

# Andrea Spinaci

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## PERSONAL

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### EDUCATION

31/03/2017

**PhD in Chemical and Pharmaceutical Sciences and Biotechnology**  
University of Camerino, Camerino

07/03/2013

**Master degree, Pharmaceutical Chemistry and Technology**  
University of Camerino, Camerino  
Vote: 105/110

06/2007

**High School Diploma – Liceo Scientifico**  
Liceo Scientifico “Galileo Galilei” Macerata

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### EMPLOYMENT

1/10/2017– Current

**Research Fellowship in Medicinal Chemistry**  
“*Sintesi innovative di farmaci adenosinici per la terapia delle malattie neurodegenerative.*”  
University of Camerino, Camerino (Italy)

1/02/2017–31/07/2017

**Post-Doc Fellowship in Medicinal Chemistry**  
University of Camerino, Camerino (Italy)

1/02/2017-31/07/2017

**Internship in Laboratory of Forensic Toxicology and Forensic Medicine**  
*University of Macerata, Macerata (Italy)*

1/02/2014 – 31/01/2017

**PhD in Chemical and Pharmaceutical Sciences and Biotechnology**  
Title of the thesis: “*Design, synthesis, and characterization of new p1 receptor ligands.*”  
University of Camerino, Camerino (Italy)

08/03/2013–07/07/2013

**Post-graduate fellowship in Medicinal Chemistry**  
University of Camerino, Camerino (Italy)

05/03/2012–05/03/2013

**Experimental Thesis in Medicinal Chemistry**  
Title of the thesis: “*Multitarget-ligands as advantageous tools in reducing the opiate withdrawal syndrome and related comorbidities.*”  
Università di Camerino, Camerino (Italy)

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01/06/2011–30/01/2012

**Apprenticeship in Pharmacy**

Farmacia Cairoli, Fabrizio Buglioni, Macerata (Italy)

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**DIDACTICAL  
EXPERIENCE**

**Supervisor of Experimental Thesis**

Master Degree in Pharmacy (LM-13); Experimental Thesis in Medicinal Chemistry “*Adenosine 2,N<sup>6</sup>-disostituite come ligandi del recettore adenosinico a<sub>3</sub> umano con potenziale attivita' antitumorale*”

Graduate Student: Ludovica Mariani,

Supervisor: Prof. Rosaria Volpini, **Dr Andrea Spinaci** (a.a. 2017-2018)

**Supervisor of Experimental Thesis**

Master Degree in Pharmacy (LM-13); Experimental Thesis in Medicinal Chemistry “*Sintesi e caratterizzazione di analoghi dell'ATP come antagonisti del recettore P2X<sub>3</sub>*”

Graduate Student: Fabiola Paoletti,

Supervisor: Prof. Catia Lambertucci, **Dr Andrea Spinaci** (a.a. 2016-2017)

**Supervisor of Experimental Thesis**

Master Degree in Pharmacy (LM-13); Experimental Thesis in Medicinal Chemistry “*Derivati disostituiti della 9-etileadenina come nuovi antagonisti del recettore adenosinico A<sub>2A</sub>*”

Graduate Student: Valeria Marcantoni,

Supervisor: Prof. Catia Lambertucci, **Dr Andrea Spinaci** (a.a. 2016-2017)

12/03/2019 – 18/04/2019

**Tutor in Medicinal Chemistry Lab**

University of Camerino, Camerino (Italy)

13/04/2015 – 29/04/2015

**Tutor in Drug Quantitative Analysis Course**

University of Camerino, Camerino (Italy)

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**INTERNATIONAL  
EXPERIENCE**

01/05/2016–11/09/2016

**Mobility periods abroad**

Institute of Medical Science (IMS)

Supervisor: Prof. Matteo Zanda

University of Aberdeen, Aberdeen (United Kingdom)

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**SKILLS**

**Language** – Good: English, mother tongue: Italian

**Computer Skills** - Microsoft Office, TopSpin and CMC-A Software for NMR spectra analysis, ChemBioDraw, Acrobat Reader, Xcalibur LC/GC-MS Software, pKcsm Software

**Synthetic Skills** - Nucleophilic aromatic substitution, cross-coupling reaction (Sonogashira coupling, Suzuki reaction), Sandmeyer reaction, nucleoside and nucleotide synthesis, oxychlorination, allylic oxidation, protecting group reaction, glycosylation, reduction and oxidation.

