

EUROPEAN JOURNAL OF NEUROLOGY

Volume 29, Supplement 1, June 2022

Abstracts of the 8th Congress of the European Academy of Neurology

Vienna, Austria

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ISSN 1468-1331 (202206)29:6+1

- Altehheld, L. 596
 Altieri, M. 725, 815
 Altmann, P. 204, 456, 860
 Altomare, D. 148
 Altomare, S. 881
 Altuna, M. 214
 Altunrende, B. 785
 Alvarenga, J. 392
 Alvares, A. 895
 Alvarez, E. 628, 631, 634
 Alvarez, V. 127
 Alvarez-Baron, E. 402
 Álvarez Bravo, G. 815, 858, 869
 Alvarez de Azevedo, C. 908
 Álvarez-Mariño, B. 585
 Álvarez Muelas, A. 540
 Álvarez Rodríguez, E. 900
 Álvarez Saucó, M. 176, 445
 Alvaro, B. 211
 Alve, P. 424
 Alves, F. 515, 840
 Alves, G. 841
 Alves, J. 666, 667, 872
 Alves, L. 570
 Alves, P. 82, 134, 638, 884
 Alves Vitorino da Silva, U. 486, 905
 Alwagdani, A. 910
 Alyavi, A. 555
 Alyavi, B. 555
 Amador, M. 606
 Amarenco, G. 918
 Amaro, S. 804
 Amatachaya, P. 877
 Amatachaya, S. 756, 877
 Amato, M. 7, 215, 261
 Amaya Pascasio, L. 758, 882
 Ambawatte, S. 342
 Ambrosino, P. 201
 Ambroz, P. 846
 Amelin, A. 178
 Ameer, A. 891
 Amin, F. 192, 417
 Amiri, M. 126, 128, 711, 713
 Amisano, P. 447
 Amoiridis, G. 239
 Amore, G. 166, 803
 Amorim, J. 893
 Amprosi, M. 114, 319, 653
 An Haack, K. 59
 Anagnostou, E. 108, 239, 515
 Anagnostouli, M. 881
 Ananeva, L. 854
 Anan, I. 656
 Anca-Herschkovitsch, M. 449
 Andabaka, M. 188
 Andelová, M. 726
 Andersen, A. 243
 Andersen, G. 596
 Andonova, S. 906
 Andrade, C. 720, 849
 Andrade, M. 873
 Andrade Zumárraga, L. 758
 Andrée Larsen, V. 126, 128
 Andreetta, F. 239
 Andreev, M. 522, 847
 Andreou, A. 13
 Andrés Bartolomé, A. 907
 Andresen, H. 185
 Andresen, O. 157
 Andrikogiannopoulos, P. 668
 Andris, C. 688
 Andrzejewska-Gorczyńska, N. 554
 Angela, C. 211
 Angelini, L. 447, 452, 765
 Angelin, L. 859
 Angeloni, B. 896
 Angelopoulou, E. 182, 530, 531, 877, 879, 889, 898, 907
 Angelov, T. 651
 Anheim, M. 117
 Aniello, M. 881
 Aniola, J. 554
 Anna Luisa, A. 209
 Annane, D. 61
 Annen, J. 196, 713, 805, 846
 Annetorp, M. 146
 Annovazzi, P. 725
 Annushkin, A. 551
 Annushkin, V. 551, 878, 882
 Ansari, S. 142, 207
 Anschau, F. 888
 Anshu, P. 896
 Antal, A. 25
 Antelmi, E. 323
 Antia, N. 854
 Antipin, V. 916
 Antochi, F. 842, 909
 Antognozzi, S. 343
 Antoli Martinez, H. 400
 Antonellou, R. 530
 Antonelou, R. 898
 Antonenko, A. 391
 Antonenko, K. 391
 Antoniazzi, E. 851
 Antonini, A. 6, 178, 439, 440, 443, 446, 450, 451, 597, 619, 897
 Antonioni, A. 683
 Antonoglou, A. 182, 853, 858
 Antonova, K. 878, 881
 Antos, A. 176, 778, 858
 Antozzi, C. 352, 469, 870
 Antunes, A. 867, 870
 Antunes, F. 838
 Anwar, M. 742
 Apap Mangion, S. 190
