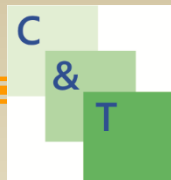




www.gliomark.eu



GLIOMARK GA673737
SME INSTRUMENT PHASE 2

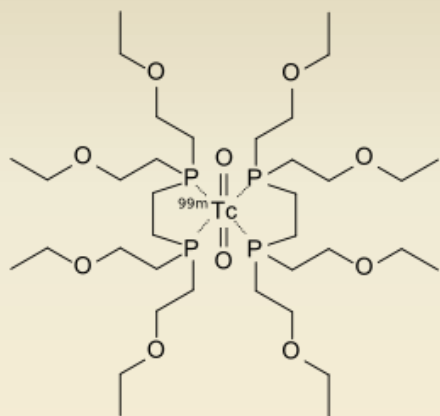
“Validation of blood-brain-barrier permeability as a glioma biomarker by means of the radiotracer ^{99m}Tc -tetrofosmin and single-photon emission computer tomography”

ΕΘΝΙΚΟ ΙΔΡΥΜΑ ΕΡΕΥΝΩΝ, 3 ΝΟΕΜΒΡΙΟΥ 2015

Presented by:

Dr. Alex Strongilos, CEO





99mTc-Tetrofosmin complex

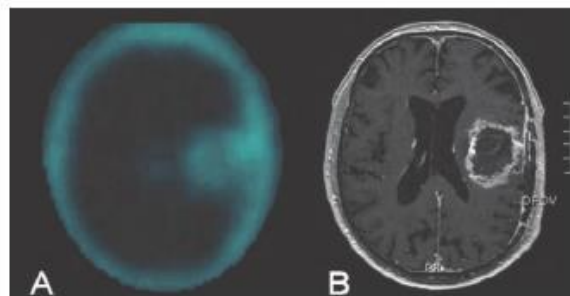


Fig. 4: A. TTF/SPECT: recurrent glioma characterized by intense TTF uptake (L/N: 8.7). B: Corresponding MRI

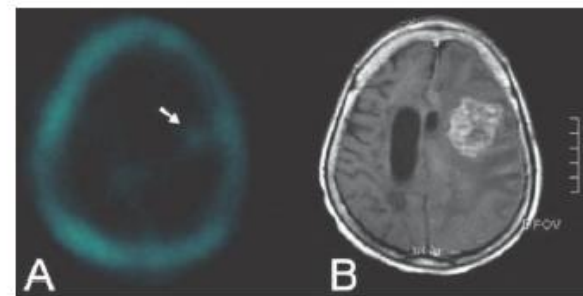


Fig. 5: A. SPECT/TTF: radiation necrosis exhibiting only mildly increased TTF uptake (L/N: 2.9). B: Corresponding MRI

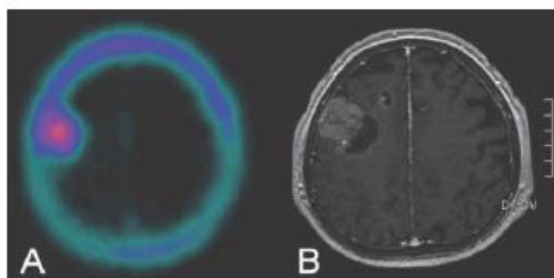


Fig. 6: A. SPECT/TTF: high grade glioma with strongly increased TTF uptake (L/N: 16.0). B: Corresponding MRI image.

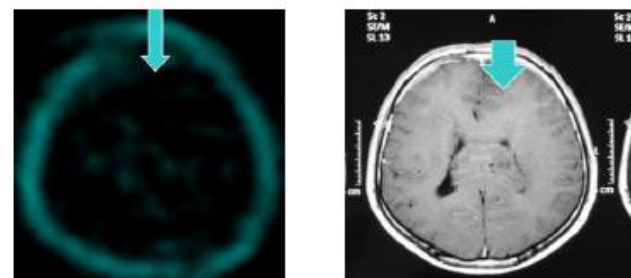


Fig. 7: Left: SPECT/TTF: low grade glioma (L/N: 1.8). Right: Corresponding MRI image.

