

2D/3D DDFV scheme for anisotropic-heterogeneous elliptic equations, application to a bio-mathematics problem: electrocardiogram simulation.

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Rodolphe TURPAULT

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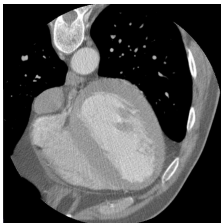
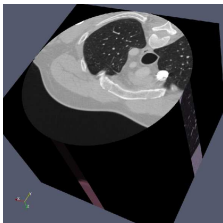
- Charles PIERRE

Outline :

1. Context, motivations.
2. 2D/3D DDFV scheme definition,
for anisotropic/heterogeneous elliptic equations.
3. Numerical analysis.
4. ECG simulation.

Medical data analysis

- Medical images (CT Scan, 3D)

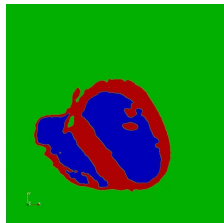
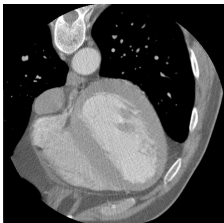


Heart Institute,
University of Ottawa.

Résolution : $1 \times 1 \times 2$ mm.

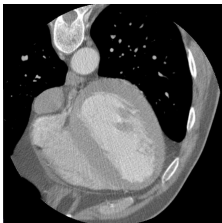
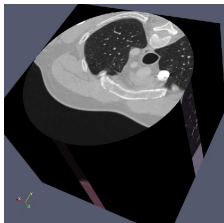
Pixels : 62 500 000

- Segmentation process (Level Set, Mumford-Shah functional)



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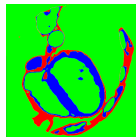
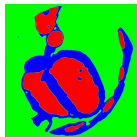
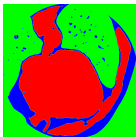
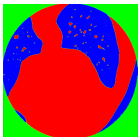
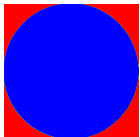


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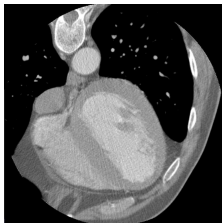
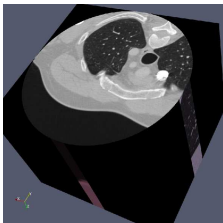
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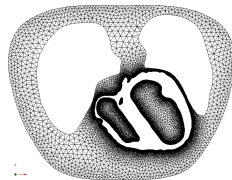
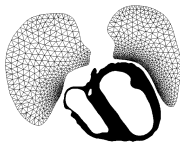
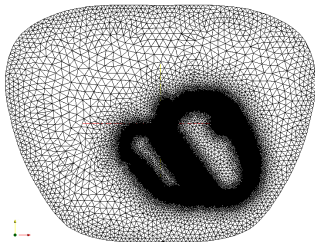


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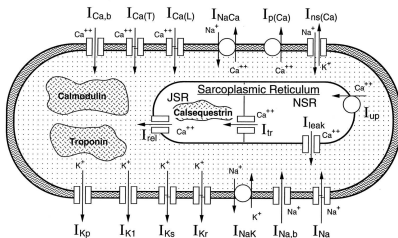
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- Physiologically relevant mesh generation



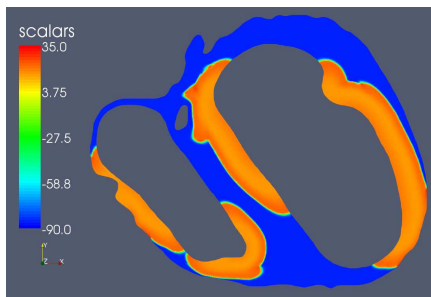
Available models

- Cardiac cells electrical activity
 - EDO system



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- Cardiac tissue electrical activity
 - Electrical potential waves propagation
 - Parabolic reaction diffusion equation
 - Description of a *membrane potential* $V_m(x, t)$ in the heart



