

Information to be requested from all CA17104 participants:



Photo

<p>Indicate your Working Group(s) in COST Action17104:</p>	<p>Pharmacokinetics and ADMET of the new therapeutic tools; safety of the targets, group 4</p>
<p>First Name:</p>	<p>Simonida</p>
<p>Surname:</p>	<p>Crvenkova</p>
<p>Department</p>	<p>Lung cancer department</p>
<p>Primary Institution</p>	<p>University Clinic of radiotherapy and oncology, Medical faculty</p>
<p>Address of Primary Institution</p>	<p>Mother Theresa 16</p>
<p>Other institutions</p>	
<p>Telephone:</p>	<p>+38970338687</p>

e-mail:	simonidac@hotmail.com
Link to webpage with biography:	https://www.pubfacts.com/author/Simonida+Crvenkova
Link to webpage with group description:	

Orcid ID or Scopus ID	https://www.scopus.com/authid/detail.uri?authorId=12763093300#top
Linkedin	https://mk.linkedin.com/in/simonida-crvenkova-91980214
Expertise relevant for this COST Action:	<p>An expanded access program of Tarceva (erlotinib) in patients with advanced stage IIB/IV non-small cell lung cancer – coinvestigator Simonida Crvenkova MO18109 2005</p> <p>- Multicenter, randomized, double-blind, phase III trial to investigate the efficacy and safety of oral BIBF 1120 plus standard pemetrexed therapy compared to placebo plus standard pemetrexed therapy in patients with stage IIB/IV or recurrent NSCLC after failure of first line chemotherapy- principal investigator- Simonida Crvenkova, 2010-12</p>
Available facilities to conduct work, relevant for this COST Action:	
Materials/Methods that could be shared with other members of this COST Action:	

NOTE: By submitting this form to the Grant Manager of CA17104, I agree that this information can be used within the scope of this COST Action (e.g. may be included on the webpage of CA17104).