The value of ^{99m}Tc-tetrofosmin brain SPECT in predicting survival in patients with glioblastoma multiforme. George A Alexiou, Spyridon Tsiouris, Athanasios P Kyritsis, George Fotakopoulos, Anna Goussia, Spyridon Voulgaris, Andreas D Fotopoulos Journal of Nuclear Medicine 11/2010; 51(12):1923-6. · 5.77 Impact Factor

SME INSTRUMENT **The SMEs “Champions League”**

Phase 1

**Concept &
Feasibility
Assessment**

**Input: “Business Plan 1”
(~ 10 pages)**

- Feasibility of concept
- Risk Assessment
- IP regime
- Partner Search
- Design Study

**Lump sum:
≈ 50,000 €
≈ 6 months**

Phase 2

**R&D,
Demonstration,
Market
Replication**

**Input: “Business Plan 2”
+description of activities
Phase 2 (~ 30 pages)**

- Development, prototype, test
- Miniaturization/design
- Clinical trials

**1 to 2,5 mn €
12-24 months**

Phase 3

Commercialization

**Input: Promotion
of idea**

- Facilitate access to private finance
- Support by means of networking, trainings, knowledge sharing and dissemination

No Funding

IDEA

MARKET



Clinical research for the validation of biomarkers and/or diagnostic medical devices (Phase 1)



European Commission - Research - Participants
Proposal Submission Forms

Horizon 2020

Call: H2020-SMEINST-1-2014

Topic: PHC-12-2014-1

Type of action: **SME-1**

Proposal number: SEP-210176060

Proposal acronym: BIOPHOS



EUROPEAN COMMISSION

H2020 SME
Head of Unit

Brussels,

Alex STRONGILOS
PRO-ACTINA CHIMIKI TECHNOLOGIA
ANONIMI ETAIRIA
ARCHIMIDOUS STREET 59
194 00 ATHINA
GREECE

Subject: information about the outcome of the evaluation
H2020 - H2020-SMEINST-1-2014
650532 - BIOPHOS



Based on the evaluation results, we regret to inform you that the Agency has decided that no financial contribution could be made for your proposal. The decision was motivated by the following:

The score obtained by your proposal does not reach the minimum threshold.

We recognise the effort you have put into submitting an application for a H2020 SME Phase 1 funding. If you want to improve your application and are considering re-submitting, we recommend that you contact an adviser from the EEN network (<http://een.ec.europa.eu/>). We hope that despite the disappointing result, you will find other ways of carrying out the business ideas reflected in your proposal.

Proposal Evaluation Form



EUROPEAN COMMISSION

Horizon 2020 - Research and Innovation Framework Programme

**Evaluation
Summary Report**

Evaluation Summary Report

Evaluation Result

Total score: 7.61 (Threshold: 13.00)

Criterion 1 - Impact

Score: **3.00** (Threshold: 4.00/5.00 , Weight: 100.00%)

Good. The proposal addresses the criterion well, although improvements would be necessary.

Criterion 2 - Excellence

Score: **2.66** (Threshold: 4.00/5.00 , Weight: 100.00%)

Fair to good. While the proposal broadly addresses the criterion, there are some weaknesses.

Criterion 3 - Quality and efficiency of implementation

Score: **1.95** (Threshold: 4.00/5.00 , Weight: 100.00%)

Poor. The criterion is addressed in an inadequate manner, or there are serious inherent weaknesses.

Clinical research for the validation of biomarkers and/or diagnostic medical devices (Phase 2)

European Commission - Research - Participants
Proposal Submission Forms

Horizon 2020

Call: H2020-SMEINST-2-2014

Topic: PHC-12-2014

Type of action: **SME-2**

Proposal number: SEP-210225391

Proposal acronym: GLIOMARK



EUROPEAN COMMISSION

Executive Agency for Small and Medium-sized Enterprises (EASME)

Unit A2

Head of Unit

Brussels,

Alex STRONGILOS
PRO-ACTINA
ANONIMI ET
ARCHIMIDOU
194 00 ATHINA
GREECE



Subject: Result of Evaluation of proposals / Invitation to grant preparation

Programme/Call: H2020 — H2020-SMEINST-2-2014

Proposal: 673737 — GLIOMARK

Dear **Alex STRONGILOS**,

We are pleased to inform you that the aforementioned proposal has been **favourably evaluated** by the Agency. Consequently, we wish to proceed to the preparation of the Grant Agreement based on your proposal.