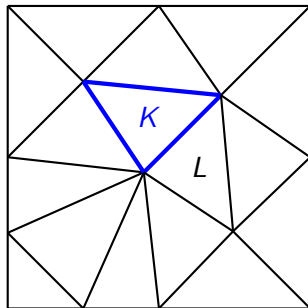
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 - Description of a *membrane potential* $V_m(x, t)$ in the heart
- Torso model
 - Elliptic anisotropic and heterogeneous equation
 - Description of a potential $\phi_T(x, t)$ in the whole torso
 - Body surface potential measurement : **ECG** simulation

$$\operatorname{div}(G \nabla \phi_T) = F(V_m)$$

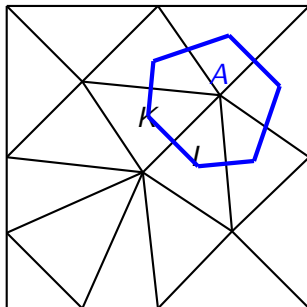
- $G = G(x)$ discontinuous (organs heterogeneity).
- G anisotropic (muscles fibres directions).

DDFV Meshes



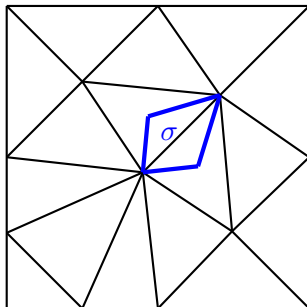
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one scalar unknown at every primal cell centre $K, L, \dots$

DDFV Meshes



- **Primal mesh :**
one scalar unknown at every primal cell centre K , L , ...
- **Dual mesh :**
one scalar unknown at every mesh vertex A (*i.e.* every dual cell).

DDFV Meshes



- **Primal mesh :**
one scalar unknown at every primal cell centre $K, L, \dots$
- **Dual mesh :**
one scalar unknown at every mesh vertex A (*i.e.* every dual cell).
- **Diamond mesh :**
one flux vector associated to every interface σ .

DDFV Scheme principle

Construction of two discrete operators

- **Discrete gradient :** scalar data $\longrightarrow$ vectorial data
- **Discrete divergence :** vectorial data $\longrightarrow$ scalar data

satisfying a duality property, mimiking the *Green formula* in the continuous context.

Property (DDFV scheme 2D-3D)

- *By construction, the discrete gradient and divergence operators fulfill a duality property : discrete Green Formula.*
- *Discretisation of elliptic equations $\implies$ **well posed** symmetric, positive definite system.*
- *Holds for general anisotropic, heterogeneous tensors both in dimension 2 and 3.*

Discrete gradient

Scalar data $(\phi_C)_{C \in \mathcal{C}} \longrightarrow$ continuous piecewise affine function $\phi :$

- $\mathcal{C}$ primal and dual cells set.
- $(\phi_C)_{C \in \mathcal{C}} :$ one data per primal, dual cells
 $\implies$ function ϕ by P^1 interpolation on diamond cells halves.
- **Discrete gradient** : $(\nabla_h \phi_C)_\sigma$ on an interface σ .

$$(\nabla_h \phi_C)_\sigma = \frac{1}{|D(\sigma)|} \int_{D(\sigma)} \nabla \phi \, dx ,$$

$D(\sigma)$ diamond cell associated to σ .

- **Discrete trace** : $(\text{Tr}_h \phi_C)_\sigma$ on boundary interfaces $\sigma \subset \partial\Omega$.

$$(\text{Tr}_h \phi_C)_\sigma = \frac{1}{|\sigma|} \int_\sigma \phi \, dx ,$$

Discrete divergence

Vector data $(\mathbf{p}_\sigma) \longrightarrow$ piecewise constant vector field $\mathbf{p}$ on the diamond cells.

- **Discrete divergence** : $(\operatorname{div}_h \mathbf{p}_\sigma)_C$ on a cell $C \in \mathcal{C}$ (primal or dual).

$$(\operatorname{div}_h \mathbf{p})_C = \frac{1}{|C|} \int_{\partial C} \mathbf{p} \cdot \mathbf{n} \, dx$$

Theorem (discrete Green formula)

Being given a vector data $(\mathbf{p}_\sigma)$ and a scalar data (ϕ_C) :

$$\int_{\Omega} \nabla_h \phi \cdot \mathbf{p} \, dx = - \langle \phi_C, \operatorname{div}_h \mathbf{p} \rangle + \int_{\partial \Omega} \operatorname{Tr}_h \phi \, \mathbf{p} \cdot \mathbf{n} \, ds,$$

$$\langle \phi_C, \operatorname{div}_h \mathbf{p} \rangle = \frac{1}{2} \sum_{C \in \mathcal{C}} (\operatorname{div}_h \mathbf{p})_C \, \phi_C \, |C|.$$

with : $\mathcal{C}$ primal and dual cells set.

