $v_{\text{aux}}(x) \geq v_{\text{aux}}$ for all $x \in \theta$.

Therewith, we are prepared to prove the announced Lemma.

**Lemma 12.** Let $u, v$ be weak solutions of (1). For all $\phi \geq 0$, there is a $\theta$ with $0 < \theta < 1$ and a time $t_1$ with

$$\int_{\Omega} u^\phi dx \leq \theta \int_{\Omega} v^\phi dx$$

for all $t \geq t_1$.

**Proof.** First, we show that there is at least one $t_1$ for which Eq. (9) is fulfilled.

Assume, there would be no such $t_1$. Then, $u$ must be equal 1 almost everywhere in $\text{supp} \phi \subset \Omega$ for all time $t$. As a solution of Eq. (1), $u$ is continuous with respect to $x$. Consequently, $u$ must be equal 1 in $\theta$ and we get the rough estimate $U(t) \geq [\theta]$. Now, the evolution of $v$ in Eq. (1) reads

$$v_t = j[u] - \eta (1 - u) v + \beta du \geq \delta U(t) v_{\text{aux}}(x) - \eta v + \beta du,$$

and after a transient phase, we get

$$v(t, x) \geq \delta U(t) v_{\text{aux}}(x) \geq \delta \theta v_{\text{aux}}(x)$$

and thus

$$v(t, x) \geq \delta \theta v(t, x) \quad \text{for all} \ x \in \theta.$$

Finally, the first equation in system (1) reads

$$u_t = u v (u - \gamma \theta v) \leq u (w(u) - \gamma \delta U(t) v_{\text{aux}}(x)) + \alpha du.$$

So, Eq. (10) implies $u_t < 0$ for all $x \in \theta$, what contradicts the assumption $u = 1$ in $\theta$. Consequently, there is at least one time instant $t_1$ fulfilling Eq. (9).

If we now assume that $u$ grows again after $t_1$ so that the estimate (9) is hurt for every $\theta < 1$ at some $t_1$, that would mean $u$ gets arbitrarily close to 1 in $\theta$. This is again a contradiction to $u(w(u) - \gamma \delta U(t) v_{\text{aux}}(x) \leq u(u) - \gamma \theta \theta v_{\text{aux}}(x) \leq 0$ at this time instant $t_1$.

In the next steps, we show, that there exists an upper bound for $v$ as well. First, we show, that $v$ is bounded in $L^1(\Omega)$ for $t \in (0, T)$. Next, we expand this property for all times $T > 0$. As an intermediate step, we show $v(t, \cdot) \in L^1(\Omega)$. Finally, by using the stationary solution of another related elliptic equation for a stationary problem, we prove that $v$ is bounded and smooth for all times $t$, $v(t, \cdot) \in L^\infty(\Omega)$.

### 3.2. $L^1(\Omega)$ Bounds

First, we determine a $L^1(\Omega)$ bound for $v$. With Theorem 6 we have a (weak) solution $(u, v)$ with $u(t, \cdot), v(t, \cdot) \in H^1(\Omega)$ for $t \in (0, T)$ with a time $T$.

In this section, we show, that $V$ is not growing to infinity for $t \in (0, T)$.

Therefore, we regard the time derivative of the functional $\Phi = \eta U + \gamma V$ which is a linear combination of the $L^1(\Omega)$-norms of $u$ and $v$. The functional $\Phi$ can be interpreted as a measure of the total harm of the infection, compare.\(^{11}\)

**Theorem 13.** Any pair $(U(0), V(0))$ with $0 \leq U(0) \leq [\Omega]$ and $0 \leq V(0) < \infty$ allows to find a trapezoid

$$\Sigma = \{(U, V) : U \in [0, [\Omega]], V \in [0, V_0 - \frac{\eta}{\gamma} U]\}$$

such that $(U(i), V(i)) \in \Sigma$ for all $i \in [0, T]$, i.e. as long as the solution $(u, v)$ of system (1) exists in $L^1(\Omega)$.

This theorem says that the $L^1(\Omega)$-norm of a solution $(u, v)$ of system (1) stays in a bounded region, namely within the trapezoid $\Sigma$, compare Fig. 5, as long as a weak solution exists.

