

P1958

Activation of Fibrinolysis System at the Main Arterial and Venous Vessels Thrombosis with the Treatment of Gene-Engineering Thrombolytic Drug 'Triasa'Kruchinsky, N¹, Plenina, L², Akulich, N¹, Tepljakov, A¹, Tsymanovich, S²¹Research Institute for Ecopathology and Occupational Diseases, Mogilev, Republic of Belarus²'Dialeck Ltd.', Minsk, Republic of Belarus

Goal: The purpose of the present research was studying safety and efficiency domestic gene-engineering Drug for thrombolysis 'Triasa' at carrying out of clinical tests (GCP) at 20 patients with thromboses main (arterial and venous) vessels.

Objectives and methods: 'Triasa' represents the albuminous complex (active substance – the enzyme triasa) received and cleared from cell-culture of a liquid imperfect saprophytic of mushroom 'Trichothecium rozeum'. This Drug for thrombolysis possesses activated fibrinolytic, thrombolytic and anti-inflammatory action, plasminogen activated, transforming it in plasmin, and renders direct fibrinolytic action. Prior to the beginning of carrying out of clinical tests of preparation 'Triasa' a series of experiments *in vitro* with use of blood of 22 volunteers for revealing an optimum doze of 'Triasa' has been lead. At each volunteer got on 50.0 ml of blood with subsequent its division on 10.0 ml, in which the 'Triasa' in various (3, 15, 150, 300 and 600 U/ml) concentration was added. Haemostasis System state research was spent before addition the drug (an initial condition), in 30 and 60 minutes after its addition in blood. The haemostasis was studied using evaluation of the platelet aggregation, coagulation, anticoagulant and fibrinolysis activity (25 parameters).

Results: The lead research *in vitro* has shown, that the optimal concentration of a preparation 'Triasa' for achievement of a comprehensible level to the fibrinolytic activity, not resulting in failure haemostasis imbalance it is possible to count 150 U/ml. The lead researches have shown good bearableness of 'Triasa' without the complications. 'Triasa' infusion in a doze of 200 000–500 000 U was promoted knocking over of hypercoagulation (statistically reliable APTT, decrease in fibrinogen concentration and D-dimers) state and fibrinolytic potential (statistically reliable of euglobulin fibrinolysis) activation.

Conclusion: Clinical tests (GSP) of 'Triasa' can be continued at a heart attack of a myocardium.

Aardestrup, IV, P1187
 Aaron, L, P1235
 Aarsman, CJ, OR327
 Aaspöllu, A, P1158
 Aballi, S, P1661
 Abashidze, A, OR139
 Abbadessa, V, P1398
 Abbate, R, OR128, OR131, P0368,
 P0456, P0479, P0496,
 P0510, P0658, P1026,
 P1095, P1123, P1124,
 P1254, P1282, P1349,
 P1563, P1564, P1603,
 P1624, P1645, P1649,
 P1678, P1684A, P1720,
 P1721, P1723, P1773,
 P1940, P2218, P2271,
 P2325
 Abbühl, B, P1231
 Abdalla, D, P0675
 Abdel Gader, A, P1949
 Abdel Hafez, A, P1868
 Abdelhaleem, M, OR360, P1519
 Abdelrahman, N, P1094
 Abdolreza, A, P1746
 Abe, Y, OR032, P0947, P2371
 Abell, L, , P0162
 Abendschein, D, OR077
 Aberbach, I, P1687
 Åberg, M, OR038
 Abeyama, K, OR170, P2382
 Abgrall, JF, P1268, P1269, P1637
 Abid Hussein, MN, P0573, P1753
 Aboelfetoh, N, P0582
 Aboud, M, P0616, P1129, P1321,
 P1376, P1983
 Abrahamsen, B, OR235
 Abrams, Z, P2024, P2112
 Abshire, T, P2029
 Abshire, TC, P0260
 Abu-Ghoush, M, P2042
 Abumiya, T, P0750
 Abu-Samra, H, P1312
 Abuzenadah, AM, P0465
 ACC members, P1711
 Accardo, PA, P0023, P1811
 Achiron, A, P1277
 Acosta, S, P1066
 Adachi, E, P2088
 Adachi, H, OR160
 Adachi, T, P1488, P1916
 Adam, M, P1813
 Adamczuk, Y, P0614, P1806
 Adamek, L, P1080
 Adams, M, P1341

Adams, TE, OR249
 Adamson, PJ, OR022
 Adamtziki, E, P0526, P1475, P2058,
 P2315
 Ádány, R, OR123
 Adcock, DM, P0711, P1901
 Adelmeijer, J, OR256, P0281, P0336
 Adess, M, P1911
 Adraktas, Th, P1050
 Aerts, JMFG, P1163
 Afrasiabi, AR, P2323
 Afuwape, A, OR137
 Agarwal, P, P2138
 Ageenkova, EV, P0236
 Ageno, W, OR175, OR311, P0450,
 P0451, P1007, P1052,
 P1630, P1648, P1650,
 P1869, P1877, P2188
 P2386
 Aggarwal, A, P1560
 Aggarwal, MA, P1273
 Aggio, M, OR175, OR273, P0395,
 P0450, P1017, P1051,
 P1627, P1643, P1646,
 P1650, P2188, P2202
 P2041
 Agnes, R, OR154
 Agneta, S, P1180
 Agostoni, P, OR384
 Ågren, A, OR385
 Aguilar, F, P0244
 Aguzzi, A, P0069
 Ahlenc-Gelas, M, P0468
 Ahlgren, A, P1270
 Ahluwalia, J, P0913
 Ahmad, F, OR372, P0913, P0982,
 P1514, P2105
 Ahmad, S, P0379
 Ahmad, SS, P0917, P0919, P0999,
 P1763
 Ahn, E, OR286
 Ahn, H, P1679
 Ahn, JY, P0917, P0919, P0999
 Ahn, Y, P1763
 Ahn, YS, P2144
 Ahnadi, CE, OR338, P1122
 Ahrens, I, P1745
 Ahrens, IG, OR241, P1480
 Ai, J, P1323, P1875, P1914
 Aiach, M, OR317
 Aillaud, MF, P0726, P1236, P2280
 Aiosa, A, P2361
 Aird, WC, P1385, P1386, P1980
 Aisina, RB, P0335, P2325
 Aiso, S, P1676
 Aixalá, M,

- | | | | |
|--------------------|--|---------------------------|---|
| Aizawa, K, | PLEN02 | Alessandrello, Liotta, A, | OR131, P0479, P2218, P2325 |
| Ajzenberg, N, | P0364 | Alessandro, G, | P1415 |
| Ajzner, E, | OR035 | Alessi, MC, | P0169, P0411, P1382, P1807,
P2369, SYM41 |
| Ajzner, É, | OR123, P1311 | Aletti, G, | P0863 |
| Akakura, N, | P2339 | Alewaeters, A, | P0876 |
| Akbar, H, | OR117 | Alexander, S, | P1359 |
| Akbulut, A, | P1909 | Alexander, WS, | SYM04 |
| Akhavan, S, | P1446, P1931 | Alexandre, MA, | P1452 |
| Akinwunmi, F, | P0429 | Alfaro, G, | P1275 |
| Akira, Y, | OR185 | Alguel, G, | OR214 |
| Akkerman, JWN, | OR013, OR051, P0298,
P0433, P0987, P1578,
P1583, P2090 | Alhaq, A, | P1451 |
| Aksoy, H, | P0899 | Alhenc Gelas, M, | P0703, P1323, P1688 |
| Aktas, B, | SOA01 | Ali, AN, | P1950 |
| Akulich, N, | P1958 | Ali, S, | P0635 |
| Al Abri, QN, | P2117 | Alieva, El, | P0236 |
| Al Askalany, H, | P0675 | Alikhan, R, | OR275 |
| Al Dieri, R, | P0422, P0703, P0889,
P1483, P1688 | Alisievitch, S, | P0483 |
| Al Haddabi, H, | P2117 | Aliyeva-Karayeva, N, | P0532 |
| Al Hajzen, A, | P2296 | Alizade, I, | P0532 |
| Al Kindi, S, | P1134, P2117 | Alizadeh, H, | P1001 |
| Al Lamki, N, | P1134 | Aljamali, MN, | H07 |
| Al Lamki, Z, | P2117 | Allaouchiche, B, | P0696 |
| Al Maarof, M, | P1134 | Allard, DA, | SYM13 |
| Ala, F, | P1389 | Allegrone, J, | P1699 |
| Ala, FA, | P1412 | Allemand, E, | P1522 |
| Aladle, D, | P2368 | Allen, D, | P0985, P1542 |
| Alatri, A, | P1707 | Allen, DL, | OR358 |
| Alaveras, M, | P0284 | Allen, G, | OR381, P0428, P1116, P1714 |
| Alban, S, | P0422, P0548, P1591 | Alliaume, N, | P1409 |
| Albanèse, J, | P1322 | Almaas, V, | P1661 |
| Alberca, I, | OR003 | Almasy, L, | P0072, P1388, P1974 |
| Alberio, L, | OR291, P0377, P0915,
P1574, P2171, SOA33 | Al-Mazedi, M, | P0161 |
| Albert, FW, | P0367 | Al-Mondhiry, HA, | OR109 |
| Albert, LM, | P1287 | Almonte, T, | P0228, P0807, P1997 |
| Albert, T, | OR208, P1829 | AL-Mugeirin, M, | P1949 |
| Albertazzi, L, | P0023, P1811 | Almus-Jacobs, F, | P2174 |
| Albertini, P, | P1449 | Al-Mutairi, M, | P0500 |
| Alberto, MF, | P0615, P1676, P2085 | Alonso, B, | P1031 |
| Albertson, T, | P1955 | Al-Rasheed, S, | P1949 |
| Albinni, S, | P0259 | Al-Salloum, A, | P1949 |
| Albisetti, M, | P0671, P1068 | Alsbjörn, B, | P0683 |
| Albisinni, R, | P0412, P1428, P1440 | Al-Tajer, S, | P1001 |
| Albrecht, C, | OR121 | Altavilla, E, | P1454 |
| Al-Buhairan, A, | P0865 | Altenburg, J, | P1332 |
| Alcalay, A, | OR056 | Altinbas, M, | P1002 |
| Alcaraz, MS, | P1206 | Altomare, C, | P1931 |
| Alcindor, T, | P1543 | Altomare, M, | P1593 |
| Alecrim, MCG, | P1452 | Altomonte, F, | P0257 |
| Aledort, L, | P0940 | Altomonte, M, | P0257 |
| Aledort, LM, | P0824, P1443 | Altuntas, F, | P0319 |
| Alentorn, F, | P1484 | Alvarez, L, | P0666, P1266 |
| Alessandrello, LA, | P1349 | Alvim, TC, | P0487 |
| | | Amagai, N, | P0784 |
| | | Amand, R, | P0051 |
| | | Amano, K, | P0800, P1832 |

- | | | | |
|------------------------|----------------------|--------------------------|----------------------|
| Amano, M, | P2247 | Andrade, LEC, | P1131, P2321 |
| Amant, C, | P0506 | Andrade, M, | P1953 |
| Amaral, A, | P0432 | Andre, P, | P1560 |
| Amaral, M, | P0276, P1003, P1473 | Andrea, L, | OR185 |
| Amaral, MM, | P0388 | Andreenko, G, | P0575 |
| Amaral, RQ, | P1131 | Andreesen, R, | P0593 |
| Amarillo, H, | P1964 | Andrei-Selmer, LC, | P2113 |
| Amato, G, | P0726, P2280 | Andreozzi, GM, | P0454, P1618, P1722 |
| Amaya, A, | P2105 | Andrews, C, | P0033, P0264 |
| Amaya, M, | OR287, OR372, P0982, | Andrews, R, | OR067, P0320, P0894, |
| | P2154 | | P1490, SYM18, |
| Amblo, J, | P0650 | Andriani, O, | P0090 |
| Ambo, H, | P0280 | Andrioti, E, | P1367, P1889 |
| Ambrosi, P, | P2285 | Angchaisuksiri, P, | P1400 |
| Ambrosio, GB, | OR175 | Angelillo-Scherrer, A, | OR233, OR368, |
| Amelot, A, | P0055 | | P1164 |
| Amengual, O, | P0024 | Angeloni, F, | P0692 |
| Ames, PRJ, | P0623, P1881 | Angerhaus, D, | P0278 |
| Amighi, J, | OR264 | Angstwurm, MWA, | P1991 |
| Amini, SH, | P0067, P1730 | Anguissola, GB, | P1180 |
| Amini-Bavil-Olyaei, S, | P2005 | Annerstedt, M, | P1375 |
| Amiral, J, | P0096, P0633, P1009, | Annichino-Bizzacchi, JM, | OR300, P1518 |
| | P1217, P1808 | Ansell, J, | P2336 |
| Amirkhosravi, A, | OR287, OR372, P0982, | Ansong, CA, | P1247 |
| | P2015, P2105, P2154 | Anthony, JC, | P1580 |
| Amlot, N, | P0471, P1666, P2262 | Anton, N, | P1726 |
| Ämmälä, P, | P2214 | Antonchuk, J, | SYM04 |
| Ammar, S, | P1097 | Antonijevic, N, | P1682 |
| Amore, C, | OR040, P1592 | Antonio, FS, | P1415 |
| Ampaiwan, C, | P0253, P0660, P1084 | Antonucci, E, | P0456, P1026, P1624, |
| An, MO, | P1081 | | P1645, P2218 |
| An, S, | P1379 | Antovic, A, | P1894, P1960 |
| An, SSA, | OR020 | Antovic, JP, | P1960 |
| Anagnostopoulos, A, | P0671 | Antrilli, TM, | P0770 |
| Anannit, V, | P1084 | Antunes, M, | P0223 |
| Ananyeva, N, | P0646, P1840 | Anucha, T, | P0253 |
| Ananyeva, NM, | P0376, P1242 | Anwar, R, | P1389 |
| Anastasio, R, | OR176, P0527, P1278, | Anzej, S, | P1894 |
| | P1398, P1706 | Apergis, G, | P1367 |
| Anastasopoulou, I, | P1050, P1937, P2023 | Appleby, JA, | OR347, P1528 |
| Ancona, E, | P0610 | Aprico, A, | P1490 |
| Ancora, MA, | P1169 | Arai, M, | P1256, P1832 |
| Andersen, CB, | P0562 | Aranda, E, | P0846, P1445, P1956 |
| Andersen, JC, | P1473A | Araújo, AG, | P2054 |
| Andersen, L, | P0095, P0810 | Araújo, F, | P0223 |
| Andersen, MD, | OR076 | Arbasi, C, | P1449 |
| Andersen, O, | OR248, P0646 | Arbeille, B, | P2126 |
| Andersen, TO, | P0492 | Arbring, K, | P1258, P2221 |
| Anderson, FA, Jr., | OR274, P1078, P1099, | Arcari, A, | P1776 |
| | P1590, P1643, P1671, | Arcas, I, , | OR234 |
| | P1673 | Aref, S, , | P0675, P1868 |
| Anderson, PJ, | P0119, P0882 | Arena, A, | P0412 |
| Anderson, S, | P2357 | Arey, BJ, | P0197, P2183 |
| Andersson, E, | P1769 | Arfin, M, | P0500 |
| Andersson, M, | P0178, P0643, P1627 | Arias-Salgado, EG, | SYM16 |
| Andersson, S, | P1114 | Ariens, R, | OR024, P1381 |
| Andrade Ferreira, I, | OR051, P0298, P0433 | Ariëns, RAS, | OR021, P1986, SYM51 |
| Andrade, CN, | P0419 | Ariens, RAS, | OR022, P1950 |

- | | | | |
|--------------------|---|---------------------|---|
| Ariga, T, | OR223, P0767 | Attanasio, M, | P1349, P1678, P1940 |
| Arikan, E, | P0586 | Attias, D, | P1602 |
| Arkin, C, | P1903 | Auboyer, C, | P0243 |
| Arkin, S, | P1443 | Aubry, P, | P0364 |
| Arit, M, | P0256 | Auerswald, G, | P0620, P1231, P1447,
P1640, P2022, P2062 |
| Armari-Alla, C, | P0847 | Auger, JM, | OR118, OR254, P0290
P0301 |
| Armour, K, | OR218 | Auger, WA, | P2266 |
| Arnal, JF, | OR010 | Aujesky, D, | P0470 |
| Arnan, M, | P0435, P1034 | Aung, MT, | OR333 |
| Arnaud, C, | P1267 | Auradé, F, | OR165 |
| Arndt, B, | P0288 | Aurousseau, MH, | P1215, P1216 |
| Arnesen, H, | P1029, P1778 | Austin, C, | P2228 |
| Arnold Spek, C, | SYM03 | Austin, H, | P0649, P1280, P1698,
P1926 |
| Arnold, D, | P0228 | Austin, RC, | P1344 |
| Arnout, J, | OR194, P0467, P0670,
P1065, P1776, P1861 | Austin, S, | P2099 |
| Arnuti, B, | P0866 | Auteri, A, | P1783 |
| Arocas, V, | OR071, P1904 | Authi, KS, | P0903 |
| Aronis, S, | P0526, P1410, P1475,
P2315 | Avcin, T, | P1215, P1216 |
| Arruda, VR, | OR047, OR300 | Averell, PM, | P11, P2379 |
| Arsenault, L, | P1634 | Avraamides, C, | P2366 |
| Arshinov, AV, | P1222 | Axelsson, T, | P1357 |
| Arteaga, R, | P1763 | Ayata, M, | P2247 |
| Arthur, J, | OR067, P0320, P0894,
P1490, SYM18 | Aydogan, G, | P1909 |
| Artmann, A, | P1686 | Ayer, M, | P1468 |
| Arya, R, | OR212, OR266, P0471,
P1069, P1666, P2262,
P2263 | Aygoren-Pursun, E, | P0602, P1204 |
| Aryurachai, K, | P2119 | Ayman, S, | P0903 |
| Arza, B, | P0285, P0467, P1064,
P1065 | Ayob, Y, | P1833 |
| Asada, Y, | P1178, P1586, P2287 | Azcan, S, | P0586, P0590 |
| Asai, F, | P1109, P1111, P1708,
P2299, P2301 | Azibi, K, | P0921 |
| Asakura, H, | OR351, P1313, P2377 | Azizova, O, | P1380 |
| Asazuma, N, | P2086, SOA27 | Aznar, J, | P0730, P0731, P1329 |
| Aschermann, M, | P2212 | Aznar, JA, | P0282, P0871, P1864,
P2013, P2073 |
| Asencio, JG, | P0266 | Baba, Y, | P0800 |
| Aseychev, A, | P1380 | Babinska, A, | P0339, P0409, P2308 |
| Ashley-Koch, A, | P0073 | Babyn, P, | P0227 |
| Ashmarin, I, | P0291 | Bach, J, | P0036, P1755, P1859 |
| Ashmelash, G, | P2019 | Bacher, P, | P2194 |
| Ashrani, AA, | P0823, P0916 | Bacheschi, L, | P2195 |
| Ashton, F, | P1694 | Bachoo, P, | P1140 |
| Asmalash, G, | P1218, P1219, P1338,
P1809, P2001 | Badenhorst, PN, | P0012 |
| Assafim, MIA, | P0140 | Bader, R, | P0215, P2059 |
| Asselta, R, | OR126, P2039 | Badereddine, L, | P1403 |
| Asti, D, | OR293 | Badimon, L, | P0156 |
| Ataullakhanov, F, | P0976 | Badlou, BA, | P1583 |
| Ataullakhanov, FI, | P0376, P0613, P1242 | Badol, P, | P1492 |
| Atichartakarn, V, | P1400 | Baert, J, | OR182 |
| Atoda, H, | P1255 | Baffour, R, | P2386 |
| Atsma, DE, | P1391 | Bagan-Triquenot, A, | P0715 |
| Atsumi, G, | P1325 | Baggio, MS, | P2055, P2235 |
| Atsumi, T, | P0024 | Baghaei, F, | P0872, P1243 |
| | | Baghaipoor, MR, | P0324, P1412 |
| | | Baghli, L, | P1403 |
| | | Baglia, FA, | OR097, OR348 |