 Apostolakopoulou, L. 878
 Appeltshauser, L. 109
 Appleton, P. 342
 Appollonio, I. 260, 379, 537, 869
 Aprile, D. 839
 Aprile, F. 806
 Aprile, M. 781
 Aquila, F. 676
 Aquilina, J. 913
 Ara, R. 850
 Ara Callizo, J. 900
 Aragon-gawinska, K. 605
 Aramini, S. 608
 Aran, K. 753
 Araújo, A. 382
 Arbia Boudjelthia, F. 863
 Arbuzova, E. 860
 Archelos-Garcia, J. 301
 Archetti, S. 248
 Ardashirova, N. 858
 Ardicli, D. 58
 Aremu, S. 870
 Arena, I. 645, 799
 Arends, M. 847
 Arenga, M. 888
 Arestova, A. 826
 Aretini, P. 209, 676
 Arévalo Bernabé, Á. 630
 Argyropoulou, A. 668
 Argyropoulou, C. 909
 Arhan, E. 405
 Arhire, N. 675, 873
 Arias-Villaran, M. 871
 Arici, D. 487
 Aridon, P. 488
 Arighi, A. 559, 846
 Aringhieri, G. 107
 Ariño, H. 98
 Aristeidou, S. 108
 Ariöz, B. 839
 Arjona Padillo, A. 758, 855
 Arkadir, D. 616
 Arlati, F. 545
 Arlati, S. 887
 Armangué, T. 241, 358
 Armenis, G. 515
 Armentano, A. 497
 Armon, C. 313
 Arnaldi, D. 94, 101, 442, 496, 886
 Arnaud-Hevi, L. 865
 Arnesen, A. 453, 610
 Arnold, D. 65, 150, 223, 285, 458
 Arnold, M. 53, 76, 78, 718, 842
 Arnould, S. 142, 207
 Arola, A. 135
 Arrambide, G. 347
 Arranz, J. 214, 534
 Arrese Regañón, I. 662
 Arriola-Infante, J. 714
 Arsava, E. 406
 Arslan, D. 426, 860
 Arteche-López, A. 649, 891
 Artemiadis, A. 907, 908, 909
 Artukoglu, B. 625
 Artusi, C. 623
 Arzimanoglou, A. 567
 Arzt, M. 862
 Arzumian, N. 459
 Ascencio Muñoz, L. 880
 Asenbaum-Nan, S. 861
 Ashina, M. 192, 417, 418
 Ashkenazi, N. 467
 Ashotn, N. 877
 Asilova, N. 164
 Asri, S. 773, 853
 Assar, H. 289
 Assenza, G. 399
 Assis, P. 888
 Assouline, A. 606
 Astrea, G. 107
 Ataniyazov, M. 697
 Atri, A. 537
 Atrous, A. 885, 901, 913
 Attaallah, B. 695
 Attarian, S. 59
 Audronytė, E. 838
 Auer, M. 188, 784
 Auger, C. 347
 Augusta Guimarães Dourado, J. 847
 Aungst, A. 284
 Aureli, F. 613
 Aurilia, C. 86, 194
 Avalos-Pavon, R. 313
 Avanzini, I. 853
 Avanzo, A. 876
 Avelar Rodrigues, M. 915
 Avenali, M. 684, 874, 896
 Averchenkov, D. 459
 Avetisyan, L. 391
 Avila, L. 689
 Ávila Rivera, M. 445
 Avino, G. 557, 840
 Avolio, C. 349
 Avoni, P. 769, 803
 Avorio, F. 914
 Axelsson, M. 157
 Axenhus, M. 68, 74
 Ay, S. 416
 Ayadi, K. 892, 917
 Ayasse, N. 375

Cerebrovascular diseases 4

EPO-493

The potential Drug-Drug Interactions (pDDIs) which include antimicrobials in patients with acute ischemic stroke

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Background and aims: Prevalence of post-stroke infection is up to 65% of patients. Potential drug–drug interactions (pDDIs) are among the leading preventable causes of adverse drug events. Antimicrobials are among the common drug groups in studies about pDDI.