**Purification Skills** – Gravimetric and flash chromatography, reversed

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phase chromatography, ion exchange chromatography, HPLC, crystallization, liq-liq extraction, distillation, preparation of dry solvents.

**Characterization Skills** – <sup>1</sup>H-NMR and <sup>13</sup>C-NMR spectra analysis, IR-spectra analysis; UV-spectra analysis; polarimetric analysis; GC-mass analysis and HPLC-MS analysis.

**Instrumentation Use Ability** - Bruker AVANCE III 400/500 NMR spectrometer, ESI-MS, LC/GC-MS Agilent, Thermo Orbitrap LC-MS, BUCHI melting point, DSX™ Automated Four-Plate ELISA System

**Biological data analysis** – Interpretation of binding affinity data and functional activity at cells culture or ex vivo assay, Elisa data analysis, preliminary multi drug test analysis for drug abuse compounds (THC, amphetamine, morphine, cocaine, etc), alcoholemia misuration.

**Sample Preparation** – Preparation and extraction from biological material (blood, hair, urine, bile) of substances of abuse by liq-liq extraction, SPE column; silyl derivatization, acid or enzymatic glucuronide hydrolysis.

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**ATTENDED  
CONFERENCE AND  
SEMINAR**

04-07/09/2019

**1<sup>st</sup> European Purine Meeting**

*Universidad Santiago de Compostela, Spain*

18/01/2019

**Riunione Annuale del Purine Club Italiano**

*Università di Firenze - Firenze*

5/10/2018

**XXXVI CONVEGNO INTERREGIONALE TUMA 2018**

*Centro Congressi “Le Benedettine”, Pisa*

25-28/10/2017

**38° Congresso nazionale società italiana di Farmacologia**

*Palacongressi Rimini*

16/5/2017

**LC-MS/MS: applicazioni per l’analisi di molecole endogene ed esogene.**

*SCIEX, SeePort Hotel, Ancona*

15/1/2016

**Riunione Annuale del Purine Club Italiano**

*Istituto Superiore della Sanità, Roma*

29/06/2014–03/07/2014

**Thematic course: European School of Medicinal Chemistry**

*University of Urbino Carlo Bo*

26/06/2015-30/06/2015

**Thematic course: European School of Medicinal Chemistry**  
*University of Urbino Carlo Bo*

27/10/2015

**Sigma-Aldrich Young Chemists Symposium (SAYCS 2015) SCI**  
*Società Chimica Italiana, Rimini*

23-25/09/2015

**TUMA XXXIV – Convegno delle sezioni Toscana Umbria Marche  
Abruzzo della Società Chimica Italiana**  
*Società Chimica Italiana, Perugia*

11/06/2015

**Statistics and Data Management**  
Speaker: Mauro Angeletti  
*University of Camerino*

16/10/2015

**Innovation management: how to manage and develop innovative  
projects at industrial level**  
Speaker: Cristina Cristalli  
*University of Camerino*

14/10/2015

**Writing a grant proposal**  
Speaker: Ciro Franco  
*University of Camerino*

15/10/2015

**Job Hunt**  
Speaker: Micheal Zebrak  
*University of Camerino*

12/10/2015

**Proprietà intellettuale e comunicazione scientifica nella società  
dell'informazione**  
Speaker: Avv. Simone Alipardi  
*University of Camerino*

10/02/2015

**Farmaci per il trattamento delle malattie virali**  
Speaker: Romano Silvestri  
*University of Camerino*

**MEMBERSHIP**

*Società Chimica Italiana (SCI)*  
*Sezione Chimica Farmaceutica*

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**PUBLICATIONS**

1) Lambertucci, C.; **Spinaci, A.**; Buccioni, M.; Dal Ben, D.; Ngnintedem, M. A. N.; Kachler, S.; Marucci, G.; Klotz, K.-N.; Volpini, R., New A<sub>2A</sub> adenosine receptor antagonists: A structure-based upside-down interaction in the receptor cavity. *Bioorganic Chemistry* **2019**, 92, art. no. 103183.

**I.F. 3.926**

2) Betti, M.; Catarzi, D.; Varano, F.; Falsini, M.; Varani, K.; Vincenzi, F.; Pasquini, S.; Di Cesare Mannelli, L.; Ghelardini, C.; Lucarini, E. Dal Ben., D.; **Spinaci, A.**; Bartolucci, G.; Menicatti, M.; Colotta, V. Modifications on the amino-3,5-dicyanopyridine core to obtain multifaceted adenosine receptor ligands with antineuropathic activity. *Journal of Medicinal Chemistry* **2019**, 62 (15), 6894-6912.