**Proof.** The time derivative of the functional $\Phi = \eta U + \gamma V$ is with system (1)

$$\Phi_t = \frac{d}{dt} \int_{\Omega} \eta u v dx = \int_{\Omega} \eta u v_t(x) + \gamma v_t(x) dx$$

$$= \int_{\Omega} \eta u v(u - \eta v + \eta \theta v) + \gamma v(u + \eta v - \eta \theta v + \beta du) dx$$

$$= \int_{\Omega} \eta u v(u - \gamma \theta v) + \gamma v(u) dx,$$

where we use the divergence theorem and homogeneous Neumann boundary conditions from Eq. (1).

The solution $u$ is meaningful for values between 0 and 1. Due to Corollary 9, it suffices to regard solutions $u$ bounded by 0 and 1, see
 Lemmas 7 and 8. Consequently, the growth function $v(u)$ in Eq. (2) is smaller than 1 for all $u \in [0, 1]$. We get

$$\Phi_j \leq \eta \int \Omega u(t, x) \, dx + \gamma \int \Omega j[u] \, dx - \eta \int \Omega v(t, x) \, dx.$$  

The inflow term $|u|$ is bounded, see Remark 4, and with Eqs. (3) and (5), we can write

$$\Phi_j \leq \eta \int \Omega (t) + \gamma \delta U(t) - \eta \int \Omega V(t) = (\eta + \gamma \delta) U(t) - \eta \int \Omega V(t),$$  

(11)

which is not positive for $U \leq |\Omega|$ and $V \geq \frac{\gamma \delta}{\eta} |\Omega|$. The derivative is non-positive for

$$\Phi \geq \eta |\Omega| + \gamma \delta \int \Omega = \left( \eta + 1 + \gamma \frac{\delta}{\eta} \right) |\Omega| = \gamma V_{up}$$  

and $U \leq |\Omega|$.

Since $U$ stays lower than $|\Omega|$, compare Corollary 11, $\Phi \geq \gamma V_{up}$ implies $\Phi_j \leq 0$, see Eq. (11). In particular, $\Phi$ cannot pass $\gamma V_{up}$ when it is once lower than $\gamma V_{up}$ with $U \leq |\Omega|$. Consequently, the $L^1(\Omega)$-norm of system (1) with $u(t, x)$ stays in $\Sigma$ when it starts in $\Sigma$.

If now $\Phi(t) = 0$, $\Phi(t)$ is smaller than $\gamma V_{up}$ for all admissible $t$. If otherwise $\Phi(0) \geq \gamma V_{up}$, we have shown that $\Phi$ decreases until $\Phi(t)$ is smaller than $\gamma V_{up}$.

Finally, $\Phi(t) \leq \max(\Phi(0), \gamma V_{up})$ is $\gamma V_{up}$ for all admissible $t$ and all initial values allow to construct a suitable $\Sigma$ where the solution stays in $\square$.

In Fig. 5, the trapezoid $\Sigma$ is shown in the phase space of $(U, V)$. The arrows show the direction of the dynamics given by the reaction term in system (1). The arrows of the dynamics point inside $\Sigma$ or at least not to the exterior, especially at the upper bound of $V$.

Remark 14. Since $|\Omega| > 0$, the $L^1(\Omega)$-norm $V(t)$ of $v$ is bounded by

$$V_{up} = \frac{1}{\gamma} \left( \eta + \frac{\gamma \delta}{\eta} \right) |\Omega|$$

(12)

for all $t \in (0, T)$.

That means that the summed strength of the immune response is bounded in a bounded time interval.

Remark 15. Let $\Sigma$ be the trapezoid in the phase space $(U, V)$, which is bounded by $U = 0, V = 0, U = |\Omega|$ and $\eta U + \gamma V \leq \gamma V_{up}$. If the $L^1(\Omega)$-norms $U(0) = \|u(0)\|_{L^1(\Omega)}$ and $V(0) = \|v(0)\|_{L^1(\Omega)}$ are inside $\Sigma$, then the linear combination $\eta U(t) + \gamma V(t)$ with $U(t)$ and $V(t)$ as the $L^1(\Omega)$-norms of the weak solutions $u(x, t), v(x, t)$ of system (1) with $t \in (0, T)$ is inside $\Sigma$ as well.

The $L^1(\Omega)$-norms $U(T - \epsilon)$ and $V(T - \epsilon)$ are for any $\epsilon > 0$ inside of $\Sigma$. $\Sigma$ depends only on the initial values, but it is independent of the time $t$ and the solutions $u$ and $v$ themselves. Consequently, the $L^1(\Omega)$-norms $U(T)$ and $V(T)$ are inside of $\Sigma$ as well. The values $u(x, t)$ and $v(x, T)$ can be seen as new initial data of system (1). By induction, the $L^1(\Omega)$-norms $U(t)$ and $V(t)$ are inside of $\Sigma$ for every $t > 0$ and the $L^1(\Omega)$-norm $V(t)$ of $v(t, x)$ is bounded by $V_{up}$ in Eq. (12) for all time $t > 0$.