Heterogeneous case

$$\operatorname{div}(G\nabla\phi) = f, \quad \text{with :}$$

G discontinuous when crossing some media-interface Γ .

The mesh is supposed to be adapted to Γ :

$$\Gamma \subset \bigcup_{\sigma \text{ mesh interface}} \sigma.$$

Adaptation. When $\sigma \subset \Gamma$:

- Two discrete tensors are associated to σ (one on each side).
 - Two discrete gradient vectors are computed on σ (one on each side).
 - One supplementary normal flux continuity condition is imposed on σ .
- Achieved by adding one auxiliary scalar unknown at the interface center, locally eliminated.

3D case : diamond cells

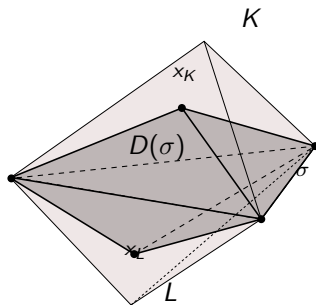


FIG.: 3D diamond cell

3D case : dual cells

$$P(A) = \bigcup_{\substack{\text{interfaces } \sigma, \\ A \in \sigma}} P_{\sigma}(A)$$

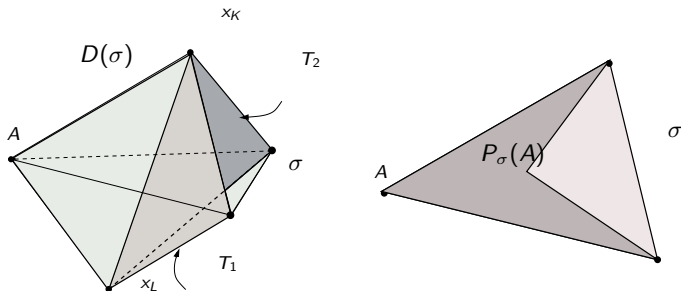
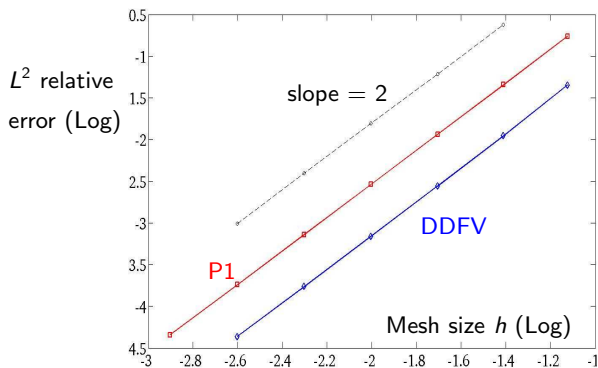


FIG.: 3D dual cells construction

2D convergence analysis

$$\begin{aligned}\Delta u &= f \quad \text{on } \Omega \\ u|_{\partial\Omega} &= 0 \quad \text{on } \partial\Omega\end{aligned}$$



- $\Omega = (-1, 1)^2$
- f trigonometric
- DDFV vs P^1 finite element,
- $177 \rightarrow 650\,000$ points triangular meshes.

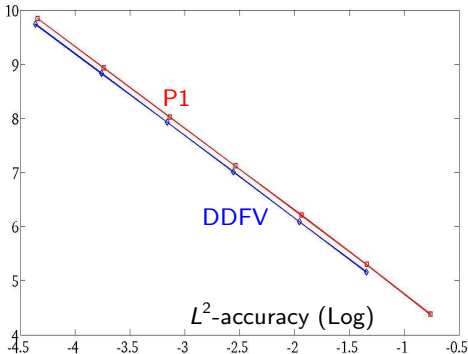
2D DDFV vs P1 : L^2 -norm convergence

2D convergence analysis

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Computational

cost (Log)