**I.F. 6.054**

3) Buccioni, M.; Dal Ben, D.; Lambertucci, C.; Navia, A. M.; Ricciutelli, M.; **Spinaci, A.**; Volpini, R.; Marucci, G. New sensible method to quantize the intestinal absorption of receptor ligands. *Bioorganic & Medicinal Chemistry* **2019**, 27 (15), 3328-3333.

**I.F. 2.802**

4) Marucci, G.; Dal Ben, D.; Lambertucci, C.; Martí Navia, A.; **Spinaci, A.**; Volpini, R.; Buccioni, M. GPR17 receptor modulators and their therapeutic implications: review of recent patents. *Expert Opinion on Therapeutic Patents* **2019**, 29 (2), 85-95.

**I.F. 3.699**

5) Dal Ben, D.; Buccioni M.; Lambertucci, C.; Marucci, G.; **Spinaci, A.**; Marchenkova, A.; Abdelrahman, A.; Nistri, A.; Müller, C. E.; Volpini, R. Investigation on 2',3'- O-Substituted ATP Derivatives and Analogs as Novel P2X<sub>3</sub> Receptor Antagonists. *ACS Medicinal Chemistry Letters* **2019**, 10 (4), 493-498.

**I.F. 3.737**

6) Amenta, F.; Buccioni, M.; Dal Ben, D.; Lambertucci, C.; Martin Navia, A.; Ngouadjeu Ngnintedem, M. A.; Ricciutelli, M.; **Spinaci, A.**; Volpini, R.; Marucci, G. Ex-vivo absorption study of lysine R-lipoate salt, a new pharmaceutical form of R-ALA. *European Journal of Pharmaceutical Sciences* **2018**, 118, 200-207.

**I.F. 3.532**

7) Lambertucci, C.; Marucci, G.; Dal Ben, D.; Buccioni, M.; **Spinaci, A.**; Kachler, S.; Klotz, K.-N.; Volpini, R. New potent and selective A<sub>1</sub> adenosine receptor antagonists as potential tools for the treatment of gastrointestinal diseases. *European Journal of Medicinal Chemistry* **2018**, 151, 199-213.

**I.F. 4.833**

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8) Zanato, C.; Pelagalli, A.; Marwick, K. F. M.; Piras, M.; Dall'Angelo, S.; **Spinaci, A.**; Pertwee, R. G.; Wyllie, D. J. A.; Hardingham, G. E.; Zanda, M. Synthesis, radio-synthesis and in vitro evaluation of terminally fluorinated derivatives of HU-210 and HU-211 as novel candidate PET tracers. *Organic and Biomolecular Chemistry* **2017**, 15 (9), 2086-2096.

**I.F. 3.490**

9) Dal Ben, D.; Marchenkova, A.; Thomas, A.; Lambertucci, C.; **Spinaci, A.**; Marucci, G.; Nistri, A.; Volpini, R. 2',3'-O-Substituted ATP derivatives as potent antagonists of purinergic P2X3 receptors and potential analgesic agents. *Purinergic Signalling* **2017**, 13 (1), 61-74.

**I.F. 3.038**

10) Dal Ben, D.; Buccioni M.; Lambertucci, C.; Marucci, G.; Santinelli, C.; **Spinaci, A.**; Thomas, A.; Volpini, R. Simulation and Comparative Analysis of Different Binding Modes of Non-nucleoside Agonists at the A<sub>2A</sub> Adenosine Receptor. *Molecular Informatics* **2016**, 35, 403-413.

**I.F. 2.375**

11) Marucci, G.; Dal Ben, D.; Lambertucci, C.; Santinelli, C.; **Spinaci, A.**; Thomas, A.; Volpini, R.; Buccioni, M. The G Protein-Coupled Receptor GPR17: Overview and Update. *ChemMedChem* **2016**, 11(23), 2567-2574.

**I.F. 3.016**

12) Thomas, A.; Buccioni M.; Dal Ben, D.; Lambertucci, C.; Marucci, G.; Santinelli, C.; **Spinaci, A.**; Kachler, S.; Klotz, K.-N.; Volpini, R. The Length and Flexibility of the 2-Substituent of 9-Ethyladenine Derivatives Modulate Affinity and Selectivity for the Human A<sub>2A</sub> Adenosine Receptor. *ChemMedChem* **2016**, 11, 1829-1839.