This results shows the boundedness of the total amount of T cells at any time, not only for a limited interval.

The amount of T cells $v$ is not only bounded in the sense of $L^1(\Omega)$ but also in the sense of $L^2(\Omega)$. This can be shown by regarding the time derivative of the functional

$$\Psi(t) = \frac{1}{2} \|v(t, x)\|_{L^2(\Omega)}^2 = \frac{1}{2} \int \Omega v^2 \, dx.$$  

Theorem 16. Let $(u, v)$ be a solution of system (1). Then, the $L^2(\Omega)$-norm of $v$ is bounded for all $t > 0$. 

Proof. The time derivative of the functional $\Psi$ is

$$\Psi' = \frac{d}{dt} \int \frac{V^2}{2} \, dx = \int \frac{\partial}{\partial t} v \cdot v \, dx = \int \partial_t v \cdot v \, dx = \int v \cdot \partial_t v \, dx = -\int (\gamma V_{up}) \cdot v \, dx.$$

Using Green’s first identity and the zero-flux boundary conditions, we get

$$\beta \int \Omega v \, dx = \beta \int \partial_t v \cdot n \, ds - \beta = \int \partial_t v \cdot v \, dx = \beta \int \Omega v \cdot v \, dx \leq 0.$$

Further, Remark 4 provides an estimate for the integral of $j[u]$, which is

$$\Psi_j \leq \int \Omega v \cdot j(u) + \eta (u - 1) \, dx \leq \delta \chi_{\max} U \int \Omega v \, dx = \eta \chi \int \Omega (u - 1)^2 \, dx,$$

with $\chi_{\max} = \max_{x \in \Omega} \chi(x)$ according to Eq. (4). Now, Remark 14 assures

$$\Psi_j \leq \delta \chi_{\max} |\Omega| \gamma V_{up} + \eta \int \Omega (u - 1)^2 \, dx = M - \eta \chi \int \Omega v \, dx,$$

with the constant $M = \delta \chi_{\max} |\Omega| \gamma V_{up}$ and the weighted mean value $\xi(t)$ defined by

$$\int \Omega (1 - u)^2 \, dx = \xi(t) \int \Omega v^2 \, dx.$$  

(13)

The mean value $\xi(t)$ fulfills $0 < 1 - \theta \leq \xi(t)$ because of Lemma 12. Finally, the functional $\Psi$ obeys the linear differential inequality

$$\Psi' = \frac{d}{dt} \int \frac{1}{2} \int \Omega v^2 \, dx - M = -\eta \chi \int \Omega v \, dx = M - 2\eta \chi \Psi(t) \Psi'$$  

(14)

with a positive decay rate $2\eta \chi$ which stays remote from 0. Eq. (14) is a first order differential inequality, compare.25 and $\Psi(t)$ is bounded by the solution of the linear first order differential equation

$$y' = M - 2\eta \xi(t)$$

with $\xi(t) \geq 1 - \theta > 0$.

Then, the largest possible accumulation point of $\Psi(t)$ is $\frac{M}{2\eta (1 - \theta)}$, and the functional $\Psi$ is bounded by $M$ after a transient phase. Later in Section 4, we will use the estimate

$$\Psi(t) \leq \Psi(0) e^{-2\eta \xi(t)} + M \leq \Psi(0) e^{-2\eta \xi(t)} \Psi(t)$$

(15)

for showing numerically the precision of the estimates.

Theorems 13 and 16 show, that the $L^1(\Omega)$- and the $L^2(\Omega)$-norms of $v$ are not only bounded for a bounded interval $(0, T)$ but for all time $t > 0$. So in these norms, the solution is not blowing up.

3.3. $L^\infty(\Omega)$ Bounds and global existence

In this section, we show the boundedness of $v$ in $L^\infty(\Omega)$ for all $t > 0$. With the boundedness of $v(t, x)$, the existence of a solution $(u, v)^T$ with finite values is shown for all $t > 0$. 

