

EUROPEAN  
CURRICULUM VITAE  
FORMAT



PERSONAL INFORMATION



Name **TZVEOVA, RENI STOYOVA**  
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Nationality Bulgarian  
Date of birth 06.10.1981

WORK EXPERIENCE

- Dates (from – to) 12.2024 - current
- Name of employer Institute of Experimental Morphology, Pathology and Anthropology with Museum IEMPAM- BAS
- Occupation or position held Assistant professor
  
- Dates (from – to) 06.2020-12.2024
- Name of employer University Multidisciplinary Hospital for Active Treatment "Tsaritsa Joanna - ISUL"
- Type of business or sector analytical, diagnostic
- Occupation or position held Molecular biologist
- Main activities and responsibilities Molecular – genetic diagnostics in patients with various oncological diseases (melanoma, colorectal carcinoma, thyroid tumors and others) in connection with referral to appropriate and effective targeted therapy. Mutational status analysis of *BRAF*, *KRAS*, *NRAS*, *PIK3CA*, *EGFR* and other genes with real-time polymerase chain reaction and digital PCR system.
  
- Dates (from – to) 2019-2020
- Name and address of employer National center of public health and analyzes, Ministry of Health, 15 Acad. I. E. Geshov bul. Sofia, Bulgaria
- Type of business or sector Research, analytical, expert
- Occupation or position held assistant
- Main activities and responsibilities DNA isolation, molecular - genetic analysis of DNA samples

- Dates (from – to) 2017-2018
- Name and address of employer Executive Environment Agency, Ministry of Environment and Water, 136, Tsar Boris III Blvd., Sofia
- Type of business or sector expert
- Occupation or position held Chief Expert
- Main activities and responsibilities DNA isolation, molecular - genetic analysis of plant samples

- Dates (from – to) 2014-2017
- Name and address of employer Molecular Medicine Center, Medical University – Sofia, 2 Zdrave str. 1431 Sofia, Bulgaria
- Type of business or sector Molecular biology and genetics
- Occupation or position held Biologist - geneticist
- Main activities and responsibilities DNA isolation, SNP genotyping, Next Generation sequencing, Sanger sequencing, Data analysis

## EDUCATION AND TRAINING

- Dates (from – to) 2011-2015
- Name and type of organization providing education and training Medical University of Sofia
- Principal subjects/occupational skills covered Pharmacogenetic studies in cardiovascular diseases
- Title of qualification awarded PhD in Molecular genetics
- Level in national classification (if appropriate)

- Dates (from – to) 2010-2012
- Name and type of organization providing education and training Sofia University “St. Kliment Ochridski”
- Principal subjects/occupational skills covered Drug chemistry, drug technology, pharmacology
- Title of qualification awarded Master degree in Medical chemistry
- Level in national classification (if appropriate)

- Dates (from – to) 2008-2010
- Name and type of organization providing education and training Sofia University “St. Kliment Ochridski”
- Principal subjects/occupational skills covered Molecular genetics, oncogenetics, genetic engineering
- Title of qualification awarded Master degree in Genetics
- Level in national classification (if appropriate)

- Dates (from – to) 1999-2006
- Name and type of organization providing education and training Sofia University “St. Kliment Ochridski”
- Principal subjects/occupational skills covered Molecular biology, molecular genetics
- Title of qualification awarded Bachelor degree in Molecular biology
- Level in national classification (if appropriate)

**PERSONAL SKILLS  
AND COMPETENCES**

*Acquired in the course of life and career  
but not necessarily covered by formal  
certificates and diplomas.*

MOTHER TONGUE

OTHER LANGUAGES

- Reading skills
- Writing skills
- Verbal skills

**SOCIAL SKILLS  
AND COMPETENCES**

*Living and working with other people, in  
multicultural environments, in positions  
where communication is important and  
situations where teamwork is essential  
(for example culture and sports), etc.*

**ORGANISATIONAL SKILLS  
AND COMPETENCES**

*Coordination and administration of  
people, projects and budgets; at work, in  
voluntary work (for example culture and  
sports) and at home, etc.*

**TECHNICAL SKILLS  
AND COMPETENCES**

*With computers, specific kinds of  
equipment, machinery, etc.*

**ARTISTIC SKILLS  
AND COMPETENCES**

*Music, writing, design, etc.*

**OTHER SKILLS  
AND COMPETENCES**

*Competences not mentioned above.*

DRIVING LICENCE(S)

**ADDITIONAL INFORMATION**

**BULGARIAN**

**ENGLISH**

EXCELLENT

GOOD

GOOD

COMMUNICATION, TEAMWORK, TOLERANCE, LOYALTY

ORGANIZATION, RESPONSIBILITY, PRECISENESS

WORK AT COMPUTER, WORK AT NEXT GENERATION SEQUENCER MISEQ (ILLUMINA)