**I.F. 3.016**

13) Lambertucci, C.; Buccioni M.; Dal Ben D.; Kachler, S.; Marucci, G.; **Spinaci, A.**; Thomas, A.; Klotz, K.-N.; Volpini, R. New substituted 9-propyladenine derivatives as A<sub>2A</sub> adenosine receptor antagonists. *MedChemComm* **2015**, 6, 963-970.

**I.F. 2.394**

*h*-INDEX: 4

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**CONFERENCE  
PRESENTATIONS**

**Spinaci A.**, Lambertucci C., Dal Ben D., Buccioni M., Ngouadjieu M. A., Marti Navia A., Marucci G., Volpini R. "New 4'-tetrazolyl adenosine derivatives as potential agents for wound healing". *1<sup>st</sup> European Purine Meeting, Universidad Santiago de Compostela - Spain -04-07/09/2019*

**Spinaci A.**, Buccioni M., Dal Ben D., Lambertucci C., Marucci G, Marti Navia A., Ngouadjieu M. A, Volpini R. "New A<sub>2A</sub> adenosine receptor agonists as potential agents for wound-healing".

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**(PERSONAL ORAL PRESENTATION)**

**Spinaci A.**, Lambertucci C., Graiff C., Buccioni M., Dal Ben D., Marucci G., Volpini R. “Synthesis and characterization of new triazolotriazinic derivatives as antagonists of A<sub>2A</sub> adenosine receptor”. XXXVI

CONVEGNO INTERREGIONALE TUMA 2018 – 5 October 2018 Centro Congressi “Le Benedettine”, Pisa

**(PERSONAL ORAL PRESENTATION)**

Ngouadjeu Ngnintedem M. A., **Spinaci A.**, Lambertucci C., Buccioni M., Dal Ben D., Marucci G., Volpini R. “Synthesis of new A<sub>3</sub> adenosine receptor antagonists”. XXXVI CONVEGNO INTERREGIONALE TUMA 2018– 5 October 2018 Centro Congressi “Le Benedettine”, Pisa

**Spinaci, A.**, Lambertucci, C., Buccioni, M., Dal Ben, D., Kachler, S., Marucci, G., Thomas, A., Klotz, K.-N., Volpini, R.

“Nucleotide derivatives as new ligands of the purinergic P2 Receptors”

38° Congresso nazionale Società Italiana Farmacologia, Palacongressi di Rimini, Rimini (Italy) October 28, 2017

**(PERSONAL ORAL PRESENTATION)**

C. Lambertucci, D. Dal Ben, M. Buccioni, G. Marucci, **A. Spinaci**, K.-N. Klotz, R. Volpini “A<sub>2A</sub>R agonists/A<sub>3</sub>R antagonists: design, synthesis, and biological evaluation of new ligands with dual activity” 7th Joint Italian-German Purine Club Meeting – “Advances in basic and translational purinergic research”, Rome. July 20-22, 2017

G. Marucci, M. Buccioni, D. Dal Ben, C. Lambertucci, **A. Spinaci**, M. P. Abbracchio, R. Volpini “Pharmacological characterization of the GPR17 receptor dual profile” XXIV National Meeting in Medicinal Chemistry -10th Young Medicinal Chemists’ Symposium, Perugia. September 11-14, 2016

A. Thomas, D. Dal Ben, C. Lambertucci, A. Marchenkova, **A. Spinaci**, G. Marucci, A. Nistri, R. Volpini “2’,3’-Substituted ATP derivatives as nanomolar antagonists of the purinergic P2X<sub>3</sub> receptors and potential analgesic agents” XXIV National Meeting in Medicinal Chemistry -10th Young Medicinal Chemists’ Symposium, Perugia. September 11-14, 2016

M. A. Ngouadjeu, M. Buccioni, D. Dal Ben, C. Lambertucci, G. Marucci, C. Santinelli, **A. Spinaci**, A. Thomas, R. Volpini “Versatile synthesis of 2,5-disubstituted triazolotriazine derivatives as A<sub>2A</sub> adenosine receptor antagonists” 33rd Camerino-Cyprus Symposium - Receptor Chemistry: Reality and Vision, Camerino. May 15-19, 2016