![Fig. 5. Trapezoid in the phase space $(U, V)$ of the $L^1(\Omega)$-norms of a solution $(u, v)$ of system (1). The scaled vector field shows the dynamics of the reaction terms in Eq. (1).](image-url)
We will prove, that there exists a value \( v_{\text{max}} \) with \( v(t,x) \leq v_{\text{max}} \) for all \( x \in \Omega \) and all \( t \in [0, \infty) \). For this purpose, a stationary problem is defined. Let \( v^* = v^*(x) \) be a solution of
\[
-\beta \Delta v = \chi_\delta(x) - \frac{1}{|\Omega|} \quad \text{for } x \in \Omega,
\]
where (16) fulfills the solvability condition because the forces are equalized, see Eq. (4) and
\[
\int_\Omega \chi_\delta(x) - \frac{1}{|\Omega|} + \beta \Delta v \, dx = \int_\Omega \chi_\delta(x) \, dx - \frac{1}{|\Omega|} \int_\Omega dx + \beta \int_{\partial \Omega} v \cdot n \, ds = 1 - \frac{|\Omega|}{|\Omega|} + 0 = 0.
\]

Remark 17. Since the right-hand side \( \chi_\delta(x) - \frac{1}{|\Omega|} \) in Eq. (16) is a bounded piecewise continuous function and thus in \( L^2(\Omega) \subset H^{-1}(\Omega) \), the existence of a weak solution \( v^* \in H^1(\Omega) \) is ensured. 

Remark 18. The solution \( v^* \) of Eq. (16) has a free additive constant as always in pure Neumann problems. In the following, we fix just one \( v^* \) with \( \|v^*(x)\|_{L^2(\Omega)} = 0 \).

Now, we will show that the population \( v = v(t,x) \) in Eq. (1) does not grow to infinity. Even having already estimates for its \( L^2(\Omega) \)-norm, cf. Theorem 13, and for its \( L^2(\Omega) \)-norm, cf. Theorem 16, it is not trivial to give a pointwise bound. Before we will do that in the later Theorem 23, we collect some auxiliary results about solutions of partial differential equations with homogeneous Neumann boundary conditions.

Lemma 19. Let \( v = v(t,x) \) be the solution of
\[
\begin{align*}
v_{\gamma} &= \beta \Delta v - a(t,x)v + f(t,x) \quad \text{for } x \in \Omega, t > 0, \\
v \cdot n &= 0 \quad \text{for } x \in \partial \Omega, t > 0, \\
v(0,x) &= 0 \quad \text{for } x \in \Omega,
\end{align*}
\]
with \( a(t,x) \geq 0 \) and \( |f(t,x)| \leq C_f \) for all \( x \in \Omega, t > 0 \). Then \( |\Delta v| \) is bounded by a constant \( C_g \in \mathbb{R} \) for all \( x \in \Omega \) and \( t > 0 \).

Remark 20. Eq. (17) is a heat conduction equation with the additional level term \( -\beta \Delta v \) and homogeneous Neumann boundary conditions, i.e. no heat flux over the boundary, and a bounded heat source \( f \). Hence, a physical point of view implies immediately a bounded curvature of \( v \).

The mathematical argumentation starts with the Green's function \( G = G(t,x,y) \) of Eq. (17), which is dominated by the singularity of the standard heat equation. Due to the Neumann boundary condition, there are no additional term at the boundary. The Laplacian \( \Delta G(x) \) is the convolution of \( \Delta G \) with the bounded function \( f \). This convolution can be estimated by a sum of spatial integrals over small domains and afterwards by time integration leading to terms in the Gauss' error function. Due to its technical effort, we omit the argumentation of the physically proven assertion of Lemma 19. By the way, another possible argumentation uses a discretization of the Eq. (17), where the eigenvalues and eigenvectors of the discretized differential operator \(-\Delta + a\) can be estimated in a similar technical argumentation. Then the limit case of a temporal step size tending to zero provides the assertion of Lemma 19 for every spatial discretization, and since \( f \) is bounded also the limit situation of a vanishing grid size.

Lemma 21. Let \( z : \Omega \to \mathbb{R} \) be a sufficiently smooth function with homogeneous Neumann boundary conditions. The function \( z \) fulfills \( |\Delta z(x)| \leq C_f \) for all \( x \in \Omega \) and \( \|z\|_{L^2(\Omega)} \leq C_f \). Then its values \( z(x) \) are bounded by some \( z_{\text{max}} < \infty \) with \( |z(x)| \leq z_{\text{max}} \) for all \( x \in \Omega \).