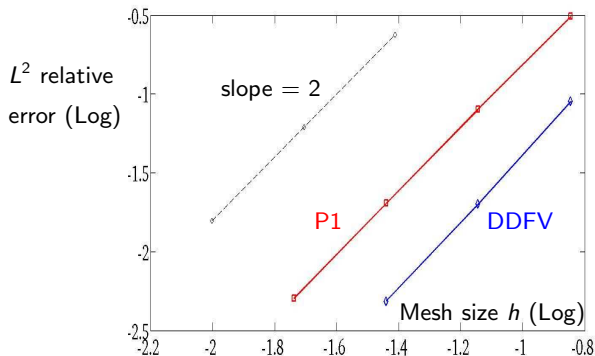


- $\Omega = (-1, 1)^2$
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2D DDFV vs P1 : comparative computational costs

3D convergence analysis

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- $\Omega = (0, 1)^3$
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- 500 $\rightarrow$ 500 000 points tetrahedral meshes.

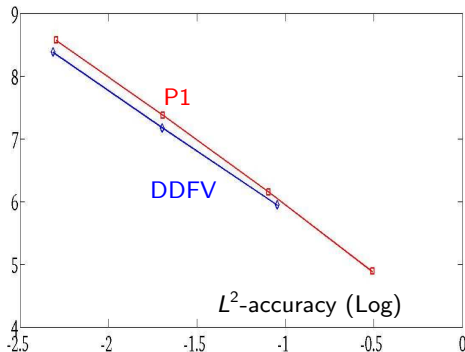
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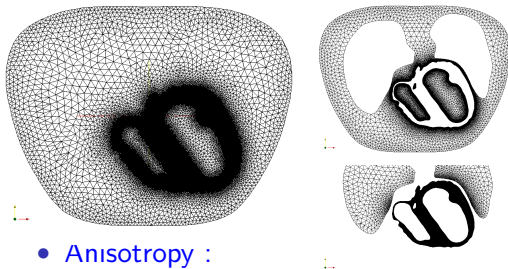


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3D DDFV vs P1 : comparative computational costs

Numerical simulations : settings

- **2D mesh** : from **DistMesh** code (P.-O. Persson, G. Strang).



→ **Heart mesh**

$\simeq$ 500 000 vertices

→ **Entire mesh** (heart+torso)

$\simeq$ 600 000 vertices

- **Anisotropy** :

$$\operatorname{div}(G \nabla \phi_T) = F(V_m) , \quad \text{at each time step.}$$

Ventricles :

$G = G(x)$: anisotropic, depending on muscular fibres directions

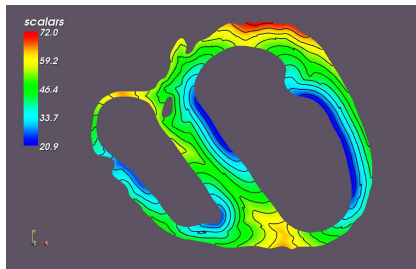
Lungs, ventricle cavity, remaining tissues :

$G = g_o \operatorname{Id}$: homogeneous, heterogeneous

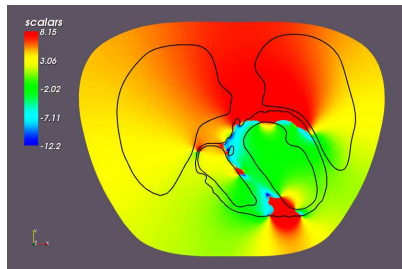
g_o discontinuous at organs boundaries.

ECG simulation : results

Simulation :

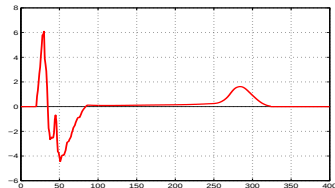
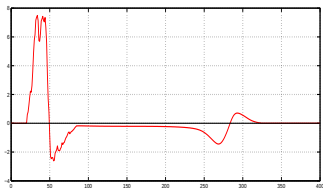


Membrane potential V_m Isochrones.



Torso potential ϕ_T .

Body surface potential recordings (ECG) :



Perspectives

1. Further numerical investigations of the DDFV scheme (behaviour in presence of strong anisotropy/heterogeneity).
2. Theoretical analysis (convergence analysis, discrete Poincaré inequality).
3. Behaviour on non conformal meshes, towards local mesh refinement.