Proposal Evaluation Form

Ref. App: 2015/1225408 - 20/02/2014



EUROPEAN COMMISSION

Horizon 2020 - Research and Innovation Framework Programme

**Evaluation
Summary Report**

Call: H2020-SMEINST-2-2014_17-12-2014
Funding scheme: SME instrument phase 2
Proposal number: 673737
Proposal acronym: GLIOMARK
Duration (months): 48
Proposal title: Validation of blood-brain-barrier permeability as a glioma biomarker by means of the radiotracer ^{99m}Tc-tetrofosmin and single-photon emission computer tomography
Activity: PHC-12-2014

Evaluation Summary Report

Evaluation Result

Total score: 13.02 (Threshold: 12.00)

Criterion 1 - Impact

Score: **4.36** (Threshold: 4.00/5.00 , Weight: 100.00%)

Criterion 2 - Excellence

Score: **4.44** (Threshold: 3.00/5.00 , Weight: 100.00%)

Criterion 3 - Quality and efficiency of implementation

Score: **4.22** (Threshold: 3.00/5.00 , Weight: 100.00%)

diagnosis and grading of gliomas as non-invasive method. This had an immediate impact on the type and aggressiveness of subsequent therapy. Gliomas are recognized as a rare tumor disease with poor prognosis and orphan diagnostic designation for GlioTect will guarantee 10 years of market exclusivity upon approval. ProActina, a Greek chemical-speciality SME having identified this innovative niche market opportunity could become global market leader in glioma diagnostics. Development is at TLR level 6-7 and will be lifted to 9. Expected market application is the s

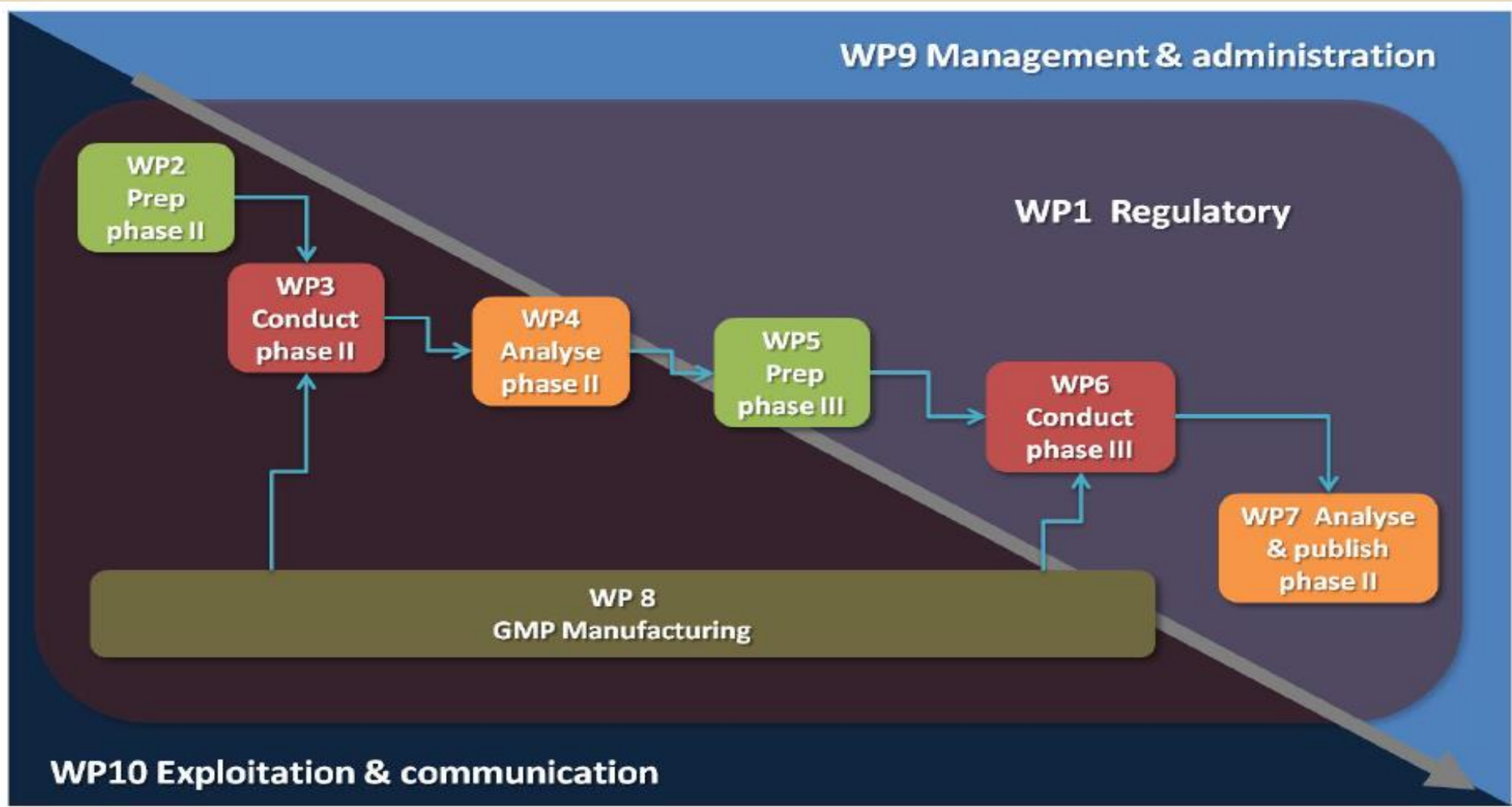
Keys to success:

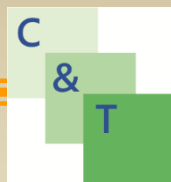
1. **Excellent & Innovative idea**
2. **Selection of partners**
3. **Solid Strategy**
4. **Impact**
 - Company
 - Country
 - European Union (patients, health)
5. **Implementation**
6. **Commercialization plan / Competition**
7. **Securing Intellectual property / Freedom to operate**
8. **Dissemination of knowledge**



EU's Main Evaluation Criteria:

- **Excellence in Innovation**
- **Economic & social impact**
- **Implementation**





Congratulations, you are among the 6% of successful proposals under the European Commission's SME Instrument funding programme. Your project is highly important to



GLIOMARK

Project reference: 673737

Funded under: [H2020-EU.2.3.1.](#), [H2020-EU.3.1.](#)

Validation of blood-brain-barrier permeability of the radiotracer 99mTc-tetrofosmin and SPECT tomography

From 2015-07-01 to 2019-07-01, ongoing project

Project details

Total cost:

EUR 4 705 344

EU contribution:

EUR 4 705 344

Coordinated in:

Germany

Topic(s):

PHC-12-2014 - Clinical research
diagnostic medical devices

Call for proposals:

H2020-SMEINST-2-5

Funding scheme:

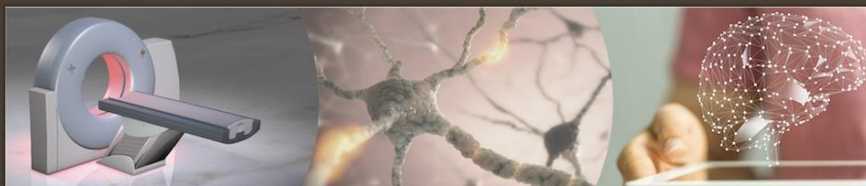
SME-2 - SME instrument

WWW.GLIOMARK.EU

GLIOMARK

NEW DIAGNOSTICS FOR GLIOMAS

- HOME
- THE PROJECT
- THE DISEASE
- NEWS
- CONTACT
- IMPRINT
- SITEMAP



Welcome to the Homepage of the GLIOMARK project!

The objective of GLIOMARK is to clinically validate the permeability of the blood brain barrier (BBB) as an *in vivo* biomarker for the diagnosis and grading of gliomas. This will be accomplished by means of the radiotracer 99mTc-tetrofosmin (TTF) and the imaging technique Single-Photon Emission Computer Tomography (SPECT).

The outcome of GLIOMARK is a diagnostic kit containing handling/diagnostic instructions and tetrofosmin. The latter has to be reconstituted with 99mTc pertechnetate to yield TTF, which will be used in combination with SPECT.

Gliomas are recognized as a rare tumor disease with poor prognosis. Often computer tomography or magnetic resonance imaging are unable to detect low-grade gliomas or distinguish glioma from other diseases. Hence, brain biopsies are necessary for confirmation of diagnosis and grading of gliomas. Our kit employs TTF/SPECT and enables reliable, fast differential diagnosis and grading of gliomas as non-invasive method. This has an immediate impact on the type and aggressiveness of subsequent therapies.