Proof. Such a function \( z \) solves a boundary value problem
\[
-\Delta z = \rho(x) \quad \text{for } x \in \Omega, \\
v \cdot n = 0 \quad \text{for } x \in \partial \Omega
\]
with a source term with \( |\rho(x)| \leq C_f \) for all \( x \in \Omega \). We choose a Green's function \( G(x,y) \) with \( G(x,y) \geq 0 \) for all \( x, y \in \Omega \).

With an additive constant \( C_g \), we have
\[
z(x) = \int_\Omega G(x,y) \rho(y) \, dy + C_g \quad \text{and} \quad |z(x) - C_g| \leq C_f \int_\Omega G(x,y) \, dy = \Phi(x)
\]
with the smooth and bounded potential \( \Phi \) for a constant source term.

There is a value \( C_{g_{\text{max}}} < \infty \) so that the condition \( \| \Phi \|_{L^2(\Omega)} \leq C_g \) is not fulfilled for any \( C_g \geq C_{g_{\text{max}}} \). Consequently, we get
\[
|z(x)| \leq C_{g_{\text{max}}} + \max_{x \in \Omega} \Phi(x) = z_{\text{max}} < \infty.
\]

This lemma shows the boundedness of a stationary problem which displays in parts the inflow of T cells in a certain region. In the next step, we use this result for showing the existence of an upper bound for \( v \) in a time-dependent setting which still abstracts from the coupled reaction diffusion system in (1).
for the standard diffusion problem in Eq. (21) is a function in \(t \) and \(x\) for fixed \(u\) and thus fixed \(U, U'\) and \(v^*\). Eq. (21) is completed with homogeneous Neumann boundary conditions for \(\tilde{v}\).

That means, we regard \(u\) to be given and investigate Eq. (21) as a diffusion problem for \(\tilde{v} = \tilde{v}(t, x)\) with the exogenous influence \(f = f(t, x)\). Since

\[
U'(t) = \int_\Omega u(t, x) \Delta x - \gamma \int_\Omega \tilde{v}(t, x) \Delta x
\]

and \(|\tilde{u}| \leq v\) with \(v \in \mathbb{R}_+\), cf. Remark 2, we see

\[
|U'(t)| \leq vU(t) + \gamma V(t)
\]

for all \(t\). Together with the boundedness of \(U\) and \(V\) for all \(t\), the exogenous influence \(f\) in Eq. (22) is bounded by some constant \(C_f \geq |f(t, x)|\) for all \(t > 0\) and \(x \in \Omega\).

Now with Eq. (21), we find

\[
V_{\text{up}} \geq \int_\Omega \tilde{v}(t, x) \Delta x = \delta U(t) \int_\Omega v^*(x) \Delta x + \int_\Omega \delta(t, x) \Delta x = \int_\Omega \delta(t, x) \Delta x. \tag{23}
\]

Now, Eq. (21) fulfills all conditions of Lemma 22, namely

\[
a(t, x) = \eta(1 - u) \geq 0 \quad \text{and} \quad |f(t, x)| \leq C_f
\]

for a heat equation with homogeneous Neumann boundary conditions. Consequently, there exists a maximal value \(\tilde{v}_{\text{max}} \geq \tilde{v}(t, x)\) for all \(t > 0\) and all \(x \in \Omega\), and we get

\[
\tilde{v}(t, x) \leq \delta[\Omega] \max v^*(x) + \tilde{v}_{\text{max}} = \tilde{v}_{\text{max}} \in \mathbb{R}. \quad \square
\]

**Remark 24.** The solution \(v\) of Eq. (1) is bounded by a finite value \(v_{\text{max}}\). Finally, the solution \(v(t, x)\) is a \(L^\infty(\Omega)\)-function.

In this section, we have proven the boundedness of the solution of Eq. (1). While the boundedness of \(u\) was a result of the used growth function and therefore allows interpretation as a concentration, the boundedness of \(v\) was not obvious. Using the oppositely acting mechanisms in the reaction functions and the boundedness of \(u\), we first showed the boundedness of \(v\) in \(L^1(\Omega)\).

We provided a bounded estimate for the \(L^2(\Omega)\)-norm of \(v\) by using the mean-value theorem of integration and the boundedness of \(\|v(t, \cdot)\|_{L^2(\Omega)}\). For proving the boundedness of \(v\) in \(L^\infty(\Omega)\), we separated \(v(t, \cdot) = \delta v^*(x) U(t) + \delta(t, \cdot)\) into different functions. One component, \(v^*\), of the functions was the solution of a stationary problem covering the space-dependent function modeling the inflow area of the liver structure. By showing the boundedness of all components of \(v\), we proved in Theorem 23 that \(v\) has a finite maximal value.