Baglin, T,	OR004	Bansal, V,	P1100, P1288, P1781
Bagot, C,	P2263	Banzet, N,	P2285
Bahnson, BJ,	H10	Bao, C-X,	P1300, P1893
Bahou, W,	OR378	Baomin, X,	P0162
Bahou, WF,	OR221	Barabás, É,	P1396
Bahr, W,	P1638, P1639	Barakat, W,	P0039
Bahram, Z,	P1746	Baranska, P,	P0555
Bahri, A,	P1097	Barbanti, M,	P0569, P1167
Bai, X,	OR133, P0315, P0391, P0579, P0855, P1153, P1507, P1509, P1946, P2344, P2136	Barbault, S,	P1719
Bai, Y,	P1873	Barbe, MF,	P1155
Baik, N,	OR326	Barber, HK,	OR120
Bailey, KR,	P0477, P1076	Barbui, T,	OR039, P1220, P1906, P2116, SYM12
Baird, TR,	OR374, OR377, P1822	Barbulescu, DM,	P1825
Bajaj, SP,	OR341, OR342	Barden, A,	P0948
Bajc, M,	P1664	Bardet, V,	P0921
Bajusz, S,	P1396	Bardin, N,	P1789, P1807
Bajzar, L,	OR396	Bárdos, H,	OR123
Baker, J,	OR068	Bareford, D,	P0160
Baker, R,	P0092, P0123, P0957	Barek, L,	P1719
Baker, RI,	P1846	Barellier, MT,	P0069
Bækgaard, N,	P0444, P2210	Bari, S,	P2085
Bakhtiari, K,	OR031	Barillari, G,	P0218, P0250, P0850
Baklan, TN,	P1035	Barker, RN,	P0026
Bal dit Sollier, C,	P0363, P0426, P1622, P1623, P2196	Barnard, D,	P0208, P0808
Balark, K,	P2029	Barnard, MR,	OR138, P0357, P0361, P0954, P0955
Balbale, S,	P1911	Barnes, C,	OR357, P0124, P0805, P1913
Balci, D,	P0590	Barnes, E,	P0630
Baldan, N,	P0610	Barnes, KD,	P1502
Baldi, G,	P0257	Barni, S,	OR060
Baldi, M,	P1684A	Baron, DW,	P1194
Baldoni, M,	P2219	Baronciani, L,	OR281
Balducci, D,	OR039, P2116	Baroni, M,	P1914
Balduini, A,	OR220	Barral, FG,	P2313
Balduini, CL,	OR220, P2115	Barrellier, MT,	P0667
Balgojevich, E,	P1649	Barrett, B,	P1404
Ball, D,	P2332, P2333	Barrett, NE,	P1541
Ballarino, GJ,	P2320	Barro, C,	P0266, P0847
Balmer, C,	P0671	Barrow, RT,	OR046
Balogh, E,	P0774	Barrowcliffe, TW, ,	P0721, P1246, P1432
Balogh, I,	OR123, P0323, P1787,	Barsom, A,	P0949
Baltrukovich, N,	P0337	Barsukov, A,	P1183
Banait, V,	P0676	Barteling, W,	P0043
Bandinelli, B,	P0479, P1564, P1678	Baruch, D,	P0333, P0364, P0877, P1540, P0678
Bandinelli, S,	OR128, P1282	Bashiruddin, J,	P1027
Banerjee, P,	P0339	Basileo, M,	P1558, P2279
Banerjee, Y,	P0154	Basili, S,	P1710
Bang, KWA,	OR288, P2143	Baskova,,	P2114
Bank, I,	P0929, P1056	Bass, H,	P2167
Banks, J,	P1089	Bassi, A,	P2195
Bannink, L,	P1653	Bassitt, R,	P0199
Banno, F,	OR026	Bassitte, RP,	OR321, OR332, OR338, P1122, P1547, P1745
Bannon, PG,	P1146	Bassler, N,	, P0345, P0697, P0812
Bansal, K,	P1839	Bassus, S,	

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|------------------|---|----------------------|--------------------------------------|
| Basta, G, | P0749 | Beckmann, MW, | P2277 |
| Bastelica, D, | P1382, P2369 | Beconcini, S, | P0368, P0658 |
| Bastida, E, | P1430 | Becton, DL, | P0246, P1436 |
| Bastien, O, | P0572 | Bedard, P, | OR135, P2355 |
| Bastos, M, | P0745 | Bedard, PW, | OR390 |
| Batchelor, AJ, | P0474, P1070 | Bedoya, G, | P0666, P1266 |
| Bates, E, | P0371, P0372 | Beeton, L, | P0587 |
| Bates, I, | P0429 | Begley, GS, | OR189 |
| Bates, SM, | P0127, P2237 | Begtrup, K, | P1363, P1934 |
| Bath, PMW, | P2373 | Béguin, S, | P0422, P0703, P0889,
P1483, P1688 |
| Batista, IFC, | P0714 | Begum, S, | P0936 |
| Battle, A, | P2330 | Behan, MWH, | P0942 |
| Battle, J, | H08, H09, OR245,
OR278, OR279, OR281,
P0265, P0269, P0270,
P0271, P1469, P1626,
P2057, P2059, | Beharry, N, | P1750 |
| Batora, I, | P1632 | Behrmann, M, | OR045, P0035 |
| Batorova, A, | P0184, P0252, P0852,
P1447, P1632 | Beijnen, JH, | OR240 |
| Battaglioli, T, | P0076, P0700 | Bein, G, | P0910 |
| Battistella, PA, | P1604 | Belik, J, | P1456 |
| Battiston, M, | P0294, P1539 | Bellatorre, AG, | P1677 |
| Battle, J, | P0865, P1461 | Bellein, V, | P0432 |
| Battler, A, | P2386 | Belli, S, | P1031 |
| Baudo, F, | OR036, P0537, P1477 | Bellia, M, | P1367, P1889 |
| Baudry, O, | P1371 | Bello, T, | P2045 |
| Bauduer, F, | P1312 | Bellucci, S, | P0363, P0921, P0922,
P2056 |
| Bauer, JS, | P0770 | Beltran, J, | P1296 |
| Bauer, K, | OR063, P1046 | Belvedere, M, | P0417 |
| Bauer, KA, | OR153, P2216 | Ben Lassoued, MA, | P1097 |
| Bauer, M, | P1751 | Ben Romdhane, N, | P1097 |
| Bauersachs, J, | OR259, P0289, P1487,
P2083 | Benavente-García, O, | P0354 |
| Bauersachs, R, | P1024, P2241 | Bendas, G, | P0548 |
| Bauersachs, RM, | OR211 | Bender, HG, | OR215, P1685, P2127,
P2277 |
| Bauman, J, | P0430 | Bender, T, | P0302 |
| Bauman, M, | P1728, P2322 | Bendezú, K, | P1031 |
| Baumbach, GA, | P1370 | Benegas, M, | P2320 |
| Baumgartner, B, | P0723 | Beneke, H, | P1324 |
| Bauters, A, | OR317 | Benfield, C, | P0873 |
| Baxi, SM, | P1213 | Benhadda, B, | P1403 |
| Baxter, J, | P2340 | Benísvy, D, | OR143 |
| Bazzan, M, | OR175, P0450,
P2188 | Bénit, L, | OR163, P0322 |
| Beard, M, | P0804 | Benjamin, RJ, | OR268 |
| Beardsley, DS, | P1511 | Bennaghmouch, A, | P0380, P1350, P1489 |
| Beaudin, S, | P0128, P0129 | Bennett, CE, | OR314 |
| Beaulieu, JF, | P1865 | Bennett, JS, | OR104, OR107 |
| Beaulieu, LM, | P1333 | Bennett, M, | P0859, P1343, P1517 |
| Beaune, P, | OR085 | Bennett, MR, | SYM13 |
| Beauregard, P, | P1543 | Ben-Shlomo, Y, | P0681 |
| Becanovic, J, | OR116 | Benson, E, | P0050 |
| Becattini, C, | OR175, P0395, P0450,
P2202 | Benson, J, | P1926 |
| Beck, DO, | P1923 | Beránek, M, | P1262 |
| Becka, M, | P1105, P1704 | Berber, E, | P0690 |
| Becker, S, | P0021, P0035 | Berckmans, RJ, | P1777 |
| | | Bercovich, D, | OR216 |
| | | Bereczky, Z, | OR123, P0774, P0926 |
| | | Béres, BJ, | P1093 |
| | | Berger, F, | P1068 |
| | | Berger, K, | P0209 |

- | | | | |
|-----------------|--|----------------------|---|
| Berger, M, | P1263 | Beurrier, P, | P1426 |
| Berger, U, | P1526 | Bevan, DH, | P0853 |
| Bergman, GE, | P0835 | Beveridge, M, | P1774 |
| Bergmann, F, | P0022 | Bevers, E, | OR345, P0380 |
| Bergmann, JF, | OR274, P1622, P1671,
P1673, P2196 | Beyzadeoglu, H, | P1467 |
| Bergmeier, W, | OR199, P0288 | Bezaf-Bouchahda, C, | P0608 |
| Bergmeister, H, | OR387 | Bezemer, ID, | P1862 |
| Bergqvist, D, | OR057, P1051, P1066,
P1738, SYM38 | Bezemer, M, | OR013, P1578 |
| Beris, A, | P1642 | Bezerra-Neto, JE, | P0199 |
| Berk, BC, | SYM46 | Bhagwanbali, V, | OR193 |
| Berland, Y, | P0597 | Bhakta, V, | P1200 |
| Bermejo, E, | P2085 | Bhatt, DL, | P0974 |
| Bermúdez, MM, | P1797 | Bhimavarapu, A, | P0647 |
| Bernard, A, | P1092 | Bhoopalam, N, | P1911 |
| Bernard, M, | P0004 | Bialonczyk, C, | OR361, P0438, P0581,
P0743, P1040 |
| Bernardi, E, | P1618, P1647 | Bianchi, M, | P1648 |
| Bernardi, F, | OR156, P0638, P0639,
P1831, P1914, P1939,
P1240 | Bianchini, EP, | OR251, P0055 |
| Bernardi, M, | P0271, P0870 | Biasini-Rebaioli, C, | P1215 |
| Bernardo, F, | P0510 | Biasoli, C, | P1449 |
| Bernasconi, A, | P1676 | Bick, R, | P0782, P2194 |
| Bernasconi, S, | OR207, P0037 | Bick, RL, | P1899 |
| Berndt, M, | OR067, P0894, P1490,
P2095, SOA27,
SYM18 | Bickeböller, H, | P1263 |
| Berndt, MC, | OR116, P2086 | Bickel, C, | P1756 |
| Berneman, Z, | OR301, P0272 | Bidligmaier, C, | OR356 |
| Bernhard, N, | P0949 | Bidlingmaier, C, | P1601, P203 |
| Bernier, M, | P1042 | Bidot, C, | P0917, P0919, P0999 |
| Berntorp, E, | P0238, P0829, P0872,
P2259, SYM19 | Bièche, I, | P1875 |
| Berrou, J, | OR227 | Bieche, I, | P1323 |
| Berruyer, M, | P0174, P1312 | Biemek, C, | P0060 |
| Berry, LR, | OR083, P0094, P0600,
P0693, P1201, P1456,
P1969 | Bierhaus, A, | P0130 |
| Bertagnin, A, | P0454 | Bierings, R, | OR008, P1767 |
| Bertelsen, V, | P0619, P1345 | Biesert, L, | P0244, P0655, P0878,
P2068 |
| Bertina, RM, | OR110, OR237, OR315,
P0059, P0149, P0730,
P0731, P1125, P1238,
P1283, P2071 | Biesma, D, | P0195, P0461 |
| Bertocco, S, | P0588, P2188 | Biesma, DH, | P0463 |
| Bertoldi, A, | OR175, P0450 | Bifarini, B, | P0249 |
| Bertomoro, A, | OR282, P0862 | Bigi, P, | P1773 |
| Bertoncello, K, | P0271, P0870 | Biglioli, P, | P0358 |
| Bertoni, A, | P1493 | Biguzzi, E, | P0725, P1905, P2376 |
| Bertrand, J-R, | OR299 | Bihoreau, N, | P0695, P1970 |
| Bertrand, MA, | P0267 | Bijlsma, MF, | OR202 |
| Bertrand, N, | P1622 | Bijsterveld, NR, | P2203 |
| Bérubé, C, | P1042 | Bild, A, | P1818 |
| Bethel, M, | P0711, P1901 | Bilgrami, SA, | P2108 |
| Bettis, B, | P0028 | Bílková, J, | P2212 |
| Betts, L, | OR103 | Billett, HH, | P1121 |
| Betz, C, | P0311 | Billing Clason, S, | P0416 |
| | | Binder, BR, | OR227, OR270, OR387,
P0826, P0875, P1157,
P1160, P1751, P1885 |
| | | Biner, B, | P0590, P1467 |
| | | Binetti, BM, | OR293 |
| | | Biondi, ML, | P2376 |
| | | Birch, B, | P0159, P0762, P0794 |
| | | | P1838 |
| | | Bird, ML, | P1911 |
| | | Birkenes, B, | P1944 |

Biró, É, P0686, P2150
 Biron-Andréani, C, OR317
 Biron-Andreani, C, P0608, P1086
 Biss, TT, P0844
 Bissery, A, P1875
 Biswas, KK, OR170, P2382
 Bitsch, N, P1489
 Bittdorf, B, P0175
 Bjørkøy, G, P0151
 Bjørnarå, BT, P0472
 Björck, M, P1066
 Black, L, OR360
 Blackburn, J, P0430
 Blacklock, L, P2268
 Blackney, M, P1715
 Blackwell, S, P1473A
 Bladbjerg, EM, P1345
 Blagojevic, J, P1645
 Blagojevich, J, P1349
 Blair, P, P1504
 Blais, L, P0425
 Blais, N, P0425, P0441, P1042
 Blamey, CJ, H10, P0304
 Blanchette, V, P0227, P1425, P1837
 Blanchette, VS, P0208, P0229, P0808, P1519
 Blanco, A, P2046
 Blanco, AN, P0263, P0615
 Blangero, J, P0072, P1388, P1974
 Blankenberg, S, P1756
 Blann, A, P0763, P1951
 Blann, AD, P0160, P1154, P1175, P1176, P2153, P2192
 Blasbery, K, P0971
 Blatny, J, P1407
 Blatny, J, P1998, P2032
 Blauth, G, P0367
 Blalek, B, P1407
 Blazek, B, P1998
 Bledzka-Krupska, K, P0337, P0931
 Bleijlevens, B, P1163
 Bliden, K, P0362, P0374, P2167,
 Bliden, KP, P0414, P0517, P1014, P2205
 Blinc, A, P0989, P1589
 Blobel, C, P0549
 Block, K, P1924
 Blocklet, D, OR306
 Bloemendal, HJ, P0329
 Blom, H, H03, P1079
 Blom, HJ, OR127, OR129, P0480
 Blom, JW, P0462
 Blombäck, M, P0183, P1960
 Blomberg, A, P1092
 Blömer, L, P2351
 Blomstrand, C, OR198, P0178
 Blostein, M, P0533
 Blouch, MT, P1268, P1269, P1637

Blumenfeld, Z, P1687
 Blumenstein, M, P0262, P1459
 Bobe, R, P2133
 Boberg, P, P1243
 Bobik, A, H15, P1193
 Bobkova, I, P0575
 Bobrovník, S, P1880
 Boccalon, H, P1276
 Boccuzzi, SJ, P1865
 Bochenek, A, P0413
 Bock, PE, OR377
 Bock, SC, OR002, P0607
 Boczowska-Radziwon, B, P0031, P1392
 Boda, Z, OR035, P0926, P1311, P1731
 Boddi, M, P1603
 Bode, C, OR321, OR332, OR338, P1122, P1547, P1745
 Bode, W, SYM26
 Boden, G, OR052, P1572
 Bodenbender, W, P2076
 Bodin, L, OR085
 Bodnar, R, P2095
 Bodó, I, P1476
 Bodzenta-Lukaszyk, A, P1372
 Boehlen, F, P1223, P2161
 Boehm, B, P0036
 Boehm, M, P0191
 Boehncke, WH, P0548, P1591
 Boekholdt, SM, P0150
 Boekhorst, J, P0220, P0818, P1418
 Boer, K, P0959, P1182
 Boeri, E, OR091
 Boerop, J, OR054
 Boertjes, RC, OR162, OR340, P0040
 Boezeman, J, OR080
 Boffa, MC, P1215, P1216
 Boffard, KD, OR379
 Bogdanov, IG, P2044
 Boggio, L, P0121, P0545, P1226, P1845
 Bohec, C, P1789
 Böhm, M, P0311, P0778, P0912, P1005
 Bohn, RL, P0824
 Böing, A, P0194, P1182
 Boinot, C, OR317, P0921, P1215
 Boissonnat, P, P0572
 Boisson-Vidal, C, OR143
 Bokarev, I, P0484, P1663
 Bokarev, IN, P1083
 Boldrin, M, P0610
 Bolis, PF, P0451
 Bolli, P, P0456, P1563, P1720
 Bolosiu, HD, P0071
 Bomgaars, L, P0138
 Bonacini, N, P1811
 Bonar, R, P1129, P1983

- | | | | |
|-------------------------|---|-----------------------|---|
| Bonardo, B, | P1161 | Bouma, B, | OR098, OR327, OR337,
P0563 |
| Boncela, J, | OR224, P2367 | Bouma, BN, | OR393 |
| Boncler, M, | P0951 | Bounameaux, H, | OR140, OR144, OR233,
OR305, P0470, P1138,
P1770, P2239, P2260,
SYM55 |
| Bonduel, M, | P1018, P2050, SYM68 | Boutière-Albanèse, B, | P1322 |
| Boneu, B, | P1267, P1276 | Bouton, MC, | P1904 |
| Bongers, TN, | OR193, P0281 | Boutou, AK, | P2242 |
| Bonifacio, G, | OR176, P0527, P1278,
P1398, P1706 | Bouziane, K, | P0694 |
| Bonnefoy, A, | OR029, P0506, P2135 | Bouzourene, K, | P1173, P1770 |
| Bonner, L, | P1069 | Boval, B, | P0363, P0922, P2056 |
| Bonnet, JL, | P0411, P2285 | Bovenschen, N, | OR162, OR225 |
| Bonnet, M, | P1297 | Bovill, EG, | OR313, P0065, P1942 |
| Bono, F, | OR141, P1790 | Bowen, D, | P0874 |
| Bono, M, | P1210 | Bowen, DJ, | P2065 |
| Boomsma, F, | OR364 | Bowers, S, | P1023 |
| Boonman-de Winter, LJM, | P0539 | Bowman, K, | OR158 |
| Booth, NA, | P0787, P1369, P1961 | Boyer, J, | P2206 |
| Borawski, J, | P0754, P0780, P1945 | Boylan, B, | OR284 |
| Borel-Derlon, A, | P0267, P0277 | Boli, M c̃, | P1113 |
| Borensztajn, K, | SYM03 | Bozic, M, | P0989 |
| Borensztajn, KS, | OR202 | Bozzao, C, | P2039 |
| Borg, JY, | OR317, P0063, P0114,
P0277, P0715, P0909 | Bozzini, C, | OR156, P0638 |
| Borgel, D, | P1914 | Brabec, P, | P2032 |
| Borggreffe, M, | OR033, P0084,
P2158 | Brackmann, HH, | P1829 |
| Bork, K, | P0669, P1794 | Bradford, HN, | P1184, P2182 |
| Borm, G, | P0480 | Bradley, CS, | P0208, P0808 |
| Bornstain, C, | P1875 | Braeunlich, CG, | P0910 |
| Borowiec, J, | P1798 | Braganza, DM, | SYM13 |
| Borrell, M, | P0466 | Brancaccio, V, | P0623 |
| Borrelli, G, | P1220, P1906 | Brandão, L, | P1214 |
| Borris, LC, | OR062, P1867 | Brandao, L, | OR360, P2185 |
| Borroni, L, | P1477 | Brandão, LR, | P0260, P1132 |
| Bortoletto, E, | P0030, P0217 | Brandao, LR, | P1924 |
| Borup, R, | P0562 | Brandjes, D, | P0203 |
| Børvik, T, | P1695 | Brandjes, DP, | OR240 |
| Bos, G, | H03, P1079 | Brandjes, DPM, | P0685 |
| Bos, MHA, | P0040 | Brandslund, I, | P1729 |
| Bosari, S, | P0086 | Brandt, J, | OR303 |
| Bosma, K, | P0052 | Brandt, JT, | P0952, P0956 |
| Bosman, L, | P0102, P1892 | Brassard, J, | P0425 |
| Bosman, RJ, | P0314 | Braun, S, | P0099 |
| Bosmans, JM, | P2250 | Braun-Dullaesus, R, | OR386 |
| Bossi, A, | P1007 | Brauner, R, | P1998 |
| Bossuyt, P, | P1658 | Brautigan, PJ, | OR298 |
| Bostock, JA, | OR049, OR314 | Bredoux, R, | P2133 |
| Boström, SL, | P0142 | Breimo, E, | P1758, P1759 |
| Bostwick, J, | P0197 | Breimo, ES, | P1943 |
| Botchkov, V, | P1160 | Breindahl, M, | P1867 |
| Botikov, AG, | P1421 | Breit, SN, | P2137 |
| Bottaro, F, | P2230 | Breitner-Ruddock, S, | OR081 |
| Bottenus, R, | P0626, P1928 | Breitwieser, A, | P0875 |
| Bouhassira, E, | OR025 | Breña, J, | P2045 |
| Boukas, AE, | P2242 | Brenkel, I, | OR239 |
| Boukour, S, | OR163 | Brennan, M, | P0085, P1752 |
| Boulanger, A, | P1406, P1444 | Brennan, MP, | P1281 |
| Boulyguine, OY, | P1980 | | |