[ Describe these competences and indicate where they were acquired. ]

- extraction and purification of nucleic acids from different biological samples
- the full range of electrophoretic techniques
- diagnostic and scanning techniques for mutation detection
- sequencing, analysis of DNA polymorphism, including RFLPs
- PCR

Category B

**Research interest in:**

- Inherited disorders and predisposition
- Pharmacogenomics
- Regulation of gene expression
- Neurodegenerative disorders
- Medical chemistry
- Pharmacoeconomics

## LIST OF PUBLICATIONS:

In foreign journals with an impact factor:

1. Yaneva-Sirakova, Teodora; Dodova, Rumiana; Kaneva, Radka; **Tzveova, Reni**; Ivanova, Raya; Vassilev, Dobrin. A STUDY OF SOME GENETIC FACTORS FOR FIBROMUSCULAR DYSPLASIA. *Journal of Hypertension*: June 2022 - Volume 40 - Issue Suppl 1 - p e220-e221 doi: 10.1097/01.hjh.0000837712.75398.19
2. Penchev V, Boueva A, Kamenarova K, Roussinov D, **Tzveova R**, Ivanova M, Dimitrova V, Kremensky I, Mitev V, Kaneva R, Beltcheva O. A familial case of severe infantile nephronophthisis explained by oligogenic inheritance. *Eur J Med Genet*. 2017 Jun;60(6):321-325.
3. Kamenarova K, Simeonov E, **Tzveova R**, Dacheva D, Penkov M, Kremensky I, Perenovska P, Mitev V, Kaneva R. Identification of a novel de novo mutation of CREBBP in a patient with Rubinstein-Taybi syndrome by targeted next-generation sequencing: a case report. *Hum Pathol*. 2016 Jan;47(1):144-9.
4. Beltcheva O, Boueva A, **Tzveova R**, Roussinov D, Marinova S, Kaneva R, Mitev V. Steroid-resistant nephrotic syndrome caused by novel WT1 mutation inherited from a mosaic parent. *Ren Fail*. 2015 Dec 1:1-4.
5. **Tzveova R.**, Naydenova G., Yaneva T., Dimitrov G., Vandeva S., Matrozova Y., Pendicheva-Duhlenka D., Popov I., Beltheva O., Naydenov C., Tarnovska-Kadrevia R., Nachev G., Mitev V., Kaneva R. Gender specific effect of *CYP2C8\*3* on the risk of essential hypertension in Bulgarian patients. *Biochem Genet*. 2015 Dec;53(11-12):319-33. IF 0.99.
6. Dimitrova-Karamfilova A., **Tzveova R.**, Chilingirova N., Goranova T., Nachev G., Mitev V., Kaneva R. Acenocoumarol pharmacogenetic dosing algorithms and their application in two Bulgarian patients with extremely low anticoagulant requirements. *Biochem Genet*. 2015 Dec;53(11-12):334-50. IF 0.99.
7. **Tzveova R.**, Dimitrova-Karamfilova A., Saraeva R., Solarova T., Naydenova G., Petrova I., Hristova N., Popov I., Nachev G., Mitev V., Kaneva K.. Estimation and validation of acenocoumarol dosing algorithms in Bulgarian patients with cardiovascular diseases. *Personalized medicine*, 12(3), 211–222. IF 1.13.

In specialized Bulgarian journals:

1. Ovcharova E, Danovska M, Marinova D, Pendicheva-Duhlenka D, Tonchev P, Atanasova M, Ruseva A, Shepherd N, **Tzveova R**. Adapted Mediterranean Diet Impact on the Symptoms of Chronic Fatigue, Serum Levels of Omega-3 Polyunsaturated Fatty Acids (PUFAs) and Interleukin 17 (IL-17) in Patients with Relapsing-Remitting Multiple Sclerosis undergoing Disease-Modifying Therapy: A Pilot Study. *J of IMAB*. 2022 Jan-Mar;28(1):4297-4304. DOI: 10.5272/jimab.2022281.4297
2. Yaneva-Sirakova T, Dodova R, Kaneva R, **Tsveova R**, Ivanova R, Vasilev D. Screening for fibromuscular dysplasia in Bulgarian patients and genetic aspects. *Cardiovascular Diseases, Cardiovascular diseases*. 2021, 52, № 2, 29-38.
3. Yordanov S., **Tzveova R.**, Goranova T., Pierini S., Genadieva M., Konov D., Popov T., Kaneva R., Rangachev Y. “Molecular Basis of Laryngeal Carcinogenesis and the Role of Promoter Hypermethylation”, *International Bulletin of Otorhinolaryngology*. 2020; Volume 11: 28-36.
4. **Tzveova R.**, Naydenova G., Yaneva T., S. Vandeva S., Pendicheva-Duhlenka D., Atanasov P., Beltcheva O., Naydenov C., Mitev V., Kaneva R., Georgieva T. „Polymorphic variants in ADIPOQ gene and coronary artery disease risk in Bulgarian population“. *Cardiovascular diseases*. 2019; 3: 11-25.
5. **Tzveova R.**, Georgieva Ts. „Application of CRISPR/CAS9 gene editing technology in the treatment of cardiovascular diseases“. *Cardiovascular diseases*. 2019; 3: 26-32.
6. Yaneva-Sirakova T, **Tzveova R**, Kaneva R, et al. The clinical point of view on whether polymorphic variants of *CYP17A1* and *PLEKHA7* genes can have a role in the development of coronary heart disease. *Cardiovascular diseases*. 2018; 1: 3-7.
7. **Tzveova R.**, Dimitrova – Karamfilova A., Saraeva R., Beltcheva O., Nachev G., Mitev V., Kaneva R. Pharmacogenetics of drug response in Acenocoumarol and warfarin treatment. *Cardiovascular diseases*, 03/2016, XLVII, p. 29-48.

8. **Tzveova R.**, Yaneva T., Dimitrov G., Pendicheva-Duhlenska D., Vandeva S., Matrozova Y., Beltcheva O., Mitkova A., Naydenov C., Zacharieva S., Tarnovska-Kadreva R., Nachev G., Mitev V., Kaneva R. Locus 9P21 and the revolution in cardiovascular genetics – contribution in vascular pathology and evaluation of coronary artery disease risk. *Cardiovascular diseases* 01/2015, XLIV, p.31-40.
9. **Tzveova R.**, Yaneva T., Dimitrov G., Naydenov K., Tarnovska – Kadreva R., Kaneva R., Nachev G., Mitev V. The role of genome-wide association studies for understanding the genetic basis of arterial hypertension – part 2. *Cardiovascular diseases*, 03/2014, XLIV, p. 30-8.
10. **Tzveova R.**, Vandeva S., Matrozova Y., Naydenova G., Pendicheva – Duhlenska D., Mitkova A., Zacharieva S., Nachev G., Mitev V., Kaneva R. Polymorphic variants in leading candidate-genes from genome-wide association studies and coronary artery disease risk – part two. *Cardiovascular diseases*, 02/2014, XLV, p. 48-57.
11. Naydenova G, **Tzveova R**, Nachev G, Mitev V, Kaneva R. The effect of polymorphisms in *PCSK 9* gene on cholesterol levels and the risk of coronary artery disease. *Cardiovascular diseases*, 01/2014, XLV, p. 28-36.
12. Paskaleva I, Dineva D, Baycheva V, **Tzveova R**, Kaneva R, Georgiev B, Gocheva N. The influence of genetic variants *CYP2C19\*2* and *CYP2C19\*17* on ADP – induced platelet aggregation in patients on thienopyridine treatment. *Bulgarian cardiology*, XIX, № 4, 2013, p. 16-26.
13. **Tzveova R.**, Yaneva T., Dimitrov G., Naydenov K., Tarnovska – Kadreva R., Kaneva R., Nachev G., Mitev V. The role of genome-wide association studies for understanding the genetic basis of arterial hypertension. *Cardiovascular diseases*, 03/2013, XLIV, p. 47-54.
14. **Tzveova, R.**, Mitkova A., Kaneva R., Nachev G., Mitev V. The role of genome-wide association studies for understanding the genetic basis of coronary artery disease. *Cardiovascular diseases*, 03/2013, XLIV, p. 55-64.
15. **Tzveova R.**, Mitkova A., Paskaleva I., Kaneva R., Mitev V.. The role of genetic factors in determining the clinical efficacy of clopidogrel. *Cardiovascular diseases*, XLIV, № 1, 2013, p. 16-29.
16. **Tzveova, R.**, Naydenova G., Kaneva R, Tzekova M., Nachev, G., Mitev, V. Role of genetic *polymorphisms in plasminogen activator inhibitor type 1 (PAI-1) and apolipoprotein E (ApoE) genes* in development of acute myocardial infarction. *Bulgarian cardiology*, **XVIII**, № 4, 2012, p. 36-42.
17. Naydenova G, **Tzveova R**, Kaneva R, Tzekova M. Genetic polymorphisms in *CYP2C8, CYP2C9, CYP2J2* and risk of coronary artery disease. *Medinfo*, 04/2012, Year XII, 4/2012, page 38-41