Methods: A 3-years retrospective study was conducted at the Clinic of Neurology, University Clinical Center Kragujevac, Serbia. A total of 696 patients with acute ischemic stroke (AIS) have been hospitalized in the neurological intensive care unit (NICU). The Micromedex software was used to determine pDDIs which include antimicrobials.

Results: From 552 (79.3%) AIS patients with antimicrobials a total of 323 (46.4%) patients were exposed to 109 different pDDIs. The most common pDDIs were Ciprofloxacin-Diclofenac (16.09% of patients), Diclofenac-Levofloxacin (10.20%) and Aspirin-Levofloxacin (9.05%). The most common contraindicated pDDIs was Ceftriaxone-Ringer Solution (6.90%). Fatal outcome was more frequent ($p<0.01$) in the group of AIS patients (43.7%/28.6%) who were exposed to pDDIs which include antimicrobials. Binary logistic regression showed that gender ($p<0.01$, $B=0.623$, 95% CI 0.439–0.884) and the number of prescribed drugs ($p<0.01$, $B=1,255$, 95% CI 1.203–1.310) were significant factors associated with his pDDIs in AIS patients.

Conclusion: A total of 46.4% of patients with AIS stroke were exposed to pDDIs which include antimicrobials and fatal outcome was more prevalent in group of AIS patients with this pDDIs. Gender and number of prescribed drugs were significant factors associated with pDDIs which include antimicrobials in AIS patients.

Disclosure: Nothing to disclose.

EPO-494

Abstract withdrawn

EPO-495

Abstract withdrawn

EPO-496

The effectiveness of anticoagulant therapy in COVID-19 associated ischemic stroke

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Background and aims: Study was aimed to study the effect of various anticoagulant agents used in COVID-19 associated ischemic stroke on hemorheological parameters.

Methods: 62 patients with COVID-19-associated acute ischemic stroke were selected for the study. These patients ($n=62$) were divided into three groups. In the group A, $n=33$ (53.12%) patients received heparin for 2 weeks at 24,000–36,000 ED per day, $n=17$ (27.4%) patients in the group B received enoxiparin 1mg/kg/day for 2 weeks, and group C $n=12$ (19.4%) patients received rivaroxaban 15–20 mg per day for 2 weeks.

Results: As a result of anticoagulant therapy in groups, the hemorheological parameters (D-dimer, fibrinogen, prothrombin time, APTT) were regressed in the groups A, B and C of patients in the following order: D-dimer from 581.4 ± 1.6 ng/ml to 334.8 ± 2.1 ng/ml; from 628.6 ± 1.4 ng/ml to 336.7 ± 2.3 ng/ml; from 541.1 ± 1.9 ng/ml to 496.6 ± 1.4 ng/ml, fibrin degradation products from 7.71 ± 1.1 μ g/ml to 3.6 ± 1.3 μ g/ml; from 7.42 ± 0.9 μ g/ml to 3.8 ± 1.19 μ g/ml, from 7.52 ± 1.2 μ g/ml to 3.71 ± 1.3 μ g/ml, prothrombin time from 15.2 ± 1.1 sec to 9.4 ± 0.8 sec; from 14.9 ± 1.1 sec to 9.6 ± 0.8 sec; from 15.6 ± 1.1 sec to 9.2 ± 0.8 sec, APTT from 31.51 ± 1.29 sec to 24.16 ± 0.8 sec; from 28.2 ± 1.71 sec to 26.9 ± 1.65 sec; from 29.76 ± 1.13 sec to 25.21 ± 1.26 sec; (respectively, $p<0.001$).

Conclusion: All anticoagulants have a significant positive effect on fibrinogen and prothrombin time, heparin and enoxiparin are effective against D-dimer, heparin and riboraxaban are effective against APTT. However, riboraxaban has almost no positive effect on D-dimer while enoxiparin has almost no positive effect on APTT.

Disclosure: Nothing to disclose.