Brenner, B, OR216, P1348, P1687, P1938
 Bresser, P, P0753, P1062
 Bressollette, L, OR272, P0432, P0668, P1048
 Brett, S, OR380
 Bretthauer, N, P1705
 Breuss, J, P1751
 Breuss, JM, OR387
 Brian, B, OR185
 Bridey, F, P0266, P0267, P0277
 Briegel, J, P0005
 Brieger, D, P1017, P1699
 Brighton, T, P2100
 Brighton, TA, P2202
 Brigulla, M, OR081
 Brill, A, OR296, P1094, P2169, P2363
 Brill, G, P0929
 Brini, M, P0023, P1811
 Briones, M, P0260
 Brisset, AC, OR010
 Brito, D, OR207
 Britt, M, P1818
 Brittenden, J, P1140, P2332, P2333
 Bröcker-Vriends, AHJT, P2009
 Bro, S, P0562
 Brockhaus, W, P0849
 Brodde, MF, OR016, P0495, P1876
 Broderick, J, P1363, P1934
 Brodin, E, P0375, P1351, P1695, P1779
 Brogi, D, P0479, P1723
 Brohée, D, P0876
 Brohee, D, P0928
 Brohult, J, P1067
 Brojer, E, P0222
 Brom, J, OR059, P1021, P1022
 Bromberg, ME, P2366
 Broni, A cç, P2314
 Brons, P, P2159
 Brons, PPT, P0220
 Bronzi, C, P0264, P0297
 Brooks, JM, OR134
 Brooks, S, P1401
 Brooks, SV, P0795
 Brophy, M, P0688
 Brophy, MT, OR089
 Brophy, T, OR106
 Brosstad, F, P0375
 Brotman, DJ, OR309
 Brouillette, S, OR290
 Brouwer, JLP, OR213, P0066, P0491, P0663, P0664, P1058, P1090, P1265, P1284, P1855, P1857, P1858, P2200, P2282
 Browett, PJ, P1694

Brown, C, OR280, P0690, P1847
 Brown, K, P1910
 Brown, L, P1847
 Brown, MA, OR189
 Brown, RC, P1819
 Brown, S, OR116, P0239
 Brown, SA, OR092, P0795, P1401, P1471, P2067
 Brownlow, SL, P1497
 Brox, J, P1758, P1759
 Broze, G, OR190
 Broze, GJ, P0718
 Brozek, J, P2353
 Brueckmann, M, P0084
 Bruel, A, P0506
 Brugge, JM, P1240
 Brüggemann, LW, P0009, P1207, P1802, P2010
 Bruggeman, LW, OR202
 Brügger Andersen, T, P1696
 Brügger-Andersen, T, P0497
 Brummel, K, P0564
 Brummel-Ziedins, K, P2353
 Brummel-Ziedens, KE, OR373, P1142
 Brun, NC, P1363, P1934
 Brunet, A, P1766
 Brunet, P, P0597
 Bruneval, P, P2296
 Bruni, F, P1783
 Brunner, P, OR387, P1157
 Brunsch, H, OR386
 Brusasco, C, P0257
 Brusi, C, H02
 Brusky, RM, P2053
 Bryl, E, P0399, P0996
 Bryniarski, L, P0564
 Brynildsrud, T, P0704
 Bryszewska, M, P1631
 Brzezinska-Kolarz, B, P1012, P1835, P2258
 Brzosko, S, P2384
 Buccholz, N, P2075
 Bucciarelli, P, P0076, P0958, P1905
 Buchacher, A, P0139
 Buchanan, GR, P1000
 Buckland, RJ, P2148, P2162
 Buckley, CD, P0934
 Buczko, W, P2180
 Buczma, A, P1419
 Budde, U, H08, H09, OR245, OR278, OR279, OR281, P0265, P0269, P0270, P0271, P0278, P0211, P0849, P0865, P1461, P1469, P2057, P2058, P2059
 Budihna, MV, P1113
 Buehler, L, OR140
 Buelens, K, P1365

- | | | | |
|----------------|----------------------|-------------------|----------------------|
| Buensuceso, C, | SYM16 | Bylock, A, | P1627, P2209 |
| Bühler, B, | P0104 | Byrne, E, | P1271 |
| Buil, A, | P1974 | Byrne, S, | P1107 |
| Bulabois, CE, | P0004 | Cabiddu, M, | OR060 |
| Bulanov, A, | P0080 | Cabral, KMS, | OR005 |
| Bulíková, A, | P1407 | Cabrera, N, | P0282, P0871, P1864, |
| Bullano, MF, | P1742 | | P2013 |
| Büller, H, | P1112, P1654, P2248 | Cabric, S, | P1804 |
| Büller, HR, | OR031, OR066, OR149, | Cachier, A, | P0364 |
| | OR240, OR289, OR364, | Caciolli, S, | P0368 |
| | P0009, P0397, P0421, | Cadavid, AP, | P0666, P1266 |
| | P0459, P0460, P0481, | Çaglayan, H, | P1468 |
| | P0685, P0687, P1055, | Cai, H, | OR231 |
| | P1056, P1057, P1059, | Cai, XH, | P0045 |
| | P1596, P1657, P1856, | Cai, X-y, | P0757 |
| | P2203, P2251, P2254, | Caille, D, | OR111 |
| | P2256 | Caine, GJ, | P2192 |
| Buller, HR, | P0424, P1060, P1062, | Calabrese, F, | P0610 |
| | P1655, P1656, P1658, | Calabrò, G, | P0726, P1236, P2280 |
| | P2190, P2252 | Calaminus, SDJ, | P0301, P0901 |
| Bullinger, M, | P0209 | Calatzis, A, | P0104, P0962 |
| Bullorsky, E, | OR087, P0198, P2230 | Callas, PW, | OR313, P1942 |
| Bullorsy, EO, | P1964 | Calmel-Tareau, C, | P0055 |
| Bülrow, R, | P1187 | Calmuntia, M, | P0485 |
| Bultelle, F, | OR319, P1760 | Calo, D, | P1534 |
| Bumbal, J, | P2212 | Calori, G, | P2199 |
| Bunjevacki, G, | P2337 | Calvo, R, | P2023 |
| Buonanno, FS, | P1607 | Calzolari, F, | P1646 |
| Burfeind, P, | H12, OR262 | Camargo, E, | P2195 |
| Burggraaf, J, | P1689, P2281 | Cambien, F, | P1719 |
| Burghaus, B, | P1600 | Cambus, F, | P1623 |
| Burlacu, A, | P1196 | Cambus, JP, | P0426, P1623 |
| Burman, H, | P1839 | Camera, M, | P0352, P0358 |
| Burman, JF, | P0161, P0224, P0972, | Cameron, C, | OR280 |
| | P1306 | Cameron, LD, | P0065 |
| Burnand, K, | P0786, P1987 | Camilleri, RS, | P1741 |
| Burnand, KG, | OR137 | Camilloni, B, | P1027 |
| Burner, J, | P0319 | Camino-López, S, | P0156 |
| Burnier, L, | OR233, P1164 | Camire, RM, | H07, OR047, OR252, |
| Burnouf, T, | P0083 | | P1249 |
| Burns, T, | P1472 | Camoin-Jau, L, | P1322 |
| Burstein, C, | P1427 | Camp, P, | P1525 |
| Burysek, L, | OR201 | Campart, D, | P0552 |
| Bush, L, | OR048 | Campbell, D, | OR196 |
| Bussel, J, | P0940 | Campbell, DM, | P0493 |
| Bussel, JB, | P0317 | Campbell, J, | OR357, P0124, |
| Busson, J, | P1104 | | P1913 |
| Bussotti, M, | P1180 | Campbell, K, | OR358, P1524 |
| Butenas, S, | OR210, OR373, P0650, | Camporese, G, | P0454, P1618, P1722 |
| | P1142, P2028 | Campos, L, | P0310 |
| Butler, J, | P1700, P1884, P2101 | Campos, M, | P0223 |
| Butler, WE, | P1607 | Camps, S, | P2091 |
| Butt, W, | OR359, P0742 | Canault, M, | P1161 |
| Buurman, W, | P0010 | Canciani, M, | P0269 |
| Buyearu, N, | P0645 | Canciani, MT, | P2324 |
| Byams, VR, | P0258 | Candolfi, R, | P0100 |
| Bydlowski, SP, | P0308 | Canella, A, | P1831, P1939 |
| Bykowska, K, | P0078, P1871 | | |

Cannegieter, SC, OR289, P1814, P1657, P2256
 Cannizzaro, V, P0671, P1068
 Cannon, M, OR276
 Cannon, S, OR276
 Cano, H, OR003, P1206, P1796, P1797
 Canobbio, I, OR220, P1493
 Canova, C, P1138
 Cantillon, CE, OR089
 Canu, P, P1539
 Cao, JY, P1339A
 Caponi, C, P0395, P1650
 Cappelli, R, P0404, P0453, P0455
 Cappucci, F, OR094, OR197
 Cappuccio, F, P1966
 Cappuccio, FP, OR194, P0670, P1776, P1861
 Capraru, M, P1147
 Caprini, J, P0430, P1046
 Caprini, JA, P1667
 Caramori, ML, P0984
 Carandina, S, P0192
 Carballo, GA, P0863
 Carbillon, L, P1215
 Carbone, A, P0148
 Carbone, S, P0017, P1683
 Carcao, M, P1425
 Cardona, H, P0666, P1266
 Cardona-Maya, W, P1266
 Cardona-Maya, WD, P0666
 Cardoso, JE, P0439, P0487, P1628, P2223
 Carella, A, P0396
 Cariou, R, P0424
 Carlo, A, P1483
 Carlsson, A, P0142
 Carluccio, M, P1169
 Carluccio, MA, OR204, P1151, P1169
 Carmassi, F, P1979, P1992
 Carmeliet, P, OR233, OR368, P1164
 Carneiro, JDA, P0673, P0994
 Carnero, J, P1626
 Carol, H, OR185
 Carole Giraud, P1888
 Carollo, C, P1008
 Carolyn, W, P2228
 Caron, C, P0277, P0876, P1461, P2056, P2059
 Carotti, A, P1931
 Carpani, G, P2376
 Carpenedo, M, OR310, P1734, P2327
 Carpinelli, M, SYM04
 Carr, P, P0382
 Carroll, WE, P1310
 Carruthers, J, P0228
 Carter, AM, OR021, OR049, OR314, P0286, P2079, SYM64

Caruso, S, P0129
 Carvalho, MG, P0431, P0439, P0487, P0745, P1628, P1690, P2223
 Carville, D, P0371, P0372, P0766
 Casais, P, P0863, P1128, P1676, P2046
 Casaña, P, P0282, P0871, P1864, P2013, P2073
 Casanegra, AI, P2320
 Casetta, B, P1757
 Casey, CR, P0247
 Casey, G, P0247
 Casini, A, P0496, P1678, P1684A
 Casonato, A, OR282, P0862
 Castaman, G, H08, H09, OR126, OR245, OR278, OR279, OR281, P0070, P0265, P0269, P0270, P0271, P0865, P0870, P1461, P1469, P2006, P2039, P2040, P2057, P2059, P0666, P1266
 Castaneda, SA, OR023, OR297
 Castel, D, P2219
 Castellani, D, P1810
 Castelli, M, P2298
 Castelli, R, OR326
 Castellino, F, OR173
 Castellino, FJ, P1924
 Castellone, DD, OR194
 Castelnuevo, AD, P1155
 Casteneda, J, P0354
 Castillo, J, P1131
 Castillo, V, P1132, P1214
 Castle, D, P1240
 Castoldi, E, P1763
 Caswell, M, P0873
 Catania, V, P0778
 Catchpole, I, P1037
 Catozzi, L, P0192
 Cattaneo, A, P1052
 Cattaneo, M, H03, P0086, P0965, P1079, P2174, P2376
 Cattini, MG, OR282, P0862
 Caturla, J, P1968
 Cauchie, P, P0876, P0928
 Caulin, C, P1622, P2196
 Cauwenberghs, N, P0002
 Cauwenberghs, S, P0987, P1584, P2179
 Cavalheiro Filho, C, P0054, P0308, P0419, P0984, P1019, P1041, P2293
 Cazenave, JP, P0108, P0739
 Cazenave, J-P, OR255
 Cazzaniga, M, OR060, OR382
 Ceccarelli, C, H10

- | | | | |
|--------------------------|---|--------------------|--|
| Ceccarelli, E, | P0404 | Chamone, DAF, | P0054, P0199, P0308, |
| Cecchi, E, | P0456, P1026, P1624,
P1723, P2218 | | P0419, P0984, P0994,
P1019, P1041, P2293,
P1409 |
| Cécile, K, | P1521 | Chamouard, V, | P1409 |
| Cederholm, I, | P2078 | Chamyan, G, | P1460 |
| Cederholm-Williams, S, | P1976 | Chan Kwo Chion, A, | OR242 |
| Cedrone, A, | P0350 | Chan, A, | OR359, P0228, P0742,
P0858, P1132, P1342 |
| Cejna, M, | OR387 | Chan, AK, | OR355, P0654, P0807 |
| Celentano, A, | P0452 | Chan, AKC, | OR083, OR360, P0094,
P0442, P0600, P0693,
P1201, P1214, P1456,
P1969, P1997 |
| Celestini, A, | P1732 | | P1480A |
| Celik, I, | P2342 | Chan, Kwo, | P0839 |
| Celikel, R, | OR258 | Chan, R, | P0516, P1115, P1116,
P1715, P2305, P2306 |
| Celiska-Löwenhoff, M nç, | P1785 | Chander, C, | OR185 |
| Celinska-Löwenhoff, M, | P0482, P2353 | Chandler, A, | OR103 |
| Celinska-Lowenhoff, M, | P0779 | Chandler, B, | P1013 |
| Cellai, AP, | OR131, P2218 | Chandler, W, | OR303 |
| Carbone, AM, | P0412, P1428, P1440,
P1651, P2041, P2326 | Chandrakumaran, K, | P1315 |
| | | Chandy, M, | P0212, P0867, P2003 |
| Cercignani, M, | P1783 | Chang, C-S, | P2063 |
| Ceresa, E, | OR394, P0167,
P1364 | Chang, M, | P2141 |
| | | Chang, N, | OR377 |
| Ceresetto, JM, | OR087, P0198, P1964,
P2230 | Chang, Y, | OR165 |
| | | Changeux, JP, | P0342 |
| Cerini, C, | P0597 | Chantarangkul, V, | OR382, P0101, P0215,
P0701 |
| Cerletti, C, | P0148, P0569, P0572,
P1167, P1594 | | P2064 |
| erne, MC~, | P1113 | Chao, T-Y, | P0922 |
| Cerny, V, | P2032 | Chaouki, M, | P1006, P2198 |
| Cerqueira, M, | P2386 | Chapman, J, | P1441 |
| Cerrato, G, | P0090, P0614,
P1806 | Chapman, W, | P1659, P1660 |
| | | Charles, P, | P2175 |
| Cerutti, C, | P1719 | Charlesworth, MC, | P1939 |
| Cesari, F, | P0658, P1254, P1940 | Charoenkwan, P, | OR288 |
| Cetin, M, | P1002 | Charron, T, | P2196 |
| Cetkovsky, P, | P0255 | Chassany, O, | P0961, P2164 |
| Cha, SH, | P1081 | Chatamra, K, | P0465 |
| Chabielska, E, | P2180 | Chaudhary, AGA, | P0385, P2138 |
| Chabot, F, | P0075 | Chaudhary, RK, | OR030 |
| Chabrak, S, | P1097 | Chauhan, AK, | OR261 |
| Chackalamannil, S, | OR286 | Chavakis, T, | P0090 |
| Chacko, C, | H06 | Chavez, L, | P2292 |
| Chafa, O, | P1446 | Chawla, R, | P1523 |
| Chagraoui, A, | P2341 | Cheesman, EJ, | P1603 |
| Chahwala, S, | P0516, P1715, P2305,
P2306 | Chellini, B, | P1511 |
| | | Chen, BG, | P0940 |
| Chai, X-Y, | P1300 | Chen, C-F, | P1805 |
| Chait, P, | P2185 | Chen, CM, | OR260, OR389 |
| Chakravarthi, P, | P2108 | Chen, D, | P0152, P1963 |
| Chakroun, T, | P2295 | Chen, F, | OR181 |
| Chalabreysse, L, | P0572 | Chen, FP, | P1805 |
| Challier, C, | OR141 | Chen, HS, | P2130 |
| Chalmers, E, | P0239, P1245 | Chen, J, | OR117 |
| Chalmers, EA, | OR092, P2066 | Chen, L, | P1075, P1192 |
| Chalmers, J, | OR196 | Chen, M, | P0932, P1531 |
| Chambers, JP, | P2084 | Chen, P, | |
| Chamley, L, | P0621 | | |
| Chammas, M, | P0751 | | |
| Chamone, D, | P2195 | | |

Chen, PS, P0137, P1331
 Chen, Q, P0531
 Chen, VMY, OR075
 Chen, W, P1192
 Chen, Y, OR181, P0187
 Chen, Y-C, P2064
 Chen, Y-H, OR324
 Chenaud, C, P1303
 Cheng, JC, OR257
 Cheng, JT, P2356
 Cheng, S, OR156
 Cheng, S-N, P2064
 Chénier, M, P0441
 Chenou, C, P0695
 Cheong, PL, P1523
 Cherkasova, K, P0291, P1680, P2274, P2275
 Chesterman, CN, P1193, SYM86
 Chetty, N, P0589, P1724, P2225
 Cheung, P, OR052
 Cheveresan, L, P0665
 Cheveresan, M, P0665
 Chevreaud, C, P0180
 Chew, H, OR055, OR056
 Chhum, S, P1922
 Chi, HS, P1221
 Chi, L, P1213
 Chiang, J-L, P2064
 Chibedi, D, OR065
 Chiles, PG, P2266
 Chindemi, PA, P1201
 Chin-Dusting, J, P1166
 Chinni, E, OR094
 Chintala, M, OR286
 Chinthammitr, Y, P0672
 Chion, A, P1480A
 Chirinos, JA, P1763
 Chironi, G, P2350
 Chistolini, A, P2189
 Chitayat, D, OR164
 Cho, L, P2292
 Cho, MR, P0596, P1709
 Choi, DH, P1081
 Choi, G, P0728, P0753
 Choi, YK, P1081
 Chong, AY, P1175
 Chong, BH, P0307, P1099, P1538, P1590, P1673
 Choolani, M, P1314, P1597
 Choong, S, P1772
 Chopard, C, P0909
 Chorna, N, P1976
 Chotisupakarn, S, P2119
 Chou, E, P1847
 Choudhury, A, P1911
 Chow, T, P2361
 Chrenek, P, P1751
 Chrétien, MH, P0063, P0114, P0715