Applied to the modeling of liver infections, the result of Remark 24 says that the amount of T cells is bounded by a finite value. The T cells attack the virus by triggering the programmed cell death. This leads to inflammation in the liver tissue and can cause secondary diseases like cancer. Besides, a too high amount of T cells might cause a sepsis.

**Remark 24** gives the fact, that the immune reaction is bounded but does not give a finite value. Therefore, **Remark 24** justifies the use of model in the sense, that the immune reaction remains bounded for all time. It is a first step towards quantitative and finer estimates for \(v_{\text{max}}\) which contain information about the occurrence of sepsis.

Therefore, the next section provides numerical evaluations on the \(L^1(\Omega)\) and \(L^2(\Omega)\) estimates.

### 4. Numerical evaluation of the estimates

Oftentimes, estimates used in analytical results are rather rough. In this section, we show numerical simulations of the estimates and the exact value. The simulations are based on the simulations in Figs. 2 and 3. Those use a semi-discretization by finite differences for the space and Runge-Kutta based methods for the time. As the finite differences discretization of the Laplacian might lead to a stiff ordinary differential equation problem, a standard solver for stiff ordinary differential equations is used in Matlab. The numerical evaluation of the estimates uses finite sums instead of the integrals in accordance with the space and time resolution used for solving the partial differential equations.

First, we evaluate the estimation of the domain \(\Sigma\) as maximal \(L^1(\Omega)\)-norms. In Fig. 6, the trajectories of the two solutions from Figs. 2 and 3 provide the total amount \(U\) and \(V\). They are compared to the estimated \(\Sigma\) following Theorem 13.

Fig. 6 shows that the upper bound of \(\eta U + \gamma V \leq \gamma V_{\text{up}}\) is a rather rough estimate for the \(L^1(\Omega)\)-norms of the solutions. In the numerical simulations in Figs. 2 and 3, which are as well used in Fig. 6, the initial conditions are \(u(0, x) = 1\) and \(v(0, x) = 0\). A solution with larger initial conditions \(V(0)\) would reach closer to the upper bound given by \(\Phi = \gamma V_{\text{up}}\). As shown in Theorem 13, the \(L^1(\Omega)\)-norms of every solution with \((U(0), V(0)) \in \Sigma\) stay in \(\Sigma\).

**Theorem 16** gives an estimate for the \(L^2(\Omega)\)-norm of \(v\). By using \(\xi(t)\) as solution of Eq. (13) and directly Eq. (14) we get the approximation

\[
\Psi(t) = \frac{1}{2} \|\xi(t, \cdot)\|_{L^2(\Omega)}^2 \leq c < 2 >^2 \|\xi(t, \cdot)\|_{L^2(\Omega)}^2 \leq c \int_0^t c \|\xi(t, \cdot)\|_{L^2(\Omega)}^2 \, dt + \Psi(0)
\]

with a parameter-dependent constant \(c = \delta[\Omega] \max \gamma[\Omega] V_{\text{up}}\). The inequality \(0 < \xi(t) \leq 1\) yields according to Lemma 12.

In the cases of the two regarded simulations in Figs. 2 and 3 of Eq. (1), the initial value \(\Psi(0)\) is zero, because \(\xi(0, x) = 0\) for all \(x \in \Omega\). Therefore, we compare the \(\Psi(t)\) to the functional

\[
E(\xi(t)) = c \|\xi(t, \cdot)\|_{L^2(\Omega)}^2 \delta[\Omega] V_{\text{up}} \int_0^t c \|\xi(t, \cdot)\|_{L^2(\Omega)}^2 \, dt.
\]

In both cases in Fig. 7, the estimation \(E\) is rather large compared to the functional \(\Psi(t)\). Nevertheless, the estimate is a good approximation of scale of the maximal value of \(\Psi\).

The numerical simulations show that the used estimates are rather loose even if they were sufficient for gaining the analytical existence results. The estimations exceed the values of the numerical solutions. In context of liver infections, the estimations can be regarded as worst case scenario. A medical treatment in the light of the worst case scenario might lead to a longer lasting infection course but decrease the risk of a sepsis.