A. Thomas, M. Buccioni, D. Dal Ben, C. Lambertucci, G. Marucci, M. A. Ngouadjeu, A. Nistri, C. Santinelli, **A. Spinaci**, R. Volpini “Constrained ribose ATP derivatives as new potent antagonists for the purinergic P2X<sub>3</sub> receptors” 33rd Camerino-Cyprus Symposium - Receptor Chemistry: Reality and Vision, Camerino. May 15-19, 2016

A. Thomas, M. Buccioni, D. Dal Ben, C. Lambertucci, G. Marucci, M.

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A. Ngouadjeu, C. Santinelli, **A. Spinaci**, S. Kachler, K.-N. Klotz, Rosaria Volpini “*Probing the 2-position of 9-ethyladenine for potent and selective A2A adenosine receptor antagonists*” 33rd Camerino-Cyprus Symposium - Receptor Chemistry: Reality and Vision, Camerino. *May 15-19, 2016*

**Spinaci, A.**, Lambertucci, C., Buccioni, M., Dal Ben, D., Kachler, S., Marucci, G., Thomas, A., Klotz, K.-N., & Volpini, R. *New Adenine Nucleoside as ligands with dual activity at the adenosine receptors*. Italian Purine Club 2015, Istituto Superiore della Sanità, Roma (Italy) *January 16, 2016*

**(PERSONAL ORAL PRESENTATION)**

**A. Spinaci**, M. Buccioni, D. Dal Ben, C. Lambertucci, C. Santinelli, A. Thomas, G. Marucci, R. Volpini “*Design, synthesis, and pharmacological characterization of 2,N<sup>6</sup>-disubstituted adenosine analogues of adenosine receptor ligands*” Purinergic Signaling, Vancouver (Canada). *January 24–28, 2016*

A. Thomas, M. P. Abbracchio, M. Buccioni, D. Dal Ben, C. Lambertucci, C. Martini, G. Marucci, C. Santinelli, **A. Spinaci**, R. Volpini “*Design, synthesis, and biological evaluation of new GPR17 ligands*” SAYCS 2015, Rimini. *October 27-29, 2015*

**Spinaci, A.**, Lambertucci, C., Buccioni, M., Dal Ben, D., Kachler, S., Marucci, G., Thomas, A., Klotz, K.-N., & Volpini, R. *New substituted 9-propyladenine derivatives as A2A adenosine receptor antagonists*. XXXIV Convegno delle Sezioni Toscana, Umbria, Marche e Abruzzo 2015 SCI - Società Chimica Italiana, Perugia (Italy) *September 24, 2015*  
**(PERSONAL ORAL PRESENTATION)**

**Spinaci, A.**, Thomas, A., Buccioni, M., Dal Ben, D., Lambertucci, C., Marucci, G., Santinelli, C., Klotz, K.-N. & Volpini, R.  
“*New N<sup>6</sup>-substituted adenosine derivatives as ligands with dual activity at adenosine receptors.*” Sigma-Aldrich Young Chemists Symposium (SAYCS 2015) SCI - Società Chimica Italiana, Rimini (Italy) *October 28, 2015*

M. Buccioni, D. Dal Ben, C. Lambertucci, G. Marucci, C. Santinelli, **A. Spinaci**, A. Thomas, K.-N. Klotz, R. Volpini “*New N<sup>6</sup>- substituted adenosine derivatives as ligands with dual activity at adenosine receptors*” XXIII Congresso National Meeting in Medicinal Chemistry (NMMC), Salerno. *September 6-9, 2015*

M. Buccioni, D. Dal Ben, C. Lambertucci, G. Marucci, C. Santinelli, **A. Spinaci**, A. Thomas, M. P. Abbracchio, R. Volpini “*Characterization of the GPR17 receptor dual pharmacological profile*” XXIII Congresso National Meeting in Medicinal Chemistry (NMMC), Salerno. *September 6 - 9 2015*

D. Dal Ben, M. Buccioni, C. Lambertucci, G. Marucci, C. Santinelli, **A. Spinaci**, A. Thomas, S. Kachler, R. Volpini “*9-Propyladenine derivatives as A2A adenosine receptor antagonists*” XXIII Congresso

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National Meeting in Medicinal Chemistry (NMMC), Salerno. *September 6 - 9 2015*

A. Thomas, M. Buccioni, D. Dal Ben, C. Lambertucci, G. Marucci, C. Santinelli, **A. Spinaci**, M. P. Abbracchio, R. Volpini “*Stable nucleotide analogues as potent ligands for the purinergic P2Y and P2Y-like receptors*” XXIII Congresso National Meeting in Medicinal Chemistry (NMMC), Salerno. *September 6 - 9 2015*