Christella, M, P0729
 Christensen, H, OR164, P1519
 Christensen, JH, P0749, P1187, P1782
 Christensen, K, P0620, P1640, P2022
 Christensen, SE, P1402
 Christersson, C, P0157, P2209
 Christian, FF, OR182
 Christian, WH, P1320
 Christiansen, K, OR132, P0095, P0840
 Christiansen, SC, P1814
 Christie, BA, P0823
 Christie, D, P2204
 Christine, H, OR185
 Christodoulos, K, P2192
 Christopherson, P, P2075
 Christopoulou-Cokkinou, V, P2286
 Chroust, K, P1998
 Chrzanowska-Wodnicka, M, P0599
 Chtourou, S, P0695, P0842, P1970, P2033
 Chu, H-L, P2011
 Chu, K, P1317
 Chuansumrit, A, P0732, P1939
 Chudasama, V, P0474, P1070
 Chudzinski-Tavassi, AM, P0713, P0714, P0883
 Chuen, CKY, P0340, P0939
 Chukhlovina, M, P1606
 Chun, F, P2383
 Chung, C-Y, P2063
 Chung, F, P0549, P0570
 Chung, HJ, P2290
 Chung, K, P1379
 Chunilal, S, P0247
 Chunilal, SD, P1659, P1660
 Church, FC, OR169, P0008, P0606, P1333
 Chyczewski, L, P1945
 Chyna, B, P0089, P1853, P1872
 Chytra, I, P0682
 Ciabattoni, G, P0958, P1557, P1558
 Ciampa, A, P0623
 Ciani, F, P1783
 Cichon, M, P0089
 Cid, A, P2073
 Cid, AR, P0282, P0871, P1864, P2013
 Cierniewski, CS, OR224, P0337, P0574, P0931, P1152, P1768, P2367
 Cieslikiewicz, D, P1610
 Cimino, E, P1428, P1440, P1651, P2041
 Cimmino, G, OR370
 Cimo, M, P1873
 Cini, M, OR177, OR179, OR180, P1644
 Cinotti, S, P1429

Cipolla, A, P0216, P0234, P0405, P0457, P1170, P1171
 Cipolla, N, P2034
 Cipriano, A, P1979
 Cirillo, F, P1651, P2041, P2326
 Cisowski, M, P0413
 Cisterna, B, OR220
 Citkowitz, A, OR077
 Ciuti, G, P1645, P1649
 Claeys, MJ, P2250
 Clark, D, P0229
 Clark, M, OR218
 Clark, P, OR239, P0493
 Clark, S, SYM76
 Clarke, MC, SYM13
 Clarke, S, OR178
 Class, D, P0261, P2370
 Clauser, S, P1875
 Clavreul, N, P1366
 Claxton, A, P0948
 Clemetson, JM, SOA02
 Clemetson, KJ, OR222, P2171, SOA02
 Clerici, C, P2219
 Clerici, M, OR382, P0215
 Clerin, V, OR390, P2355
 Clohisy, DR, P0823
 Clongstaff, C, P1910
 Cloutier, S, P0127, P1561
 Cludts, K, P0895
 Clutterbuck, R, OR319, P0977, P1159, P1536, P2354
 Cobain, TJ, P0957
 Cobb, R, P1359
 Cocci, F, P1608
 Cochery-Nouvellon, E, P0068, P0634
 Cochrane, A, P0859, P1343, P1517
 Cockerill, PN, OR203
 Coen Herak, D, P1291, P1292
 Coen, D, P1004
 Coen, M, P0639
 Coetzee, M, P0155
 Coghlan, D, P2234
 Cohen, A, OR275, P1627
 Cohen, AT, P0400, P1051, P1621, P1712, P2303
 Cohen, ER, OR133
 Cohen, H, OR090, P0543, P1141, P1741, P2307, P2331
 Cohen, M, P1611
 Colaizzo, D, OR094, OR197
 Colbatzky, F, P1702
 Cole, K, P2261
 Cole, SA, P1388
 Cole, V, P0123
 Cole, VJ, P1846
 Coleman, M, OR212
 Colle, S, P0875

Collen, D, OR233, P0032, P0551, P1164
 Collet, J-P, H17
 Collic-Jouault, S, OR143, P2296
 Collins, J, P1496
 Collins, P, OR036, P0327, P0874, P2332, P2333, P2354, P2383
 Collot, S, OR319, P1760
 Colman, RF, P1502
 Colman, RW, P0163, P0306, P1184, P1287, P1502, P2182
 Colman, S, OR196
 Colombo, M, P0218, P2191, P2330
 Colucci, G, OR291, P0377, P0915, P1574, P2171
 Colucci, M, OR293, P1108, P1573
 Colwell, CW, P1716
 Colwell, CW, Jr, OR061, P1667
 Combe, S, P0428, P1115, P1116, P1614, P1714
 Combot, C, P1296
 Combrié, R, P1520
 Comeglio, M, P1123
 Comfurius, P, OR345
 Comp, P, P1046
 Compernelle, G, OR394, P0168
 Compes, M, P0848
 Comuzzie, AG, P1388
 Conard, J, OR085, OR318, P0069, P0180, P0490, P0662, P0667, P1045, P1085, P1262, P1312, P2297
 Concolato, A, P1025, P2244
 Conkie, J, OR178
 Conlan, MG, OR268
 Conley, PB, P1496
 Connor, DE, P0122, P0945, P1194
 Connor, M, P0407
 Conradi, A, P1728
 Consiglio, P, P0726, P1236, P2280
 Consoli, A, P2352
 Constantopoulos, S, P0448
 Conti, AA, P1282
 Conti, S, P0257, P1027
 Conway, EM, P1164, SYM22
 Cook, PM, OR395
 Cook, SA, P1370
 Cooke, VG, OR295
 Cooley, SM, P1089
 Coombs, CJ, P1130
 Cooper, A, P0159, P0762, P0794, P1838
 Cooper, JA, OR153, P0401, P0475, P0997
 Cooper, PC, P0605, P0708, P1304
 Cooper, SM, P0605, P0708
 Cooper, ST, OR169

- | | | | |
|-----------------|--|-----------------|--|
| Coppens, M, | P0459, P0481, P1055,
P1056, P1856 | Cozzi, MG, | P1482 |
| Coppo, M, | P1123 | Cozzi, MR, | P0294, P1539 |
| Coppola, A, | OR207, P0218, P1428,
P1440, P1651, P2041,
P2326 | Craciun, A, | P0071 |
| Coppola, R, | P0766, P2324 | Craig, D, | P1820 |
| Corash, L, | OR268, P0108 | Crain, E, | P0162, P0197, P1852 |
| Corazzi, T, | OR370 | Cramer, E, | P0921 |
| Corbière, C, | P2341 | Cramer, EM, | OR163, OR165, P0322,
P1523 |
| Corbin, J, | SYM04 | Crandall, D, | P1981 |
| Cordova, R, | P1618 | Crandall, DL, | OR134, P0770 |
| Corinaldesi, C, | P0326, P0406, P1595,
P1692 | Cranford, LC, | P0606 |
| Corinaldesi, G, | P1595, P1692 | Craven, S, | P0504 |
| Cornelli, U, | P0519, P0991, P1118,
P2310 | Craven, SJ, | P0349, P0350, P1491 |
| Cornillon, J, | P0004 | Crawford, GL, | OR120 |
| Corno, AR, | P1477 | Crawley, JTB, | OR203, OR242, P0881,
P1480A |
| Cornuz, J, | OR305, P0470 | Crean, CS, | P2206 |
| Corral, J, | OR003, OR102, OR119,
OR234, P0529, P1127,
P1206, P1374, P1796,
P1797, P1968, P2048,
P2118, P2329 | Cree, A, | P2231 |
| Corrocher, R, | OR156, P0638 | Cremona, LS, | P2320 |
| Corsi, AM, | OR128, OR131, P0479,
P1282, P2271 | Cremonesi, M, | OR060 |
| Cortina, V, | P0282, P1864, P2013,
P2073 | Crescenzo, N, | P1593 |
| Corvazier, E, | P2133 | Crippa, L, | P0927 |
| Cosemans, JMEM, | OR368 | Crittenden, JR, | OR199 |
| Cosmi, B, | H02, OR177, OR179,
OR180, P1644 | Crookston, K, | OR268, P2268 |
| Costa, C, | P1403 | Crosby, D, | OR068 |
| Costa, F, | SYM71 | Crosdale, DJ, | P2172 |
| Costa, FF, | P1518 | Crotti, S, | P0451 |
| Costa, JM, | P0220 | Crowther, M, | OR311, P0443, P1898 |
| Costa, MC, | P0308 | Crowther, MA, | P2237 |
| Costanzo, M, | P0510, P1563, P1564 | Crudder, S, | OR383, P0258, P0973 |
| Cotrina, R, | P2328 | Cruz, MA, | P2130 |
| Cottini, S, | P1303 | Csóti, E, | P1093 |
| Coughlin, SR, | P0890 | Ctercteko, G, | P1039 |
| Cournoyer, D, | P2361 | Cubillo, P, | P1329 |
| Coutinho, A, | P1880 | Cueto, C, | P0626 |
| Couturad, F, | P0447 | Cugno, M, | P1180 |
| Couturaud, F, | P0446, P0668, P1048,
P1268, P1269, P1637 | Cui, J, | P0179, P0674 |
| Couvret, C, | P0309 | Cui, Y, | P0315 |
| Couzens, S, | P0327 | Cuisset, T, | P0411 |
| Covic, L, | OR015 | Culakova, E, | OR058, P1598 |
| Cox, D, | P0085, P1281, P1752,
P1726 | Cullen, J, | P0599 |
| Cox, DM, | P0328 | Culligan, K, | OR106, P0983 |
| Coyle, L, | P0616, P1321, P1376 | Culligan, KG, | P2359 |
| Coyne, E, | P2311 | Cultrera, D, | P0216, P0234, P0405,
P0457, P1170, P1171,
P2034 |
| Coyne, S, | P0892 | Cummine, J, | P2231 |
| Cozzi, E, | P0610 | Cumming, A, | P0874 |
| | | Cunanan, J, | P0520, P0521, P0710,
P0782, P1036, P1120,
P1717, P1288 |
| | | Cunningham, M, | OR303 |
| | | Cunto, E, | P0016 |
| | | Cupanik, V, | P0852 |
| | | Cupit, LD, | OR221 |
| | | Curmi, PMG, | P2137 |
| | | Curnow, J, | P1376 |
| | | Curtin, J, | P2155 |

Curtin, P, P0437
 Curtin, R, P2129
 Curtis, D, H15
 Curtiss, LK, OR323
 Curvers, J, P0386, P0987, P1584, P2179
 Cushman, M, P1669
 Cutini, C, P0297
 Cutter, J, SYM13
 Cuzon, G, P0083
 Cymbalista, CM, OR314
 Czerwicz, E, OR189
 Czestochowska, E, P0079, P0398, P0399, P0783, P0996, P1352
 Czlonkowska, A, P0078
 Czuriga, I, P0774
 Czwalinna, A, P1377
 D'Agnano, G, P1624
 D'Amico, E, P2195
 D'Amico, EA, P0199, P0673, P0994
 D'Angelo, A, P0927, P1914, P2199
 D'Arpa, S, P1171
 D'Atri, LP, P0941, P2132
 D'Atri, P, P1761
 D'Incà, M, P1449
 D'Rozario, J, P1990
 D'Souza, R, P1614
 D'Ursi, P, P0725
 D Borgel, D, P1875
 D. Belvini, D, P2006
 Dabbous, OH, P1017, P1693
 Dabrio, CG, P0435
 Daddi, G, P1646
 Daenens, K, OR029, P0744, P1701
 Dager, W, P0318, P1955
 Dahan, K, P0876
 Dahl, O, P1627
 Dahl, OE, OR062, P0472, P1667
 Dahlbäck, B, P0640, P1330, P1787, P1919
 Dahm, AEA, P1944
 Dai, J, P1378, P1420
 Dai, K, P2095
 Daikur, J, P0521
 Dailiana, Z, P1234
 Daleskog, M, P0355, P0436, P0604, P1527, P2147, P2226
 Dallongeville, J, OR155
 Daly, ME, P0378, P1478
 Dambrink, JHE, P0968
 Dämgen-von Brevern, G, P0098
 Damodar, P, P0394
 Damon, LI, OR268
 Danchev, D, P2294
 Dandie, G, P2231
 Danesh, J, SYM43
 Daniel, J-M, OR386, P1744

Danielsson, M, P0706, P0707
 Dannenberg, H, P2363
 Darby, A, OR390
 Dardik, R, OR297
 Dargaud, Y, P0174, P0866, P1409, P1984
 Darnell, K, P0049
 Darstein, C, P0956
 Dart, A, P1166
 Das, R, P1270
 Das, SS, P0385, P2138
 Dashevsky, O, OR296, P2363
 Dashkova, N, P0541
 Dashti, AA, P0077, P1237
 Date, H, P2287
 Dateki, T, OR209
 Daurès, JP, P0068
 Daures, JP, P1086
 Dautat, M, P0068
 Davagnanam, I, P2153
 Davalos, A, P1363
 Davey, M, P1659, P1660
 Davi, G, P0958, P0964, P1557, P1558, P2279
 David, D, P0223
 David, L, OR185
 David, T, OR255
 David-Dufilho, M, P1766
 Davido, A, P2260
 Davidson, AO, OR112
 Davidson, BL, OR061, OR366, P0424, P1712, P1716, P2303
 Davidson, SJ, P0161, P0224, P0972, P1306
 Davie, E, OR157, P1827
 Davie, EW, P1843
 Davies, DA, P0961, P1668, P2164
 Davies, J, P1315
 Davies, LW, OR070, P1524, P2176
 Davies, SW, P0972
 Davis, EF, P1621
 Davis, S, P1363, P1934
 Dawson, D, P0473, P2264
 Dawson, DJ, P0476, P1072, P1074
 Day, D, P0770
 de Boer, HC, P0329
 De Bonis, S, P0452
 de Bruijne, ELE, P1579
 De Bruin, K, P2102
 De Candia, E, OR376, P2115
 De Caterina, R, OR142, OR204, P0749, P1151, P1169, P1198, P1608, P2352
 De Cristofaro, R, OR376, P0279, P0879, P1931, P2115
 De Curtis, A, P0988, P1776
 De Filippi, F, P0218
 De Filippis, V, OR376

- | | | | |
|---------------------|--|--------------------|---|
| de Gaetano, G, | OR194, P0148, P0569,
P0670, P0988, P1167,
P1372, P1594, P1776,
P1861 | De Visser, MCH, | OR110, OR315, P0149,
P1283, P2071 |
| De Giorgi, A, | P1992 | De Vos, R, | H11, OR029 |
| de Graaf, J, | OR127 | de Vries, M, | P0818 |
| De Graeff, PA, | P0491, P2282 | De Vriese, AS, | OR129 |
| de Groot, PG, | OR028, OR256, OR294,
OR349, P0025, P0281,
P0336, P1812, P1813,
P2072 | de Winter, RJ, | P2150 |
| de Groot, PhG, | OR098, OR327, OR337,
P0563, P2008, P2253 | de Wolf, JThM, | P0102, P1892 |
| de Jonge, E, | OR031 | Deane, C, | P1069 |
| de Jonge, I, | OR318 | Dear, AE, | P1977 |
| de Kam, ML, | P1689, P2281 | DeBernardo, S, | OR390, P2355 |
| De Kort, W, | P2179 | Debili, N, | OR165, P2133 |
| De Kruif, MD, | P1878 | Debray, M, | P0277 |
| de Laat, HB, | P1813 | Decker, K, | P0228, P0807, P1997 |
| de Larrañaga, G, | P0016, P1031 | Deckers, NM, | P0557 |
| de Leeuw, PW, | P1384 | Decking, UM, | P1800 |
| de Lorgeril, M, | OR194, P0670 | Deckmyn, H, | OR244, OR331, OR363,
P0002, OR182, P0243,
P0310, P0331, P0334,
P0341, P0380, P0877,
P0935, P1972, P2069,
P2102, P2124, P2125,
P2157 |
| de Maat, MPM, | OR193, P0281, P1579 | Declerck, PJ, | OR394, P0167, P0168,
P1364, P1365 |
| De Maeyer, M, | P2069 | Decousus, H, | OR274, P0424, P1099,
P1590, P2313, SYM44 |
| de Maistre, E, | P0923 | Defoor, J, | H11 |
| De Marco, L, | P0294, P1539 | Degen, JL, | OR019 |
| De Mattei, M, | P0192 | Degerstrom, J, | P0583 |
| De Mattia, D, | P0396, P1454, P1593 | DeGrado, WF, | OR104 |
| De Mazancourt, P, | P0174 | Deguchi, H, | P0233, P0986, P1189
P2227, P2379 |
| de Mazancourt, P, | P0180, P0847 | Dejaeghere, D, | P0002 |
| de Mestre, A, | P1193, | del Aguila, MA, | P1865 |
| De Meyer, SF, | OR182 | Del Re, L, | P0958 |
| De Michele, R, | P1236 | del Rey, ML, | P0529, P2048, P2329 |
| De Michele, S, | P0726, P1236, P2280 | Del Rosso, M, | P1721 |
| De Moerloose, P, | OR111 | Del Turco, S, | P0749 |
| de Moerloose, P, | OR294, P0039, P0143,
P0184, P0252, P0356,
P0467, P1065, P1223,
P1303, P2161 | Del Vecchio, GC, | P1593 |
| de Moerloose, PH, | P0470, P1224 | DeLa Cadena, R, | P1750 |
| De Mol, B, | P0421 | DeLa Cadena, RA, | P1155 |
| de Nanteuil, G, | P1371 | Dela Cruz, RG, | P0607 |
| De Negri, F, | P1979, P1992 | Delahousse, B, | P0074 |
| De Palma, M, | P0192 | Delain, M, | P0004 |
| De Paula, EV, | OR300, P1518 | Delecroix, E, | P0696, P1984 |
| De Piano, LPA, | P0054, P0419, P1019,
P1041 | Delesque, N, | OR141, P1790 |
| de Pont, AC, | OR031 | Delgado, AV, | P0803, P1402, P2084 |
| De Prost, D, | OR317 | Delisle, L, | P1543 |
| De Punzio, I, | P1027 | Deliu, A, | OR394, P0168 |
| de Roos, A, | OR362 | Della Valle, P, | P0927 |
| De Rop, C, | P1506 | Dellgren, G, | P1748 |
| de Saint Martin, L, | P0432 | DeLucca, I, | P0162 |
| De Saint Martin, L, | P1789 | Delves, G, | P0794, P1838 |
| De Simone, C, | P0412, P1428,
P1440, P2041 | Dementieva, I, | P0081, P0541, P1380 |
| De Stefano, V, | P0064 | Demers, C, | P1561 |
| De Tinteniach, A, | OR272 | Demeulemeester, D, | P0551 |
| | | Demina, E, | P1126 |
| | | Demir, | P2193 |