### 5. Conclusions

With the aim to modeling the dynamics of liver infections as an interplay between virus and T cells, a reaction diffusion system was presented in Ref. 9. A non-local term in the reaction function describes the inflow of T cells depending on the total virus amount in the domain. The model abstracts from the cell scale with many unknown mechanisms to a mesoscopic length scale. On this scale, the mathematical description contains a space-dependent term which leads to a new problem concerning the analysis of reaction diffusion equations. Additionally, the reaction terms contain oppositely acting mechanisms resulting in a feedback loop for the increase of T cells. The longtime behavior of the solutions of the model depends on the used parameters. On the one hand, there are solutions with a strong tendency towards zero. These solutions are interpreted as healing infection courses. On the other hand, there are solutions with a tendency towards a spatial inhomogeneous steady state. These solutions are interpreted as chronic infection courses with some virus persisting in the liver. One quantity of the model gives the amount of T cells which trigger the cell death of infected liver cells. This effect leads to further mechanisms known as inflammation of the tissue. Inflammation might be critical as the danger of sepsis is connected to strong inflammation. Therefore, it is important to know about upper bounds for the modeled quantities.

The aim of this paper was to prove the existence of bounded solutions for all time and to provide a proof with interpretable intermediate steps. First, some results on the existence theory for parabolic partial differential equations are summarized. As the focus lies on the
interpretation of the gained results are therefore, the steps for proving longtime existence and boundedness of solutions are explained.

We started with a local existence theorem and some properties of a weak solution. Then, we showed the boundedness of the solution in the $L^1(\Omega)$-norm and in the $L^2(\Omega)$-norm. Both results are based on the interplay of the two species in the population dynamics model and the oppositely acting mechanisms of growth and decay. While the local boundedness of $u$ was a result of scaling and a limited growth, the dynamics of the second state variable are more complex and depending locally on the integral of $u$. For proving boundedness of the second variable, the pure boundedness of the first variable is not enough: We need working predator–prey mechanisms leading to a decay of the first variable away from the maximal capacity. Additionally, the non-negativity of both state variable is shown. This is important for interpreting the state variables biologically. The $L^1$-bound for $v$ gives a maximal total amount of T cells in the regarded domain and therefore information on the maximal inflammation. We then defined a stationary problem for showing the boundedness of the solution in the $L^\infty(\Omega)$-norm. This approach was based on a decomposition of the solution for the T cells into a stationary and a time-dependent problem.

In Section 4, we evaluated the sharpness of the used estimates in the proofs. The numerical simulations show that the estimates are rather loose for the regarded cases. Nevertheless, the estimations provide insight:

In the light of the application, modeling liver infections, the estimation can be seen as a worst case scenario. The boundedness of the $L^\infty(\Omega)$-norm of $v$ is a feature uprating the model for liver infections.

A further investigation could be the improving of the used estimated such that the difference between estimate and real value of the functionals becomes smaller. Another possible extension is the application of the estimates for a wider class of integro-partial differential equations. The gained estimations can be used for planning treatment more effectively and with less harm for the patients. First results on modeling treatment plans by hands of Eq. (1) can be found in Ref. 27.

Financial disclosure

None reported.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Acknowledgment

We acknowledge support by the Open Access Publication Funds of Technische Universität Braunschweig.
References

1. Thomas E, Liang T. Experimental models of hepatitis B and C – new insights and progress. Nat Rev Gastroenterol Hepatol. 2016;13(6):362–374. http://dx.doi.org/10.1038/nrgastro.2016.37.

2. Ibragimov A, McNeal C, Ritter L, Walton J. A mathematical model of atherogenesis as an inflammatory response. Math Med Biol. 2005;22(4):305–333. http://dx.doi.org/10.1093/imammb/dqi011.

3. Volpert V. Elliptic Partial Differential Equations. (no. 104). Basel: Birkhäuser; 2014. In: Monographs in Mathematics.

4. Rezounenko A, Karazin V. Viral infection model with diffusion and state-dependent delay: Stability of classical solutions. Discrete Cont Dyn-B. 2018;23(3):1091–1105. http://dx.doi.org/10.3934/dcdsb.2018143.

5. Tadmon C, Foko S, Rendall AD. Global stability analysis of a delay cell-population model of hepatitis B infection with humoral immune response. Dyn Syst. 2021;36(4):537–559. http://dx.doi.org/10.1080/14689367.2021.1940868.