A. Thomas, M. Buccioni, D. Dal Ben, C. Lambertucci, G. Marucci, C. Santinelli, **A. Spinaci**, K.-N. Klotz, R. Volpini. “*Upside-down adenine scaffold for the design of A<sub>2A</sub> adenosine receptors ligands*” XXIII Congresso National Meeting in Medicinal Chemistry (NMMC), Salerno. *September 6 - 9 2015*

G. Marucci, M. Buccioni, D. Dal Ben, C. Lambertucci, C. Santinelli, **A. Spinaci**, A. Thomas, K.-N. Klotz, R. Volpini “*Synthesis and biological characterization of 9-propyladenine derivatives as ligands for adenosine receptors*” 6th Joint Italian-German Purine Club Meeting, Hamburg (Germany). *July 23-25, 2015*

C. Lambertucci, M. Buccioni, G. Marucci, D. Dal Ben, **A. Spinaci**, A. Thomas, K.-N. Klotz, R. Volpini “*New ligands for adenosine receptors*” 6th Joint Italian-German Purine Club Meeting, Hamburg (Germany). *July 23-25, 2015*

C. Lambertucci, M. Buccioni, G. Marucci, D. Dal Ben, **A. Spinaci**, A. Thomas, A. Nistri, R. Volpini “*New purine derivatives as antagonists of P2X3 receptors*” 6th Joint Italian-German Purine Club Meeting, Hamburg (Germany). *July 23-25, 2015*

R. Volpini, M. Buccioni, D. Dal Ben, C. Lambertucci, G. Marucci, C. Santinelli, **A. Spinaci**, A. Thomas “*New ligands for the dual receptor GPR17*” XXV National Congress of the Italian Chemical Society SCI 2014, Arcavacata di Rende. *September 7-12, 2014*

D. Dal Ben, M. Buccioni, C. Lambertucci, G. Marucci, C. Santinelli, **A. Spinaci**, A. Thomas, R. Volpini “*Molecular modeling studies on A<sub>2B</sub> adenosine receptor and its ligands*” XXV National Congress of the Italian Chemical Society SCI 2014, Arcavacata di Rende. *September 7-12, 2014*

C. Lambertucci, M. Buccioni, D. Dal Ben, G. Marucci, C. Santinelli, **A. Spinaci**, A. Thomas, R. Volpini “*P1 Receptor Ligands: Design, Synthesis, and Biological Activity of New Adenosine Analogues*” XXV National Congress of the Italian Chemical Society SCI 2014, Arcavacata di Rende. *September 7-12, 2014*

D. Dal Ben, M. Buccioni, C. Lambertucci, G. Marucci, **A. Spinaci**, A. Thomas, R. Volpini “*Development of structural models of purinergic P2X receptors and analysis of interaction with agonists and antagonists*” Purines 2014 – International Conference on Nucleotides, Nucleosides and Nucleobases, Bonn (Germany). *July 23-27, 2014*

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D. Dal Ben, M. Buccioni, C. Lambertucci, G. Marucci, **A. Spinaci**, A. Thomas, K.-N. Klotz, R. Volpini “*Different efficacy of adenosine and NECA derivatives at the human A3 adenosine receptor: molecular modeling analysis and insight into the receptor activation switch*” Purines 2014 – International Conference on Nucleotides, Nucleosides and Nucleobases, Bonn (Germany). July 23-27, 2014

A. Thomas, M. Buccioni, C. Lambertucci, D. Dal Ben, G. Marucci, C. Santinelli, **A. Spinaci**, R. Volpini “*Design, synthesis, and pharmacological characterization of 2,N<sup>6</sup>-disubstituted adenosine analogues as P1 receptor ligands*” Purines 2014 – International Conference on Nucleotides, Nucleosides and Nucleobases, , Bonn (Germany). July 23-27, 2014

C. Lambertucci, M. Buccioni, D. Dal Ben, G. Marucci, **A. Spinaci**, A. Thomas, R. Volpini “*Upside-down adenine derivatives as new antagonists for A<sub>2A</sub> receptors*” Purines 2014 – International Conference on Nucleotides, Nucleosides and Nucleobases, Bonn (Germany). July 23-27, 2014

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Camerino, 3 settembre 2019