- | | | | |
|--------------------|---|-------------------|---|
| Demir, M, | P0586, P0590, P1288,
P1467 | Di Maio, C, | P0726, P1236, P2280, |
| Demiral, HM, | P1397 | Di Minno, G, | OR094, P0412, P1428,
P1440, P2041, P2112,
P2326 |
| Demirel, GY, | P0899, P2152 | Di Minno, M, | P0412, P0452, P1440 |
| Dempfle, CE, | OR033, P0458, P0469,
P1991, P2158, P2240 | Di Nisio, M, | P0397, P1596, P2190,
P2203 |
| Demyanets, S, | P0561, P0580 | Di Paola, J, | OR346, P1472 |
| den Heijer, M, | H03, OR127, OR129,
P0480, P1079 | Di Renzo, L, | P0295, P1732 |
| Denecke, B, | OR330 | Di Santo, A, | OR040, P1592 |
| Deneys, V, | P0241, P1092 | Di Santo, S, | P0509 |
| Denis, CV, | P0134, P2072, SYM85 | Di Sanza, C, | OR111 |
| Denisa, W, | P0949 | Di Stasi, F, | P1413, P1414 |
| Denninger, H, | P1312 | Di Stasio, E, | P0879 |
| Dennison, D, | P2117 | Di, MG, | P1651 |
| DeNofrio, JC, | SYM76 | Diakur, J, | OR335 |
| Dent, J, | OR119 | Diakur, JM, | P2106 |
| Dentali, F, | P0451, P1052, P1630 | Diamandis, E, | P0773 |
| Depasse, F, | P1104, P2295 | Diamant, M, | OR054, P1319 |
| Depka, M von, | P1506 | Diaz, D, | OR082 |
| Depraetere, H, | P0331 | Diaz, M, | P1911 |
| Depunzio, I, | P0249 | DiBenedetto, D, | P1699 |
| Derenne, S, | P1215 | Diby, M, | P1303 |
| Derksen, RHWM, | OR256, OR349, P0025,
P1812, P1813 | Dickneite, G, | OR297, P0172 |
| DeRosa, L, | OR359, P0742 | Dickneitte, G, | OR023 |
| Desai, H, | OR372, P0982, P2105 | Didon, S, | P0651 |
| Desai, S, | P2263 | Diehl, JL, | P1875 |
| Descourt, R, | P0446, P1048, P1268,
P1269, P1637 | Diehl, S, | P1591 |
| Deshayes, A, | P0063 | Diehm, C, | P2208 |
| Desjardins, L, | P1634 | Dielis, AWJH, | P1384 |
| Desmarais, S, | P1634 | Diener, HC, | P1022 |
| Desmond, JF, | P1580 | Dietcher, S, | P1599, P2193 |
| Desmyter, L, | P1092 | Dietrich, CP, | P2311 |
| Desvard, F, | P1323 | Dietrich, U, | OR034, OR191, P1929 |
| Deutsch, V, | OR139, P1537, P1890 | Dietz, C, | P2342 |
| deVeber, GA, | OR355 | Diez, J, | P1023 |
| Develter, J, | P1365 | Differding, JA, | P0082 |
| Devenish, R, | P2047 | Diggs, B, | P0082 |
| Deville, A, | P0921 | diGirolamo, N, | P1150 |
| Dewaele, J, | P0032 | Dignat-George, F, | OR144, P0597, P1322,
P1770, P1789, P1807,
P2285 |
| Dewald, G, | P0669, P1794 | Dikkeschei, B, | P0884, P1299 |
| Dewar, L, | P0349, P0504, P1491 | Diks, S, | SYM03 |
| Dewerchin, M, | P0032 | Dillow, A, | P1362 |
| Dexter, M, | P2231 | Dimichele, DM, | P0056 |
| Dhanasekharan, D, | P0305 | DiMichele, DM, | P1924 |
| Dhanjal, TS, | P0934 | Din, C-P, | P0757 |
| Dhundale, A, | OR221 | Din, Y, | P0757 |
| Di Castelnuovo, A, | P0670, P1372, P1776,
P1861 | Ding, K, | P1153, P1946, P2344 |
| Di Cera, E, | OR048 | Ding, K-Y, | P0579 |
| Di Cuonzo, F, | P0396 | Ding, QL, | P1378, P1420 |
| Di Febbo, C, | P0958 | Ding, S, | P1508 |
| Di Giusto, DA, | OR333 | Dinisia, M, | P0009 |
| di lasio, mg, | P1914 | Diniz, EMA, | P0673 |
| Di Iorio, A, | P0958 | Diniz, MJ, | P0223 |
| | | Dinyari, M, | P1726 |
| | | Dippel, DWJ, | OR193 |

- | | | | |
|------------------|---|--------------------------|--------------------------------------|
| Diquattro, M, | P0726, P1236 | Douketis, J, | OR147, OR307, OR308,
P1020 |
| Dirckx, C, | OR194, P0670 | Douketis, JD, | OR364 |
| Diringer, MN, | P1363, P1934 | Dova, L, | P0448, P1234, P1642 |
| Dirks, C, | P1861 | Dovifat, C, | P2336 |
| Dissanayake, MP, | OR022 | Dowling, D, | P2228 |
| Distante, A, | OR204, P1151, P1169 | Dowling, GJ, | P1130 |
| Divoka, M, | P0809 | Dowling, N, | OR383, P0973, P1926 |
| Djemal, S, | P0427 | Dowling, NF, | P0258, P0649, P1280 |
| Djordjevic, V, | P1674, P2337 | Downey, C, | OR174, P0034, P1793 |
| Djuric, DM, | P0891 | Dr Torres, O, | P0226 |
| Dlott, J, | P1310 | Dr. Pollola, J, | P0226 |
| Dmitrieva, A, | P2272 | Dra Cermelj, M, | P0226, P1824 |
| Dobaczewski, G, | P1231 | Dra Ferro Ana, M-C, | P1824 |
| Dobrovolsky, A, | P1195, P1681, P2215 | Dra Ferro, AM, | P0226 |
| Dockal, M, | P0907 | Dra Ouviaña, S, | P0226, P1824 |
| Dodds, A, | P0617 | Dra Paz, A, | P0226, P1824 |
| Dodt, J, | P1246 | Dragani, A, | P0218 |
| Doering, CB, | OR184 | Dragomir, E, | P1147 |
| Doerr, H, | P0699 | Drechsler, WI, | P1434 |
| Doggen, CJM, | OR237, P0462, P0672,
P1061, P1063, P1238,
P1653, P1862 | Drenth, AF, | P0918, P2107 |
| | P1379 | Dressler, M, | P1202 |
| Doh, H, | P0176, P0418 | Dressman, H, | P1818 |
| Dohiguchi, Y, | P0086 | Dreven ek, Gš, | P1113 |
| Doi, P, | P0448, P1234 | Drexler, A, | P1016, P1514, P2109 |
| Dokou, E, | OR113, P0942 | Dreyfus, M, | P0921 |
| Dolan, G, | OR176 | Drigo, P, | P1604 |
| Dolce, A, | P1785 | Drijfhout, JW, | P1207 |
| Domagala, T, | P0482, P1080 | Drogemuller, CJ, | OR298 |
| Domagala, TB, | P0435, P1034 | Dropi ski, J nç, | P0482 |
| Domenech, P, | P1869 | Dropinski, J, | P0413, P1012, P1609,
P1785 |
| Donadini, M, | P1230 | | P0363, P0426, P1622,
P1623, P2196 |
| Donat, F ẽe, | P1180 | Drouet, L, | H06 |
| Donatelli, F, | P1592 | Drown, P, | P0071 |
| Donati, M, | OR040, OR194, P0148,
P0569, P0670, P0988,
P1167, P1372, P1594,
P1776, P1861, P1966,
P2084 | Drugan, C, | OR012 |
| Donati, MB, | P0273, P0315, P0855
P1507, P2136 | Drummond, GR, | P2365 |
| | P0917 | Drummond, O, | P0740, P1302, P1803 |
| Donna, E, | P1089 | Druzhinin, AV, | OR356, P1601 |
| Donnelly, JC, | P2129 | Düring, C, | P0152 |
| Dooley, M, | P0880 | Du, J, | OR181 |
| Doornebal, CW, | P1661 | Du, JW, | P2095, SYM78 |
| Dörje, C, | OR260, OR389 | Du, X, | OR253 |
| Dorling, A, | P0773 | Du, XY, | OR163 |
| Dorn, J, | P0848 | Dubart-Kupperschmitt, A, | OR367, P0347, P1500 |
| Dörner, T, | OR072, P1501, P2096 | Dubois, C, | OR087, P0198, P1964,
P2230 |
| Dorsam, RT, | OR087, P0198, P1964,
P2230 | Duboscq, C, | P1482 |
| Doti, C, | P0977, P1159, P1536 | | OR161, P0632 |
| Dotsenko, O, | P0597 | Duca, ML, | P0114, P0715 |
| Dou, L, | P1216 | Ducasse, C, | P1634 |
| Douard, P, | P0328 | Ducruet, T, | P2362 |
| Douglas, CWI, | P2242 | Dudani, A, | P0382 |
| Dougou-Tzaou, S, | | Dudbridge, F, | OR211 |
| | | Dudenhhausen, J, | P0917 |
| | | Dudkiewicz, P, | OR281 |
| | | Dudlova, J, | P0109, P1717 |
| | | Duff, R, | |

Dufresne, J, P1543
 Duga, S, OR126, P2039, P2040
 Duggan, C, P0427, P0429
 Dujols, C, P2296
 Dulí ek, P D ě ě, P1505
 Dulí ek, P ě ě, P1261
 Dulí ek, P ě ě, P2284
 Dulí cek, P, P1866
 Dumont, B, OR071
 Duncan, A, P0050, P0165, P1227
 Duncan, E, P1659, P1660
 Duncan, EM, P0247, P0804, P1825
 Duncan, MK, P1514
 Dundar, S, P1909
 Dunkley, S, P0320, P0631
 Dunn, A, P2029
 Dunn, AS, OR308
 Dunn, EJ, P1986
 Dunoyer-Geindre, S, OR140, P1223, P1224
 Dunsch, D, P1929
 Dunska, V, P1545
 Dupe, R, P0516, P2305
 Dupouy, D, OR010
 Durackova, Z, P2146
 Durand, E, P2296
 Duranovic, V, P1004
 Durante-Mangoni, E, P1881
 Dureau, G, P0572
 Dureuil, B, P0909
 Durham, T, P2206
 Durin-Assolant, A, P1426
 Du ek, L s ě, P1407
 Dusanter-Fourt, I, P0322
 Dusek, L, P1998, P2032
 Dusse, LMS, P0431, P0439, P0487,
 P0745, P1628, P2223
 Dusting, GJ, OR012, P0571
 Dutra, A, P2195
 Dutrillaux, F, OR317, P0923
 Dutt, T, P0873
 Dvorak, AM, SOA06
 Dvorak, HF, SOA06
 Dworakowska, D, P0079, P0398, P0399,
 P0783, P0996, P1352
 Dworakowski, R, P0079, P0398, P0399,
 P0783, P0996, P1352
 Dwyre, L, P0200
 Dyck, J, P1728
 Dyr, JE, P1505
 Dyson, JM, OR116
 Dyszkiewicz-Korpanty, A, P1012
 Dytrychová, V, P1293
 Dziedzina, S, P0413
 Dzoljic, M, P0753
 É, P2150
 Eagle, KA, P1699
 Eastham, D, P0786

Easton, L, P0864
 Ebbesen, LS, OR132
 Eberl, W, P1860
 Eble, JA, OR018, P1494
 Eby, C, OR303
 Echelard, Y, P0610
 Eckes, B, OR254
 Eckly, A, OR255
 Eckman, JR, P0073
 Eckstein, R, P0175
 Edelmiro, M, P1580
 Edenberg, H, P1360
 Édes, I, P0774
 Edlund, M, P1769
 Edmunds, LH, Jr., P0163
 Edouard, A, P1297
 Edson, B, P0709, P1309
 Edwards, S, P1141
 Edwards, VD, P1519
 Efron, E, P0198
 Egan, B, P0805
 Egberts, JFM, P1712, P2303
 Egorina, E, P0151
 Eguchi, Y, P2346, P2348
 Ehrlich, HJ, P0138
 Ehrlich, YH, P0339, P0409, P2308,
 H03, OR270, OR361,
 P0438, P0743, P0992,
 P1040, P1935, P2184
 Eid, S, P2042
 Eidam, A, P0099
 Eidingen, GW, P1021
 Eifrig, B, P0256, P2015, P2154
 Eigenthaler, M, P0289, P1487, P2083
 Eikelboom, JW, P0948, P1846, P2202
 Eikenboom, J, H09, H08, P0270, P0271,
 P0265
 Eikenboom, HCJ, OR315, P1125, P1238,
 P2071
 Eikenboom, J, OR245, OR278, OR279,
 OR281, P0865, P1461,
 P1469, P2057
 Eikenboom, JCJ, P0269, P2059
 Eikvar, L, P1029
 Eilertsen, AL, P0492
 Eilertsen, KE, P1758, P1759
 Einarson, T, P0227
 Eisert, R, P0210, P0986, P1377
 Eisert, RM, P0233
 Eissner, G, P0593
 Ejiri, S, P2348
 Ekert, H, P0805
 Ekholm, C, OR277
 Ekström, B, P1375
 El Amara, W, P0364
 El Ayachi, S, P2369
 El Kheir, D, P1654, P2248, P2254
 El Nemer, W, P2338

- | | | | |
|--------------------|---|----------------------------|---|
| El Rouby, S, | P1023, P1145, P1611,
P1743, P2336 | Eriksen, K, | P0683 |
| El Sherbiny, M, | P0675 | Eriksson, B, | P1627 |
| Elalamy, I, | P0490, P0662, P1045,
P1085, P1262, P2295,
P2297 | Eriksson, BI, | OR062, P2216 |
| El-Ali, M, | P2286 | Eriksson, H, | OR057, P0416, P1738 |
| Elawady, M, | P2368 | Eriksson-Berg, M, | P0604 |
| Elbaz, O, | P0582, P2368 | Ernest, CS, | P0952, P0956 |
| Elchaninov, P, | P1606 | Ertl, G, | P0289, P1487, P2083 |
| Elford, H, | P1772 | Erwich, JJHM, | OR213, P1090 |
| Elfvinge, P, | P2027 | Erwig, LP, | P0026 |
| Elgue, G, | P0158 | Esbensen, K, | P1965 |
| Elias, DJ, | P1189, P2227, P2379 | Eschrich, R, | P1885 |
| Eliasson, Å, | P1066 | Escuriola, C, | OR191, P1929 |
| Elice, F, | P1334 | Escuriola-Ettinghausen, C, | P0812 |
| El-Idrissi, A, | P0409 | Eser, B, | P1002 |
| Elleman-Olesen, M, | P1619 | Eshghi, P, | P1389 |
| Eller, Th, | P1675 | Eskiocak, M, | P1467 |
| Ellery, P, | P1341 | Esmon, C, | P0125 |
| Ellingsen, C, | P2228 | Esmon, CT, | P0105, P0734 |
| Elliott, JL, | OR121 | España, F, | P0730, P0731, P1328,
P1329 |
| Elliott, MA, | P0111, P1308, P2173 | Espinola, R, | P1184 |
| Ellis, A, | P0858, P1342 | Esposito, BA, | OR133 |
| Ellis, SA, | OR112 | Esposito, E, | P0726 |
| Ellwanger, C, | P1745 | Essayagh, S, | OR010 |
| Elnoor, IB, | P1134 | Estellés, A, | P0730, P0731, P1328,
P1329 |
| Elokdah, H, | P1981 | Etienne, E, | OR272 |
| Elvevoll, EO, | P1758, P1759 | Eto, T, | P1586 |
| Emanuel, V, | P0528 | Ettema, HB, | OR066, P1057 |
| Emmerich, J, | OR065, P0667, P1276 | Eugène, S, | P0552 |
| Emsley, J, | H04, OR097 | Evangelista, M, | P0749 |
| Enayat, MS, | P0821, P1412, P1470,
P2066 | Evangelisti, L, | P1649, P1721, P1940 |
| Enayat, S, | P0874 | Evans, G, | P0720, P1339, P1433,
P1434, P1897 |
| Enders, G, | OR042 | Evans, S, | P0320, P2100 |
| Endoh, I, | OR231 | Evers, EJ, | P2241 |
| Engelhardt, W, | P0097, P0099, P0698,
P0699, P0885 | Evers, LM, | P0573, P1753 |
| Engelmann, B, | OR100 | Everse, SJ, | OR249 |
| Engelmann, L, | OR079, P0717, P0812,
P2177 | Evgen, M, | P1394 |
| Engelsen, J, | P1294 | Evgueni, S, | P1394 |
| Engl, W, | P1231 | Ewenstein, B, | P0829, P1837 |
| Engová, D, | P0429 | Ewing, N, | P2036 |
| Enjolras, N, | OR161, P0632 | Ewing, NP, | P1436 |
| Ennamany, R, | P0520 | Exner, M, | OR264 |
| Enomoto, TM, | P0082 | Exner, T, | OR084, P0113, P0122,
P0617, P0945, P1297 |
| Enouf, J, | P2133 | Externest, D, | P2020 |
| Ensslen, S, | OR330 | Eyries, M, | P0595, P1362 |
| Eppihimer, MJ, | OR390 | Ezekowitz, M, | OR271 |
| Eppe, G, | P0186 | Fabbrini, N, | P1911 |
| Epstein, SE, | P2386 | Fabre, A, | P2268 |
| Er, O, | P1002 | Fabris, F, | P0030, P1522 |
| Erahbor, T, | P0635 | Faden, D2, | P1215 |
| Eram, SM, | P1411 | Fadin, M, | P0588, P1264, P1810 |
| Erber, WN, | P1336 | Fagnani, F, | OR065 |
| Erger, RA, | P1332 | Fagon, JY, | P1875 |
| | | Fahmy, RG, | OR388, P1150 |

Fahs, S, P2075
 Fahs, SA, H05, OR186
 Fain, O, P1215, P1216
 Fainboim, H, P1031
 Faioni, EM, P0086, P0725, P2376
 Falanga, A, OR039, OR060, P2116
 Falaschi, F, P1706
 Falati, S, P1541
 Falciani, M, OR131, P0479, P1026,
 P1124, P2325
 Falcinelli, E, OR370, P154
 Falco, A, P0958, P1557
 Falet, H, OR115, OR116
 Fan, Q-S, P2011
 Fang, J, OR108, OR186
 Fang, Y, P1378, P2011
 Faradji, A, P1426, P2021
 Faradz, S, P0203
 Farah, RA, P0039, P0751
 Fard, P, P1412
 Fareed, D, P0110, P0520, P0521,
 P1177, P1186, P1288,
 P1337, P2194
 Fareed, J, OR335, P0014, P0089,
 P0109, P0110, P0518,
 P0519, P0520, P0521,
 P0547, P0609, P0710,
 P0782, P0913, P0991,
 P1023, P1100, P1118,
 P1120, P1177, P1185,
 P1186, P1288, P1337,
 P1599, P1611, P1717,
 P1781, P1853, P1872,
 P1899, P1902, P1988,
 P2193, P2194, P2310,
 P2312, P2334
 Farge, D, P1216
 Faria, F, P0883
 Fariás, C, P0276, P0863, P1003,
 P1473
 Fariás, CE, P0388
 Farid, NA, P0952, P0956
 Faridi, A, OR211
 Farina, A, P1648
 Farinaro, E, P1966
 Farndale, R, P0299
 Farndale, RW, OR070, OR347, P0331,
 P0935, P0981, P1524,
 P1528, P2090, P2094,
 P2176, SYM59
 Farndale, W, P2093
 Farouque, HMO, P1560
 Farrell, DH, OR109
 Farren, T, OR212, P0346
 Farris, D, OR135, P1981
 Farrugia, C, P0083
 Fass, DN, P0650
 Fath, J, P2037