Horizon 2020 SME Instrument beneficiaries

Last update: 07/09/2015

Phase 1 | Phase 2



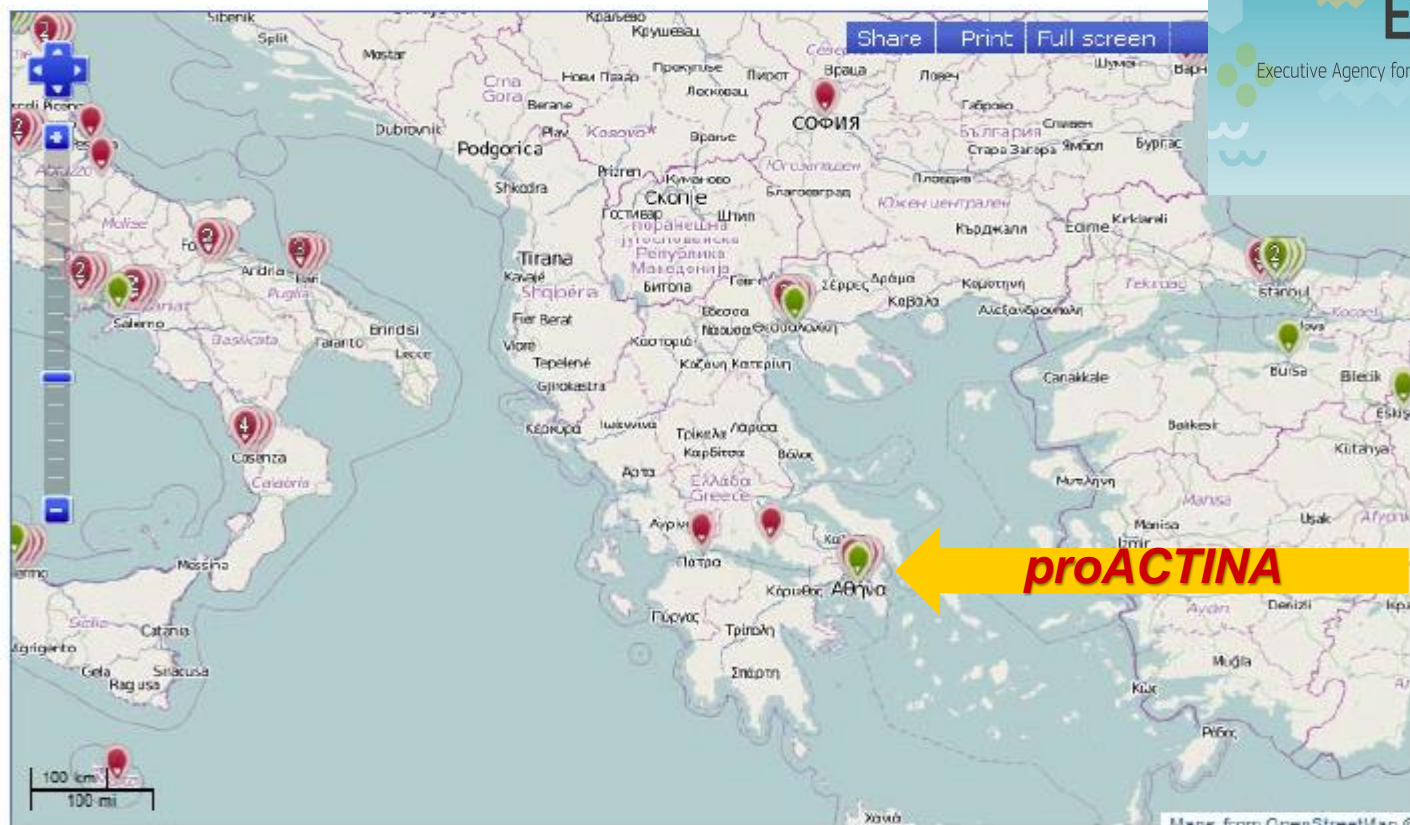
EASME
Executive Agency for Small and Medium-sized Enterprises

<https://ec.europa.eu/easme/en/sme-instrument-beneficiaries>

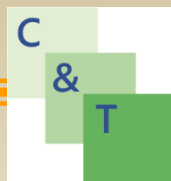
Horizon 2020 SME Instrument beneficiaries

Last update: 07/09/2015

Phase 1 | Phase 2



<https://ec.europa.eu/easme/en/sme-instrument-beneficiaries>



Thank
You!