6. Aston P. A new model for the dynamics of hepatitis c infection: Derivation, analysis and implications. Viruses. 2018;10(4). http://dx.doi.org/10.3390/v10040195.

7. Dahari H, Lo A, Ribeiro R, Perelson A. Modeling hepatitis c virus dynamics: Liver regeneration and critical drug efficacy. J Theoret Biol. 2007;247(2):371–381. http://dx.doi.org/10.1016/j.jtbi.2007.03.006.

8. Nangue A, Rendall AD, Tcheugam BK, Simo PSK. Analysis of an initial value problem for an extracellular and intracellular model of hepatitis C virus infection. Int J Biomath. 2022;15(06):2250041. http://dx.doi.org/10.1142/S1793524522500413.

9. Kerl H, Langemann D, Vollrath A. Reaction diffusion equations and the chronification of liver infections. J Theoret Biol. 2007;247(2):371–381. http://dx.doi.org/10.1016/j.jtbi.2007.03.006.

10. Reisch C, Langemann D, Vollrath A. Reaction diffusion equations and the chronification of liver infections. J Theoret Biol. 2007;247(2):371–381. http://dx.doi.org/10.1016/j.jtbi.2007.03.006.

11. Reisch C, Langemann D, Vollrath A. Reaction diffusion equations and the chronification of liver infections. J Theoret Biol. 2007;247(2):371–381. http://dx.doi.org/10.1016/j.jtbi.2007.03.006.

12. Reisch C, Langemann D. Modeling the chronification tendency of liver infections as an evolutionary advantage. B Math Biol. 2019;81(11):4743–4760. http://dx.doi.org/10.1007/s11538-019-05956-y.

13. Reisch C. Reaktions-Diffusions-Gleichungen und Modellfamilien zur Analyse von Entzündungsprozessen. Göttingen: Coviller, TU Braunschweig; 2020.

14. Allee W. Principles of Animal Ecology. Philadelphia: Saunders Co.; 1949. http://dx.doi.org/10.5962/bhl.title.7325.

15. Kanel GC. Pathology of Liver Diseases. Hoboken, NJ: John Wiley & Sons, Inc; 2017.

16. Evans L. Partial Differential Equations. 2nd ed. (no. v. 19). Providence, RI: American Mathematical Society; 2010. In: Graduate Studies in Mathematics.

17. Amann H. Linear and Quasilinear Parabolic Problems: Volume I: Abstract Linear Theory. (no. 69). Basel: Birkhäuser; 1995. Monographs in Mathematics; http://dx.doi.org/10.1007/978-3-0348-9221-6.

18. Lunardi A. Analytic Semigroups and Optimal Regularity in Parabolic Problems. Basel: Birkhäuser; 2012. In: Modern Birkhäuser Classics.

19. Roubíček T. Nonlinear Partial Differential Equations with Applications. second ed. (no. 153). Basel: Birkhäuser; 2013. In: International series of numerical mathematics.

20. Showalter R. Monotone Operators in Banach Space and Nonlinear Partial Differential Equations. (no. v. 49). Providence, RI: American Mathematical Society; 1997. Mathematical Surveys and Monographs.

21. Henry D. Geometric Theory of Semilinear Parabolic Equations. (no. 80). Berlin, Heidelberg: Springer; 1981. In: Lecture Notes in Mathematics.

22. Laamri E, Pierre M. Global existence for reaction-diffusion systems with nonlinear diffusion and control of mass. Ann I H Poincare-AN. 2017;34(3):571–591. http://dx.doi.org/10.1016/j.anihpc.2016.03.002.

23. Rouchon P. Universal bounds for global solutions of a diffusion equation with a nonlocal reaction term. J Differential Equations. 2003;193(1):75–94. http://dx.doi.org/10.1016/S0022-0396(03)00039-1.

24. Anguiano M, Kloeden P, Lorenz T. Asymptotic behaviour of nonlocal reaction-diffusion equations. Nonlinear Anal-Theor. 2010;73(9):3044–3057. http://dx.doi.org/10.1016/j.na.2010.06.071.

25. Walter W. Ordinary Differential Equations. New York: Springer; 1998. http://dx.doi.org/10.1007/978-1-4612-0601-9.

26. Renardy M, Rogers R. An Introduction to Partial Differential Equations. New York: Springer; 2010.

27. Reisch C. Modelling health impacts of hepatitis – model selection and treatment plans. Math Comput Model Dyn Syst. 2022;28(1):28–54. http://dx.doi.org/10.1080/13873954.2021.2020296.