Fatini, C, P0479, P0658, P1254,
 P1684A, P1720, P1773
 Fattorini, A, P0927
 Fattorini, L, P0249
 Faulkner, LJ, P0047
 Faure, V, P0597
 Fauvel-Lafeve, F, OR037, P0506, P2126
 Favaloro, EJ, P0864, P1129, P1983
 Favaloro, JWC, P0864
 Favier, R, P0322, P0724
 Faxälv, L, P0886
 Fay, PJ, P1247, SYM34
 Fearnside, M, P2231
 Fearon, WF, P1560
 Featherstone, J, P0762
 Federici, A, H08, H09, OR245,
 OR278, OR279, OR281,
 P0265, P0270, P0271,
 P1461, P1469, P2057
 Federici, AB, P0269, P0869, P0870,
 P2059
 Federici, F, P0865
 Federico, H, H13
 Fedi, S, OR131, P0479, P0658,
 P1254, P1603, P1678,
 P2325
 Fedotkina, Yu, P2215
 Fedotova, I, P0777, P2291
 Fefer, P, P0950
 Feijge, MAH, OR051, OR344, P0987,
 P1540, P1584, P1701
 Feilotter, H, P0690
 Feindt, P, P0797
 Feingold, J, P1241
 Feistritz, C, OR230, OR375
 Feldman, B, P0227
 Feldman, LJ, P0364
 Feldo, M, P1959
 Felici, A, P0395
 Felicissimo, P, P1102, P1103
 Feliú, A, P1018, P2050
 Fellows, GA, P0400
 Fenech, M, P0247
 Feng, R, P0359
 Feng, Y, P1662
 Fenneteau, O, P0921
 Fens, D, OR153
 Fenton II, J, P1491
 Fenyvesi, T, P1015, P1106, P1705,
 P1801
 Feraud, O, OR165
 Ferec, C, P1268, P1269, P1637
 Feren ak, G c̃, P2314
 Fernández, G, P1761
 Fernández, JA, OR048, P1925, P2227
 Fernandes, AP, P0431, P0439, P0487,
 P0745, P1628, P1690,
 P2223

- | | | | |
|---------------------|----------------------|----------------------|----------------------|
| Fernandes, JO, | P1865 | Fischer, K, | OR205, P2007, P2008, |
| Fernandez, JA, | P1332 | Fischer, R, | P0232, P1231, P2021 |
| Fernandez-Borja, M, | OR008 | Fischer, T, | OR211 |
| Ferrando, F, | P0730, P0731 | Fish, R, | OR140 |
| Ferrante, F, | P0249, P1027, P1613, | Fish, RJ, | P1223 |
| | P1733 | Fishbein, MC, | P0566 |
| Ferrara, JD, | OR103 | Fishel, S, | P2279 |
| Ferraresi, P, | P1939 | Fisher, R, | P1525 |
| Ferrari, E, | P1869 | Fisher, WD, | OR063 |
| Ferrari, S, | P0907, P1563 | Fissha, M, | P0517, P2205 |
| Ferraris, SP, | P1525 | Fitzgerald, A, | P2129 |
| Ferraris, VA, | P1525 | Fitzgerald, DJ, | P2129 |
| Ferraro, L, | P2280 | FitzGerald, G, | OR274, P1078, P1590, |
| Ferrazzi, P, | P1677 | | P1643, P1671, P1673 |
| Ferreira, MFR, | P0745 | Fitzgerald, JR, | P0085 |
| Ferreira, MS, | P0140 | Fitzmaurice, DA, | P0107 |
| Ferrero, S, | P0086 | Fkih, M, | P1097 |
| Ferrini, S, | P1349, P2271 | Fläring, A, | OR154 |
| Ferris, NML, | P0427 | Flan, B, | P0842, P2033 |
| Ferro, D, | P0509 | Flaujac, C, | OR085, P0662, P1045 |
| Ferroni, P, | P1557, P2279 | | P2297 |
| Ferrucci, L, | OR128, OR131, P0479, | Flaumenhaft, R, | OR253 |
| | P1282, P2271 | Fleseriu, I, | P1196 |
| Fex, B, | OR135 | Fletcher, J, | OR276, P0437, P0560, |
| Feyen, JHM, | P0197, P2183 | | P0594, P1039, P2231 |
| Feys, HB, | P2102 | Fletcher, JP, | P2233 |
| Ffrench, PO, | P0174 | Flierl, U, | P0289 |
| Fibbi, G, | P1721 | Flight, S, | P1975 |
| Fichelson, S, | P1523 | Flipiak, KJ, | P0177, P1610 |
| Fiedler, F, | OR033, P2158 | Floccard, B, | P0696 |
| Figg, N, | SYM13 | Flodgaard, H, | OR262 |
| Figueiredo, MS, | P0093, P0618 | Flood, VH, | OR109 |
| Figura, S, | P0021, P1929 | Flores, MPA, | P0713 |
| Fijnheer, R, | OR028 | Florian, M, | P0089, P0520 |
| Filatov, V, | P0196, P0792 | Floriani, I, | OR060 |
| Filer, K, | P0795 | Florian-Kujawski, M, | OR335, P0782, P1118, |
| Filho, CC, | P2236 | | P1781, P1853, P1872, |
| Fillion, ML, | P1200 | Floros, G, | P1988, P2310, |
| Filippucci, E, | P1643, P2219 | Florquin, S, | P1132, P1425 |
| Filipuzzi, L, | P0923 | Flounders, K, | OR391 |
| Filoni, C, | P1684A, P2325 | Flynn, M, | P0092 |
| Finazzi, G, | P1220, P1906, P2116 | Fogato, L, | P0543 |
| Finazzi, S, | P1007 | Fok, TF, | P0192 |
| Finch, S, | P0957 | Foka, ZI, | P0340, P1538 |
| Findlay, JBC, | P0286, P2079 | Foley, JH, | P2002 |
| Fink, L, | OR041, P1334 | Folkeringa, N, | OR096 |
| Finn, B, | P2230 | Follows, GA, | OR213, P1090 |
| Finocchi, A, | P0509 | Folsom, AR, | OR203 |
| Fiore, | P0688 | Fondu, P, | P1669 |
| Fiore, LD, | OR089 | Fons, P, | P2125 |
| Firbas, C, | P0001, P0722, P1874 | Fonseca Neto, CP, | OR141, P1790 |
| Fisac Herrero, R, | P0485 | Fontaine, A, | P1690 |
| Fisac Martin, M, | P0485 | Fontaine, S, | P1543 |
| Fischer, AM, | OR143, P2296 | Fontaine, S, | P0724, P1297 |
| Fischer, A-M, | P1446 | Fontana, G, | P0086, P2376 |
| Fischer, D, | P0131 | Fontana, P, | P0356, P1323, P2161 |
| Fischer, E, | P1093 | Fontana, V, | P0917, P0919, P0999 |
| Fischer, H, | P1021, P1748 | Fontayne, A, | P0002, P2157 |

Fontcuberta, J, OR318, P0466, P1974
 Forastiero, R, P0015, P0016, P0090, P0614, P1806
 Ford, E, OR266
 Ford, I, P0026, P0474, P1140, P2261, P2332, P2333
 Forgan, M, P0493
 Foroud, T, P1360
 Forslund, M, P0930
 Fortenberry, YM, OR169, P0606
 Fortini, P, P0192
 Fortuna, S, P0132
 Fossum, T, P2305
 Foster, C, OR286
 Foster, P, P0060
 Foster, TJ, P0085, P0328, P1281
 Foti, L, OR094, OR197
 Fouda, M, P0582, P0675
 Fournier, M, P0425
 Fox, E, P0440
 Fox, KAA, P1017, P1693, P1699
 Fox, ML, P0954, P0955
 Fox, SC, OR334, P0942, P1498, P2149
 Fraccarollo, D, P1487, P2083
 Frampton, J, OR042, OR322
 Francis, CW, OR058, P1598, P1747
 Francis, JL, OR287, OR372, P0982, P1016, P1514, P2015, P2105, P2109, P2110, P2154
 Francis, SE, P2172
 Franco, HA, P0487
 Franco, RF, P0093, P0618, P2054, P2055, P2235
 Franco, RM, P0487
 Franken, T, P0655
 Franks, P, P0478
 Franow, H, P1729
 Frantz, S, P2083
 Franz, F, P1618, P1722
 Franzoi, M, P1604
 Frattani, FS, P0140
 Fredenburgh, JC, P1957
 Frederix, K, OR043, P0752, P1208, P1209
 Freedman, J, OR288, P0985, P1491, P1531, P1542, P2143
 Freedman, JE, P1504
 Freeley, M, P2359
 Freeman, MS, OR049, OR314
 Freeman, R, P0091
 Freestone, B, P1175
 Fregin, A, OR187
 Frelinger, AL, OR138, P0357, P0954, P0955
 Frelinger, III AL, P0361
 French, JK, P1694

Frere, C, P0169, P0411, P1382
 Freson, K, H11, P0321, P0920
 Fressinaud, E, OR278, P0265, P0267, P0270, P0271, P0277, P1426
 Freund, M, OR059, OR255, P1427
 Frey, M, P0561
 Freyburger, G, OR317
 Friberg, B, OR277
 Fribourg, C, OR188, P0659
 Friedl, P, SYM42
 Friedrich, W, P0925
 Friedrichs, B, P1162
 Frigerio, M, P0352, P0358
 Friso, S, OR156, P0638
 Fritsch, S, P0138, P0829
 Fritsche-Danielson, R, H14
 Fritzen, M, P0713
 Fritzsche, J, P0548
 Froehlich, JB, OR274, P1078, P1671
 Froesner, G, P0832
 Frölke, JPM, P1397
 Frontoni, R, P0701
 Fronroth, JP, P1018, P2050
 Frossard, M, P2142
 Frost, C, P1318
 Frost, FJ, OR308
 Frostick, SP, P1667
 Frulla, M, P1025, P2244
 Fu, B, P0152
 Fu, J, P1946
 Fu, QH, OR124, P0007, P0387, P1378
 Fuchs, I, P2142
 Fuchs, S, P2386
 Fuertbauer, E, P1751
 Fuhrmann, J, OR243
 Fujii, G, SYM71
 Fujii, K, P0133
 Fujii, T, P0553
 Fujimoto, M, P0791, P2103
 Fujimura, Y, P1178, P2103
 Fukuda, A, P0799
 Fukuda, K, P1652
 Fukuda, T, P0511, P0512, P0612
 Fukui, H, P2103
 Fukushima, N, P0784
 Fukutake, K, P0800, P1256, P1832
 Fuller, G, P1494
 Fuller, GLJ, OR018
 Fuller, N, P1826
 Fulton, J, P2265
 Fumeaux, T, OR233
 Funahashi, T, OR285
 Fung, S, OR211
 Funiaková, S, P0360
 Funk, M, P0581
 Furber, A, OR304, P1049, P2260

- | | | | |
|------------------|--|------------------------|---|
| Furie, B, | OR189, OR253, OR367,
OR374, OR377, P0347,
P0890, P1500, P1541 | Gan, A, | OR359, P0858, P0742,
P1342 |
| Furie, Ba, | P1822 | Gan, TE, | P1700, P1884, P2101,
P2211 |
| Furie, BC, | OR189, OR253, OR367
OR374, OR377, P0347,
P0890, P1500, P1541,
SYM60 | Ganapathy, P, | P2316 |
| Furie, Br, | P1822 | Ganderton, T, | SYM86 |
| Furman, MI, | OR138, P0357, P0954,
P0955 | Gandini, G, | P2006 |
| Furmedge, J, | OR359, P0742, P0858,
P0859, P1342, P1343,
P1517 | Gando, S, | P1994 |
| Furtado, C, | P0971 | Gandrille, S, | P1914 |
| Furugohri, T, | P0511, P1110 | Gane, P, | P2338 |
| Furukoji, E, | P1178 | Ganesan, V, | OR355 |
| Fuster, JL, | P1797 | Gangadharan, B, | OR184 |
| Futers, TS, | OR314 | Ganjanakarn, E, | P2119 |
| Gähler, A, | P0776 | Ganser, A, | P0210, P0233, P0986,
P1377 |
| Gabazza, EC, | P0133 | GAO, W, | P1509 |
| Gabi, K, | P0776 | Garancini, P, | P2199 |
| Gabriel, DA, | P1260 | García-Barberá, N, | OR102, P1374 |
| Gachet, C, | OR255, OR369, P0739,
P1490, P2174, SYM17 | García De La Torre, I, | P0565 |
| Gadisseur, A, | P0272 | Garcia de Lorenzo, A, | P1585 |
| Gaffney, PJ, | P1975 | Garcia, A, | P1494, P2096 |
| Gafou, A, | P1889 | Garcia, AA, | P2054, P2055 |
| Gagnon, D, | OR089 | Garcia, D, | OR311, P2268 |
| Gaidarov, B, | P0827 | Garcia, JCF, | P1690 |
| Gaikovaya, L, | P1183 | Garcia, JF, | OR272 |
| Gailani, D, | H12, OR097, OR341,
OR342, P1822 | Garcia, ML, | P0431, P1628, P2223 |
| Galanakis, DK, | OR114 | Garcia, N, | P1031 |
| Galar, M, | P0170, P1368, P2385 | Garcia, RA, | OR300 |
| Gale, AJ, | P1423 | Garcia-Talavera, J, | P2045 |
| Galea, B, | P2286 | Gardiner, C, | OR090, P2307 |
| Gallagher, J, | OR004 | Gardiner, E, | OR067, P0894, SYM18 |
| Galli, M, | OR311, P1052, P1220,
P1869, P1906, SYM12 | Gardner, A, | OR298 |
| Gallimore, MJ, | P0720, P0768, P0769,
P1339, P1897 | Garewal, G, | P1270 |
| Gallinaro, L, | OR282, P0862 | Garghetti, E, | P1482 |
| Gallo, L, | P1891 | Gari, MA, | P0465 |
| Gallo, N, | P0690 | Garin, G, | OR385, P2383 |
| Gallotta, G, | P0452 | Garner, SF, | P1524 |
| Gallus, A, | OR064, OR265, P0424,
P2202, P2234, P2246,
SYM11, SYM37 | Garo, F, | P2141 |
| Gallus, AG, | P1712 | Garrard, CS, | P0728 |
| Gallus, AS, | P1051, P1643, P2303 | Garth, S, | OR276 |
| Galton, DJ, | P0475 | Gartner, TK, | P1494 |
| Galy-Fauroux, I, | OR143 | Gary, EG, | P1320 |
| Gamberalle, R, | P2085 | Gasem, MH, | P1995 |
| Gamberini, M, | P2360 | Gaspar, L, | P1632 |
| Gamble, JR, | OR298 | Gast, MCW, | OR240 |
| Gamzatov, T, | P1183 | Gattens, M, | P0544 |
| | | Gatti, S, | P0639 |
| | | Gatzemeier, U, | OR059 |
| | | Gaudin, P, | P0266 |
| | | Gaus, K, | P0893 |
| | | Gaussem, P, | P0378, P1323 |
| | | Gavalaki, M, | P1050 |
| | | Gavasso, G, | P1722 |
| | | Gavasso, S, | P0132, P0250, P0610,
P1604, P1810, P1915 |
| | | Gavazova, S, | P2075 |
| | | Gavornik, P, | P1632 |
| | | Gawaz, M, | OR259, OR320, OR322 |

Home Author Index Keyword Search

Gay, V, P0847
 Gayen-Betal, S, P0550, P1149
 Gazzini, A, P1940
 Gérald, B, P1521
 Géronimi, F, OR141
 Gézsi, A, P1476
 Geary, MP, P1089
 Gebbink, MFBG, OR098, OR327, OR337,
 P0329, P0563
 Geczy, C, P1146
 Geczy, CL, OR231
 Geeraedts, LMG, P1397
 Geerts, W, P2237
 Geha, RS, OR115
 Gehrisch, G, P1233
 Geil, J, P0138
 Geisel, J, P2351
 Geisen, C, OR101, OR187, P0056,
 P0525, P0848
 Geisen, P, P0599
 Geisen, U, OR211
 Geiter, S, OR270, P0875
 Gelmont, D, P0138
 Gemmati, D, P0192, P1831
 Gemmell, R, P2100
 Geneux, MG, P0317
 Geng, J, P1772
 Geng, J-G, P1075
 Geng, Y-J, OR142, P1198
 Gennari, LC, P0615
 Gensini, F, P1124, P1723
 Gensini, GF, OR128, OR131, P0368,
 P0456, P0479, P0496,
 P0510, P0658, P1026,
 P1095, P1123, P1124,
 P1254, P1282, P1349,
 P1563, P1564, P1603,
 P1624, P1645, P1649,
 P1678, P1684A, P1720,
 P1723, P1773, P1940,
 P2218, P2271
 Gent, M, OR063, OR147, P0424,
 P1020, P1046, P1051
 Gent, YJJ, OR327
 Gentilomo, C, P1604
 Gentry, HR, OR103
 George, B, P0212, P0867, P2003
 George, J, OR139, OR268, P0940
 George, MR, P1272
 George, MW, P1326
 Gera, R, P0052
 Gerard, B, P0921
 Gerard, RD, OR011
 Gerasimidis, T, P1617, P2197
 Gercovich, D, P0226, P1824
 Gerdes, V, P1654, P2237, P2248
 Gerdes, VEA, P0150, P0685, P0687
 P1060, P2251

Gerenli, M, P0586
 Gerhard, D, P0978
 Gerhardt, A, OR215, OR316, P0535,
 P1685, P2127, P2277
 Gerland, V, P1409
 Germer, M, OR045, P0035
 Gerotziafas, GT, P1104, P2295
 Gerrits, WBJ, H03
 Gershkovich, KB, P1385
 Gerstle, T, P0393
 Gessner, R, P0186
 Gestel, D, P1416
 Gevaert, K, P2141
 Gevenois, PA, OR306
 Gevers, T, P1834
 Ghaly, M, P1527
 Ghamari, A, P0067, P1730
 Ghanima, W, P1661
 Gharavi, AE, P0028
 Ghaznawi, F, P0339
 Ghent, T, P0689
 Gheorghiu, E, P0665
 Ghevaert, C, OR218, OR358
 Ghezzi, A, P1783
 Ghezzi, F, P0451
 Ghiotto, R, OR126, P2039, P2040
 Ghirarduzzi, A, H02, OR366, P0023,
 P0257, P0537, P1630,
 P1647, P1811
 Ghosh, K, P0268, P0676, P0677,
 P1464
 Giacomelli, S, P0070
 Gianelli, U, P0086
 Giangrande, P, P0209, P1241
 Giannarelli, C, P1979, P1992
 Gianniello, F, P0869
 Giansily-Blaizot, M, P0146, P0751
 Giarratano, A, P1706
 Gibb, CB, P1924
 Gibb, S, P1037
 Gibbins, JM, OR017, P0900, P1541
 Gibbs, H, P1037, P1038
 Gibson, C, P0916
 Gibson, NS, P1060, P2251
 Giddings, J, P0327
 Giddings, JC, P0961, P1668, P216
 Giese, CH, P1015, P1106, P1705,
 P1801
 Giesler, U, P1122
 Giglioli, C, P0510, P1095, P1254,
 P1563
 Gijzen, KA, OR008
 Gilbert, GE, P0647
 Gill, J, P2075
 Gille, J, P1591
 Gillen, J, P1089
 Gilles, JG, OR159
 Gilmore, G, P0948, P0957

- | | | | |
|-----------------|--|-----------------------|---|
| Gils, A, | OR394, P1364, P1365,
P0167, P0168 | Golovina, OG, | P0679 |
| Ginsberg, JS, | OR147, P0127, P1020,
P2237 | Goltsova, K, | P1681, P2215 |
| Ginsburg, D, | OR025, OR027, OR030,
P0578, PLEN5 | Golubeva, M, | P0819 |
| Giordano, G, | P1594 | Gomaa, M, | P1764 |
| Giordano, KF, | P2267 | Gomes, L, | P0134 |
| Giordano, L, | P0988 | Gomes, N, | OR037 |
| Giordano, P, | P0396, P1454, P1593 | Gomez, GEB, | OR193 |
| Girard, P, | P0445, P0447, P1044,
P1047 | Gomez, JG, | P0666 |
| Girardi, G, | OR352 | Gomez, JM, | P0666, P1034 |
| Girelli, D, | OR156, P0638 | Gomez, K, | P0761 |
| Girisch, M, | P0861 | Gomez, N, | P0016 |
| Girndt, M, | P2351 | Gomez, RM, | P0941, P2132 |
| Girolami, A, | OR366, P0030, P0217,
P0251, P0814, P1450,
P2244, P2245 | Gomperts, ED, | P2028 |
| Girolami, B, | P0814, P2245 | Goncalves, I, | P2081 |
| Girtovitis, F, | P0116, P0312, P0384,
P1641, P2002, P2121,
P2242 | Gonchar, V, | P2026 |
| Gissel, MT, | P2028 | Goncharov, I, | P1890 |
| Gissen, P, | OR164 | Gong, X, | P2131 |
| Gitel, S, | P2169 | González, B, | P1484 |
| Giuliano, V, | P2219 | González-Conejero, R, | OR102, P0529, P1796,
P1968, P2048, P2329 |
| Giuseppe, T, | P1415 | González-Enríquez, E, | P1626 |
| Giusti, B, | OR128, P1095, P1124,
P1721, P2271 | Gonzales, A, | P1611 |
| Giusti, C, | P0958 | Gonzalez Vucovik, M, | OR087 |
| Giustolisi, R, | P0216, P0234, P0405,
P0457, P1170, P1171,
P2034 | Gonzalez-Conejero, R, | P1127, P1374, P2118 |
| Gizzi, S, | P2219 | Goodall, AH, | OR290, OR347, P0105,
P0474, P0721, P0734,
P1070, P1528, P1553 |
| Gladysheva, I, | P1962 | Goodall, GJ, | OR298 |
| Glenn, JR, | OR334, P0942,
P2134 | Goodeve, A, | H08, H09, OR245,
OR278, OR279, OR281,
P0265, P0269, P0270,
P0271, P0865, P1461,
P1469, P2057, P2059 |
| Glenn, P, | OR182 | Goodfellow, KJ, | P0708 |
| Glowczynska, R, | P0177, P1610 | Goodnough, LT, | OR268 |
| Gnatenko, D, | OR378 | Goodwin, C, | P2304 |
| Gnatenko, DV, | OR221 | Gopinadhan, M, | OR217 |
| Gnau, O, | P0284, P2076 | Gorbacheva, L, | P2349 |
| Gómez, RM, | P2360 | Gorbatenko, A, | P0080 |
| Goda, T, | P0675 | Gore, JM, | P1017, P1699 |
| Goecke, TW, | P2277 | Gore, T, | P2035 |
| Goemans, N, | P0321 | Gorelik, R, | OR104 |
| Goglidze, I, | OR149 | Gori, AM, | OR128, OR131, P0479,
P1095, P1123, P1282,
P1349, P1678, P1684A,
P1723, P1940, P2271,
P1995 |
| Gokoz, G, | P1002 | Goris, M, | P1995 |
| Gokudan, S, | OR209 | Gorman, R, | P2204 |
| Golanski, J, | P2146 | Gorospe, JR, | P2120 |
| Gold, SH, | P0836 | Gorp van, ECM, | P1995 |
| Goldberg, RJ, | P1699 | Gorter, G, | OR013, OR051, P0433,
P2090 |
| Goldenberg, NA, | P0546, P1251 | Goshen, G, | P1526 |
| Goldshmidt, O, | P1938 | Goss, D, | P1069 |
| Goldsmith, G, | P0198 | Gossage, J, | OR137, P1987 |
| Goldsmith, R, | P0228, P1997 | Gossage, JA, | P0786 |
| Golino, P, | OR370 | Gosselin, R, | P0318, P1571, P1955 |
| | | Gosselin, RC, | P1225 |

Goto, S, P0975, P1550, P1784, P2287
 Gotoh, M, P0799
 Goudemand, J, H08, H09, OR155, OR245, OR278, OR279, OR281, P0265, P0269, P0270, P0271, P0865, P1426, P1461, P1469, P2057, P2059
 Gough, PJ, SYM14
 Gougopoulou, D, P2286
 Gouin, I, P0426, P1623
 Gould, WR, P0648, P2309
 Gourdiar, AL, P2260
 Gouw, SC, OR093, P0219, P1417
 Govers-Riemslog, JWP, OR153, P1285, P1384
 Goyal, A, P1838
 Goycoolea, M, P0846, P1275, P1445, P1956
 Gozal, Y, OR296
 Gräber, S, OR330
 Grabarczyk, K, P0222
 Grabowski, EF, P0164, P0764, P1607
 Grabowski, M, P0177, P1610
 Gradwell, DP, P0474, P1070
 Graffigna, M, P1031
 Grafos, N, P0275, P1479
 Graham, G, OR017
 Grailhe, R, P0342
 Grailly, S, OR159
 Gram, J, OR235, P1965
 Grancha, S, P1246, P2073
 Grandini, A, P1831
 Grandone, E, OR094, OR094, OR197
 Granier, C, P0146, P1828
 Grant, A, P2144
 Grant, P, OR024, P1381
 Grant, PE, P1607
 Grant, PJ, OR021, OR022, OR049, OR314, P0286, P1950, P1986, P2079
 Grant, R, P2185
 Grant, RM, P0393
 Grass, L, P0773
 Grassilli, S, P0639
 Gravell, D, P2117
 Graw, J, P1829
 Gray, E, P0721, P1271, P1316, P1713
 Graybiel, AM, OR199
 Greaves, M, OR239, P0026, P0474, P0493, P1140, P2261, P2332, P2333
 Grebe, R, P2358
 Greber, S, P0259
 Greco, NJ, P1242
 Greco, V, P0726

Green, D, P0545, P1226, P1312, P2335
 Green, FR, OR112, P0587
 Green, R, P0674
 Greenfield, R, P0159, P0762, P0794, P1838
 Greenfield, RS, P0275, P1479, P1817
 Greensmith, H, P0853
 Greer, IA, OR239, P0493, P1091
 Gregor, J, P2284
 Gregorini, L, P1180
 Greinacher, A, OR081, OR217, P0504, P1727, SYM90
 Greinstetter, S, P0139
 Grenner, Y, P2351
 Grener, P s, P1631
 Gresele, P, OR283, OR370, P0701, P0964, P1548, P1573
 Grezchnik, E, P0864
 Griesshammer, M, P1324
 Griffin, B, P0060
 Griffin, JH, OR048, OR172, P1189, P1332, P1423, P1925, P2227, P2379
 Griffioen, A, P1471, P2067
 Griffith, B, P0522, P2166
 Griffiths, ABM, P0017, P1683
 Grigorescu-Sido, P, P0071
 Grigorjeva, M, P0819
 Grill, DE, P2267
 Grillari, J, P2141
 Grimaudo, S, P0527, P1278
 Grimsley, A, SOA03
 Gringel, E, P0097, P0885
 Gringeri, A, OR207, P0029, P0209, P0813, P0815, P1413, P1414, P1999
 Grinnemo, KH, P1748
 Gris, JC, P0068, P0634
 Grisar, D, P1537
 Gritsch, H, P0126, P1438
 Gritsenko, PG, P1393, P1985
 Gritti, D, OR375
 Grizzi, F, P0538
 Groeseneken, T, P0895
 Gröner, A, P0284
 Groner, A, P1582, P2076
 Gronowska, A, P0222
 Groot de, PhG, P0329
 Gros, A, P0696
 Grosley, M, P1296
 Gross, C, P1359
 Gross, PL, OR133, P1541
 Grossi, P, P1877
 Grossmann, R, P1555
 Grosso, S, P0615, P2046
 Grosso, SH, P0263
 Grüner, S, H12, OR254

- | | | | |
|------------------|---|-------------------|---|
| Grünewald, M, | OR101, P1324 | Hársfalvi, J, | P0334, P0963, P1972 |
| Gruber, A, | OR048 | Haas, A, | P2380 |
| Gruber, G, | P0878, P1474, P2068,
P2070 | Haas, F, | P0195, P0461 |
| Gruel, Y, | P0004, P0074, P0309,
SYM88 | Haas, FJLM, | P0463 |
| Grundt, H, | P0497, P1615, P1736 | Haas, S, | P1667 |
| Grundy, P, | P0874 | Haas, SK, | OR059, OR062 |
| Gücükoglu, A, | P0644 | Haas, TA, | P0332 |
| Günther, JU, | OR217 | Haasnoot, K, | P0194 |
| Guagnano, MT, | P1557, P1558 | Habart, D, | H08, H09, OR245,
OR278, OR279, P0265,
P0269, P0270, P0271,
P0865, P1461, P1469,
P2040, P2057, P2059 |
| Guanti, G, | P1454 | Haberichter, S, | P2075 |
| Guardascione, M, | P1881 | Haberichter, SL, | H09 |
| Guazzaloca, G, | OR175, OR177, OR179,
OR180, P0450 | Hack, C, | P0203 |
| Gudmundsen, TE, | P1644 | Hack, CE, | OR153, P0150, P0686,
P2150 |
| Guéguen, G, | P0472 | Hackeng, T, | P0380, P1350 |
| Guercini, F, | OR141 | Hackeng, TM, | P0134, P0729 |
| Guerrero, F, | P1027, P1613 | Hadders, M, | P2069 |
| Guerrero, JA, | P1267 | Hadjar, E, | P0678 |
| Guertin, S, | OR003, P0354, P1127,
P2118 | Hadley, A, | OR218 |
| Guglielmini, G, | P2037 | Haerberli, A, | OR291 |
| Guido, T, | OR283 | Haertel, S, | P2021 |
| Guidobono, R, | P1326 | Hafizi, S, | P1787 |
| Guiffre, A, | P2229 | Hafner, G, | P0097, P0885 |
| Guillaumon, AT, | P0560, P0594 | Hagedorn, A, | P1015, P1106, P1801 |
| Guillet, B, | OR300 | Hageman, G, | P0987 |
| Guilliatt, A, | P1409, P1984 | Hagemeyer, CE, | OR338 |
| Guilliatt, AM, | OR281, P0865, P0874 | Hagg, D, | P0082 |
| Guillin, MC, | P1470, P2066 | Hagger, D, | P0543 |
| Guillo, P, | P0364 | Haghighi, N, | P2243, P2300 |
| Guillonneau, S, | P0432 | Hagihara, K, | P1109 |
| Guimarães, AHC, | P1688 | Hagiwara, H, | OR223 |
| Guimarães, DAM, | OR392 | Hagiwara, N, | P2224 |
| Guinto, E, | P0439, P0487 | Hague, W, | P1683 |
| Guinto, ER, | P1479 | Hahn, CN, | OR298 |
| Guldiken, S, | P0275, P1817 | Haiyan, J, | OR182 |
| Gun-Munro, J, | P0586 | Haj manová, Z s', | P0255, P0682 |
| Gunneman, T, | P0443 | Hakobyan, N, | P0013 |
| Gunnemann, T, | P0709, P1309 | Halawani, S, | P1140 |
| Gunther, KE, | P1145 | Halder, C, | OR281 |
| Guo, R, | P1510 | Halimeh, S, | P0403, P1133, P2276 |
| Guo, Y, | P0437 | Hall, AJ, | P1259 |
| Gupta, PK, | H16, P0402, P1893 | Hall, L, | P0274 |
| Gurbel, P, | P2058 | Halldén, C, | P0274, P1330, P1466,
P1919 |
| Gurbel, PA, | P0362 | Halona, M, | P0240 |
| Gurfinkel, EP, | P0374, P0414, P0517,
P1014, P2167, P2205 | Hamacher, J, | P1638, P1639 |
| Gursel, T, | P1017 | Hamaguchi, M, | P0640 |
| Guschlbauer, E, | P1909 | Hamaguchi, N, | P2014 |
| Gustafsson, KM, | P1290 | Hamaia, S, | P0935 |
| Gustafsson, T, | P1258 | Hamato, N, | P0006 |
| Gutgobert, C, | P1748 | Hamdy, M, | P1868 |
| Guyer, K, | P1268, P1269, P1637 | Hamilton, JR, | P0890 |
| Guzal, M, | P0371, P0372, P0374,
P2167 | Hammer, F, | P0241 |
| | P0502 | Hammerstingl, C, | P0420 |

- | | | | |
|-----------------|--|----------------|--------------------------------------|
| Hammerstrøm, J, | P1814 | Harris, SL, | P0768, P0769 |
| Hampton, KK, | P0254, P2049 | Harrison, JA, | P2030 |
| Hamulyák, K, | OR043, P0459, P1055,
P1056, P1209, P1384 | Harrison, N, | P0520, P0853 |
| Hamulyak, K, | OR053, OR153, P0481,
P0752, P1285, P1856 | Harrison, P, | P0971, P1566, P1567,
P2365 |
| Han, F-Y, | P0674 | Harrison, SJ, | OR113 |
| Han, J, | P0970, P2160 | Hart, GW, | OR120 |
| Han, JH, | P1082 | Hartl, K, | P0197, P1228 |
| Han, Y, | P0499 | Hartmut, K, | OR219 |
| Handa, M, | P0335, P2086, P2087,
P2139, P2140 | Hartnett, ME, | P0599 |
| Handler, S, | P1290 | Hartwig, I, | P2336 |
| Hanfland, P, | P2114 | Hartwig, J, | SYM05 |
| Hankey, GJ, | P0948, P1846, SOA36 | Hartwig, JH, | OR115, OR116 |
| Hanley, JP, | P0844 | Harvald, T, | P2187 |
| Hann, A, | P1404 | Harvey, D, | OR055, OR056 |
| Hansell, S, | OR182 | Harvey, EJ, | P2361 |
| Hansen, H, | P1878 | Harvey, M, | P0179, P1404 |
| Hansen, HR, | OR053, P1285 | Hasebe, Y, | P0767 |
| Hansen, J, | P1013 | Hasegawa, H, | P2346 |
| Hansen, JB, | P0375, P1009, P1351,
P1695, P1779 | Hasegawa, M, | P1109, P1708 |
| Hansen, L, | P1289 | Hashemi, M, | OR281, P1469 |
| Hanslik, A, | P0259, P0860 | Hashiguchi, T, | OR170, P2382 |
| Hanson, SR, | OR048 | Hassan, S, | P2182 |
| Hanss, M, | P0847, P1984 | Hassell, KL, | P0062 |
| Hanss, MML, | P0174 | Hassler, M, | P0993 |
| Hansson, K, | P1787 | Hassouna, HI, | OR232 |
| Hansson, S, | P1739 | Hasstedt, SJ, | OR313, P1942 |
| Hantson, P, | P0241 | Hatakeyama, K, | P1178, P2287 |
| Hao, X, | P1748 | Hathaway, WE, | P1251 |
| Haond, C, | OR165 | Hathcock, J, | P1953 |
| Haque, W, | OR335, P0521, P2106 | Hathcock, JJ, | P1912 |
| Harada, N, | P1205 | Hattori, A, | OR001 |
| Harari, G, | P1277 | Hattori, K, | OR160, P0038 |
| Harbeck, N, | P0773 | Hattori, T, | P0163 |
| Harbeck-Seu, A, | P2058 | Hau, C, | P0959 |
| Harbour, SN, | P0892 | Hau, RCM, | P0573, P1753 |
| Harbrecht, U, | P2114 | Haubelt, H, | P0036, P0366, P1562,
P1755, P1859 |
| Hardikar, W, | P0124, P1913 | Hauch, O, | P1742, P1865 |
| Harding, K, | OR390 | Hauel, N, | OR336, P0507, P1703 |
| Hardy, AR, | P1496 | Hauer, A, | P0434 |
| Hardy, K, | P1341 | Haumer, M, | P0259 |
| Harenberg, J, | OR211, P1015, P1021,
P1022, P1106, P1705,
P1801, P2208 | Hauschner, H, | P1533 |
| Haridas, S, | P1259 | Hauser, H-P, | P0632 |
| Harish, VC, | P0061, P0171, P1513 | Hauser, HP, | OR161 |
| Harlan, J, | SOA17 | Havekes, LM, | OR225 |
| Harousseau, JL, | P0004 | Havik, S, | OR391 |
| Harper, AGS, | P0300 | Havik, SR, | OR393 |
| Harper, J, | P1038 | Hawk, S, | P0838 |
| Harper, MT, | P1552 | Hawkins, JR, | P1271 |
| Harper, P, | OR365, P1694 | Hawkins, M, | P1271 |
| Harper, PL, | OR302 | Hawkins, SF, | P1435 |
| Harris, AL, | P2192 | Hay, C, | P0874 |
| Harris, SB, | P0050 | Haya, S, | P0282, P0871, P1864,
P2013, P2073 |
| | | Hayakawa, M, | P1994 |
| | | Hayakawa, T, | P0767 |
| | | Hayashi, H, | P1488 |

- | | | | |
|--------------------|--|------------------------|---|
| Hayashi, K, | P0058, P0556 | Heller, C, | P1601 |
| Hayashi, M, | P1488 | Helley, D, | P2296 |
| Hayashi, T, | OR001, P0133, P0967,
P2088, P2178 | Hellstern, P, | P0036, P0366, P1562,
P1755, P1859 |
| Hayashida, H, | P2247 | Helm, CA, | OR217 |
| Hayes, K, | P1014 | Hemberger, M, | P0289 |
| Hayes, KK, | P0414 | Hemker, HC, | P0019, P0422, P0703,
P0889, P1483, P1688 |
| Hayes, PD, | P1553 | Hempel, M, | P0022 |
| Hayes, T, | OR303 | Henckel, T, | OR079, P0698, P0995,
P1088 |
| Hayoz, D, | P1138, P1173, P1770 | Henderson, RB, | OR118 |
| Hays, T, | P0138, P0546 | Hendler, MF, | P1676 |
| Hayward, CPM, | P1826 | Hendriks, D, | P0166, P1445, P1956 |
| Hazenberg, BPC, | OR098 | Hendrikson, SB, | H16, P0402 |
| He, J, | P0387 | Hendry, M, | P0179, P1404 |
| He, R, | P1356 | Henke, P, | OR135, P1981 |
| He, S, | P0152, P0183, P0705,
P1431 | Hennan, JK, | OR134 |
| He, SL, | P0153, P0558, P1354,
P1355, P1356, P0755,
P0756, P1947 | Hennig, J, | OR321 |
| He, XF, | P0153, P0558, P1354,
P1355, P0755, P0756,
P1947 | Henning, S, | P2342 |
| Healey, JF, | OR046 | Henny, ChP, | OR066, P0042 |
| Hebbel, RP, | P1075, P1772 | Henriet, C, | P0267 |
| Heck, AJR, | OR327 | Henriksson, CE, | P1879 |
| Hedger, S, | P2234 | Henschen, A, | OR114 |
| Hedner, U, | OR095, P1431 | Henschler, R, | P1591 |
| Heeb, MJ, | OR005, P0736 | Hepner, M, | P1018, P2050 |
| Heemskerk, JW, | OR051 | Heptinstall, S, | OR334, P0942, P0953,
P1498, P2134, P2149,
P2373 |
| Heemskerk, JWM, | OR254, OR343, OR344,
OR368, P0290, P0341,
P0906A, P0987, P1540,
P1584, P1701, P2170,
P2179 | Heras, I, | P1797 |
| Heeremans, J, | P2179 | Hérault, J-P, | OR141, P1790 |
| Heeringa, P, | P1208 | Herbel, P, | P1299 |
| Hegadorn, C, | OR280, P0264 | Herbert, J-M, | OR141, P1164, P1790 |
| Heger, A, | P0244 | Herchenhan, E, | OR211 |
| Heguilen, R, | P1676 | Heresi, GA, | P1763 |
| Heiden van der, Y, | P1995 | Hergenroeder, GW, | P2110 |
| Heifetz, S, | P1743 | Hermans, C, | P0241, P1092 |
| Heijnen, HFG, | P2090 | Hermesen, CC, | P0960 |
| Heikki, Vanannan, | P1912 | Hermesen, JT, | OR301 |
| Heil, SG, | OR129 | Hermus, AR, | OR127, P0480 |
| Heilmann, L, | OR059 | Hernández, F, | P0088 |
| Heim, M, | P1253 | Hernández-Espinosa, D, | OR003, OR234, P1206,
P1796, P1797 |
| Hein, L, | P1577 | Hernandez, F, | P1422, P1837 |
| Heine, GH, | P2351 | Hernandez, I, | H16 |
| Heine, RJ, | OR054, P1319 | Hernandez, J, | P0485 |
| Heinz, S, | P0035 | Hernanz, P, | P2045 |
| Heisteringer, M, | OR187 | Herrera, M, | P2045 |
| Heit, JA, | P0111, P0258, P0477,
P1076, P1308, P2173,
SOA15 | Herrmann, FH, | P0211, P0852, P1447 |
| Hekim, N, | P2152 | Hersemeyer, K, | OR261, OR386, P1744 |
| Hekmatian, N, | P1277 | Hertenstein, B, | P1506 |
| Helgeson, E, | P1246 | Hertfelder, H-J, | P2114 |
| | | Hervig, T, | P0108 |
| | | Herzog, A, | P0907 |
| | | Herzog, S, | OR332, P0130 |
| | | Herzog, W, | P0374 |
| | | Hess, O, | P1534 |
| | | Hess, Z, | P1636 |

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|------------------------|---|-------------------------|--|
| Hesse, B, | P1347 | Hirosawa, S, | P1917 |
| Hetherington, R, | P1037 | Hirose, M, | P0006, P0603 |
| Hetland, Ø, | P1696 | Hiroshi, K, | P0798 |
| Hevelow, ME, | P0373 | Hiroshi, S, | OR032, P1054 |
| Hevessy, Z, | P1556 | Hirschl, M, | OR361, P0438, P0581,
P0743, P1040 |
| Heward, J, | P1159 | Hirsh, J, | OR147, P1020 |
| Hewett, J, | P0349 | Hitos, K, | OR276, P0437, P2231,
P2233 |
| Hewitt, J, | P1847 | Hjemdahl, P, | P0355, P0436, P0604,
P0930, P1527, P2123,
P2147, P2226 |
| Hewson, J, | P2155 | Hjortoe, GM, | P0746 |
| Heyland, D, | P0200 | H awacz-Paszczyk, B I', | P1353 |
| Heysen, J, | P0804 | Hlusi, A, | P0809, P2168 |
| Hickey, MJ, | P0567 | Hmaguchi, M, | P1916 |
| Hicks, G, | P1213 | Ho, P, | P2360 |
| Hideo, W, | OR032, P1054 | Ho, PL, | P0714, P0883 |
| Hiebert, L, | P0505 | Hobbs, JE, | P1954 |
| Hiebert, LM, | P1032 | Hockman, S, | P2098 |
| Hiebinger, C, | P1231 | Hod, H, | P0950, P2145 |
| Higaki, N, | P2247 | Hoefelmann, C, | OR077 |
| Higashihara, M, | P2089 | Hoegge-de Nobel, E, | P1391 |
| Higgins, CF, | OR121 | Hoek, FJ, | P1319 |
| Higgins, J, | P0085 | Hoekstra, JBL, | OR050 |
| Higgins, LA, | OR371 | Hoekstra, MMCL, | P0539 |
| High, KA, | H07, P1317 | Hoffer, L, | P0655, P0656 |
| Higuchi, T, | P0684 | Hoffman, EP, | P2120 |
| Hijkata-Okunomiya, PA, | P1932 | Hoffman, M, | OR095, OR130, OR381,
SYM63 |
| Hilbert, L, | OR245, P0265, P0266,
P0865, P1461 | Hoffmann, C, | OR320 |
| Hilbrands, LB, | P0854 | Hoffmann, D, | OR208 |
| Hilgartner, M, | P0837 | Hoffmann, JN, | P0005 |
| Hill, F, | H08, H09, OR245,
OR278, OR279, OR281,
P0265, P0269, P0270,
P0271, P0865, P0874,
P1461, P1469, P2057,
P2059 | Hoffmann, U, | P0084 |
| Hill, FGH, | OR092, P0821, P1470,
P2066 | Hoffmeister, KM, | SYM07 |
| Hill, M, | OR113 | Hofmann, H, | OR321 |
| Hillarp, A, | OR277, P1330, P1919 | Hofstra, L, | P0557, P1350, P1489 |
| Hillege, HL, | OR086, P1735, P2289 | Hogg, K, | P0473, P2264 |
| Hilliard, P, | P0227 | Hogg, PJ, | OR075, SYM86 |
| Hillman, A, | P2133 | Hogwood, J, | P0105, P0734, P1713 |
| Hillmann, AG, | P1544 | Hohensinner, PJ, | P0580 |
| Hillmayer, K, | P0168 | Hohm, T, | OR208 |
| Hillyer, C, | OR268 | Höijer, P, | P1244, P1665 |
| Hilpert, M, | P1751 | Hoke, M, | P0743 |
| Hilton, A, | SYM04 | Holanda, RARR, | P0206 |
| Hilton, DJ, | SYM04 | Holbova, R, | OR297 |
| Hiltunen, L, | P0489 | Holdsworth, SR, | P0567 |
| Himbert, D, | P0364 | Holler, E, | P0593 |
| Hinder, M, | P0115, P0716 | Holly, P, | P0360 |
| Hirai, M, | P1053 | Holm, JP, | OR213, P1090 |
| Hirano, K, | P1313 | Holmberg, L, | H08, H09, OR245,
OR278, OR279, OR281,
P0265, P0269, P0270,
P0271, P0865, P1461,
P1469, P1739, P2057,
P2059, P2061 |
| Hirashima, K, | P1488 | Holmen, J, | P1814 |
| Hirayama, F, | P0603, P0967, P2178 | Holmen, LO, | P1661 |
| Hirmerová, J, | P1636 | | |
| Hirnle, T, | P1697 | | |
| Hirohata, K, | P1652 | | |

- | | | | |
|-------------------|---|--------------------------|---|
| Holmström, M, | OR384, P0238, P0828,
P0872, P2027 | Housman, DE, | OR199 |
| Holyoake, AJ, | P1523 | Houston, DS, | P0691, P1405, P1886 |
| Holzhauser, S, | P1555 | Houwing-Duistermaat, JJ, | OR110 |
| Homeister, J, | OR027 | Hovens, MMC, | P2249 |
| Honda, S, | OR026, OR285, P0057 | Hovy, L, | P2019 |
| Honda, Y, | P0511, P0512, P0612,
P1110 | Howard, I, | P0346 |
| Hong, JT, | P0596 | Howard, TE, | P0642 |
| Hong, T, | P1460 | Howarth, DJ, | P0401, P0475 |
| Hongli, W, | P1948 | Howarth, N, | P2260 |
| Hönig, M, | P0925 | Howe, B, | P0709, P1309 |
| Honnart, D, | P1049 | Howes, J-M, | P0286, P2079 |
| Hoogsteden, H, | P2257 | Hoylaerts, M, | P0987 |
| Hook, CC, | P0652 | Hoylaerts, MF, | OR029, P0321, P0331,
P0341, P0744, P0895,
P1701, P2082, P2135 |
| Hooper, W, | P2228 | | P1407, P1505 |
| Hooper, WC, | P0649, P1280, P1698,
P1926, P2375 | Hrachovinová, I, | P0585 |
| Hoppe, CC, | P0073 | Hrafnkelsdottir, T, | P0817 |
| Hoppe, K, | OR042 | Hristov, D, | P0438, P1040, P1935,
P2184 |
| Hoppener, MR, | OR066, P0042, P1057 | Hron, G, | P0584, P1327 |
| Hoppensteadt, D, | OR335, P0014, P0089,
P0110, P0518, P0519,
P0520, P0521, P0609,
P0710, P0782, P0991,
P1023, P1036, P1100,
P1118, P1120, P1177,
P1185, P1186, P1288,
P1337, P1599, P1611,
P1717, P1781, P1853,
P1872, P1899, P1902,
P1988, P2193, P2194,
P2312 | Hryszko, T, | P1358 |
| Hoppensteadt, DA, | P2334 | Hsiao, G, | P0338 |
| Horák, J, | P2212 | Hsu, CC, | P0772 |
| Horellou, MH, | OR085, P0180, P0490,
P0662, P1045, P1085,
P1262, P2297 | Hsu, M-Y, | P0822 |
| Hori, Y, | P0967, P2178 | Hsu, T-C, | P0579 |
| Horie, S, | P0727, P0784, P1325 | Hu X-H, | P0895, P1551, P1662,
P1701 |
| Horn, A, | P2187 | Hu, H, | OR225 |
| Horn, E, | P1245 | Hu, L, | OR078, P1153, P2344 |
| Horn, M, | OR016, P1876 | Hu, X, | P2269 |
| Horstman, L, | P0917, P0919, P0999 | HU, Y, | P1378 |
| Horstman, LL, | P1763 | Hu, YQ, | P1300 |
| Horter, S, | P0986 | Huan, W-M, | P1893 |
| Horton, S, | P0859, P1343, P1517 | Huan, X-m, | OR257 |
| Horvath, A, | OR303 | Huang, CL, | P1331 |
| Horvathova, D, | P0184, P0252 | Huang, CW, | OR221 |
| Hosch, S, | P1486 | Huang, EC, | P1331 |
| Hoshiba, Y, | P2287 | Huang, HC, | P1996 |
| Hoshino, H, | P1994 | Huang, J, | P1718 |
| Hosokawa, K, | P0513 | Huang, Q, | P0283 |
| Hotamisligil, GS, | P1633 | Huang, R-H, | P0338, P0944 |
| Houben, B, | P1384 | Huang, TF, | P0144 |
| Hough, C, | OR157, OR280, P0033,
P0264, P0690, P1232 | Huang, THW, | P1099 |
| Houng, A, | P1387, P1962 | Huang, W, | P0340 |
| | | Huang, WZ, | P0152 |
| | | Huang, X, | P1996 |
| | | Huang, X-p, | P1246 |
| | | Hubbard, AR, | OR017 |
| | | Hubbard, GP, | OR114 |
| | | Hubbs-T, D, | P1537 |
| | | Hubel, E, | OR264, P0561, P0580 |
| | | Huber, K, | P0419, P1019 |
| | | Hueb, WA, | P0061, P0171, P1513 |
| | | Huff, JD, | P2141 |
| | | Hugelier, K, | OR018, P2357 |
| | | Hughan, S, | OR014, P0902 |
| | | Hughan, SC, | P1216 |
| | | Hughes, GRV, | |

Hughes, P, P0145
 Hughes, Q, P0123
 Huh, HJ, P1221
 Huhle, G, P0084
 Huijgen, R, P0042
 Huisman, MV, H01, OR062, OR225,
 OR362, P0458, P0469,
 P2240, P2249, P2252
 Huisse, MG, P0364
 Huizinga, E, P2069
 Hulett, MD, P1193
 Hull, RD, P1667
 Hulstein, JJJ, OR028
 Humardewayanti Asdie, R, P1978
 Humphries, J, OR137, P0786, P1987
 Humphries, SE, P0105, P0734
 Humpich, M, P0020, P1849
 Hunault-Berger, M, P0004
 Hung, SH, P1502
 Hunt, BJ, P0423, P1725, P1952
 Hunt, J, P0781
 Huntington, J, OR004
 Huntington, JA, SOA11
 Huntscha, S, P0130
 Huntsman, DG, P1847
 Hunzelmann, N, P0831
 Huo, J, P0611
 Huppert, P, P2241
 Hurlen, M, P1029
 Hurley, BL, P0565, P1881
 Hurlstone, A, OR385
 Hursting, MJ, P0547, P1515
 Hurtaud-Roux, MF, P0921
 Husain Asdie, A, P1978
 Hutajulu, SH, P1978
 Huth-Kühne, A, OR036, P0056, P0232,
 P1830
 Hutson, SM, P0661
 Hutspardol, S, P2119
 Hutten, BA, OR031
 Huugen, D, P1208
 Hvitfeldt Poulsen, L, P0095, P0810
 Hyland, C, SYM0
 I ecki, Ml' z ., P1353
 Iaboni, S, P1027, P1733
 Iacobelli, M, P0593
 Iacone, R, P1966
 Iacoviello, L, OR194, P0670, P0988,
 P1372, P1776, P1861,
 P1966
 Iannaccaro, P, P0816, P1448
 Iannaccone, L, P0623
 Iatsenko, EA, P0236, P1737
 Ichihara, M, P1587
 Ichimura, K, OR228
 Ichinose, A, OR019, P1967, SYM50
 Ide, L, OR184

Ieko, M, P0024, P0202
 Ifrah, N, P0004
 Iglesias Varela, ML, P0614, P1806
 Ignjatovic, V, OR359, P0124, P0742,
 P0858, P0859, P1342,
 P1343, P1517, P1913
 Ihara, A, P1096
 Ihara, H, OR226, P1327
 Iijima, K, P1832
 Ikebe, M, P2089
 Ikeda, Y, P0280
 Ikeda, M, P2247
 Ikeda, S, P1383
 Ikeda, Y, P0335, P0966, P2139,
 P2140
 Ikejiri, M, P1488
 Ikemoto, Y, P0185
 Ilan, N, P1938
 Ilancheran, A, P1314, P1597
 Ilari, I, P2218
 Iles, DE, P1950
 Ilia, T, P1235
 Illiceto, S, OR366
 Illán, F, P1206
 Ilveskero, S, P1295
 Ilzecki, M, P1959
 Imamura, T, P1586, P2287
 Imamura, Y, P2088
 Imberti, D, OR175, P0450, P1648,
 P1650, P2188
 Inácio, RAS, P2235
 Inaba, H, P1256, P1832
 Inbal, A, OR023, OR297, P1277
 Ince, H, P1427
 Ing, C, OR360
 Ingelfinger, JR, P0164, P0764
 Ingerslev, J, H08, H09, OR132
 OR245, OR278, OR279,
 OR281, P0095, P0242,
 P0265, P0269, P0270,
 P0271, P0810, P0840,
 P0865, P1461, P1469,
 P2057, P2059
 Ingersoll, S, P0982, P2015
 Ingersoll, SB, OR287
 Inguenaud, C, P1267
 Iniesta, JA, P1374, P2048
 Innocenti, D, P1124, P2325
 Inoue, O, P0898, P1494, P2086
 Ioannidis, G, P0116, P0312, P0384,
 P1641, P2002, P2121,
 P2242
 Ionita, H, P0665
 Ionova, V, P0353, P1190, P1786
 P2201, P2364
 Iorio, A, H02, P0218, P0249
 P0257, P1027, P1613
 P1733, P2219

- lorio, AM, P1027
lotti, M, OR366, P0023, P1811
lqbal, A, P1911
lqbal, O, P0110, P0518, P0520,
P0521, P0782, P0991,
P1100, P1177, P1186,
P1288, P1337, P1599,
P1902, P2193, P2194,
P2312
Irani, M, P2165
Ireland, H, P0105, P0734
Iriundo, A, P2191, P2330
IRO, P2043
Isaac, G, P1530
Isermann, B, P0130
Isgro, F, P1755
Ishida, H, P1784
Ishida, N, P0784, P1325
Ishida, T, OR228
Ishihara, H, P0280, P0966
Ishii, H, P0684, P0727, P1193
Ishikawa, M, P2103
Ishiwata, A, OR183
Isobe, H, P1160
Isobe, K, P0511, P0512, P1110
Isobe, T, P0611, P2299, P2301
Isordia-Salas, I, P1184, P1287, P2182
Israels, S, P0227
Israels, SJ, P0896
Isturiz, M, P1761, P2132
Itakura, H, P0073
Italiano, J, SYM05
Itani, Y, P2247
Ito, T, OR170
Ivanciu, L, OR011
Ivanková, J, P0360
Ivaskevicius, V, OR187, P0261, P0525
Iversen, N, P1351, P1944
Ivey, L, P1289
Iwamoto, M, P0603
Iwamoto, T, P2103
Iwaniec, T, P0482, P1785, P1835,
P2258, P2317
Iwashita, M, P0966
Iyengar, SS, P1179, P1629, P1691,
P2316
Izraeli, S, P0325
Izui, S, P1164
Izzo, M, P0192
J, Jacoby, R, P1571
Jørgensen, M, P0444, P2210
Jackowski, R, P1697
Jackson, C, H14
Jackson, CL, P1752
Jackson, DE, P0892, P1529
Jackson, RD, P1310
Jackson, S, OR147, P1020, P1166
Jackson, SP, OR369, P1165, P2081
Jacob, L, P1235
Jacobs, DR, P1669
Jacobs, LG, P1121
Jacobs, M, OR189
Jacobs, N, P0960
Jacobs, R, P0421
Jacobson, A, OR308, P0709, P1145,
P1309, P1900, P2222
P0407
Jacobson, BF, P1251
Jacobson, L, P1375
Jacobsson, A, P1431
Jacobsson, EG, OR159, P0032, P0647,
P1843
Jacquemin, M, P2250, P2257
Jacques Michiels, J, P0077, P1237
Jadaon, MM, P1033
Jaeckel, J, P0920
Jaeken, J, OR309
Jaffer, AK, OR330
Jahnhen-Dechent, W, P1134
Jain, R, P2058
Jaiprakash, M, OR002, P0607
Jairajpuri, MA, P1609
Jakiela, B, P0891
Jakovljevic, V, P0952, P0956, P1109,
P1111, P1708, P2299
Jakubowski, JA, P2301
P0819
Jakunina, L, P0305
Jalagadugula, G, P2243, P2300
Jalali, F, P2283
James, A, OR280, P0264, P1848
James, PD, OR332, P1122
Jaminet, P, P2283
Jamison, M, P1326
Jan, R, P1320
Jan, TR, P1571
Janatpour, K, P1225
Janatpour, KA, P0318
Janatpur, K, OR071, P0294, P0341,
P1904
Jane, S, H15
Jang, SS, P1221
Jankovicova, D, P0852
Jankowski, M, P1250, P1252
Jansen, M, P0952
Janson, T, P0704
Janssen, HLA, P0281
Janssen, M, P0710
Jansson, L, P0142
Janz, K, OR390, P2355
Jardine, L, P0654
Jarvis, GE, P0981
Jaskolowski, S, P1884
Jason, Y, P1568, P1569, P1570
Jatene, FB, P0984

- | | | | |
|-----------------|---|-----------------------|--|
| Javen, D, | P2232 | Johansson, U, | P1172 |
| Jayandharan, G, | P0212, P2003 | Johnell, M, | P0759 |
| Jayasuriya, K, | P1456 | Johnsen, J, | P0578 |
| Jazebi, H, | P1389 | Johnson, A, | P0942, P2099, P2134 |
| Jazebi, M, | P1412 | Johnson, BD, | OR108, OR186 |
| Jeimy, SB, | P1826 | Johnson, C, | P0845 |
| Jenkins, A, | OR196 | Johnson, FL, | P2206 |
| Jennings, GM, | P0961, P1668, P2164 | Johnson, GJ, | P0302 |
| Jennings, I, | P0107, P1271, P1305,
P1898 | Johnson, J, | OR117, OR312, P0534 |
| Jennings, LK, | P1013 | Johnson, WD, | P0627 |
| Jerczynska, H, | P0555 | Johnston, A, | P1321 |
| Jern, C, | OR198, P0178 | Johnston, K, | OR374 |
| Jern, S, | P0585 | Johnston, L, | P0503 |
| Jeske, J, | P0109, P0110 | Johnston, M, | P0127, P0692, P0979,
P1774, P1957 |
| Jeske, W, | P0089, P1186 | Johnston, N, | P0117, P1357, P1620 |
| Jeske, WP, | P0918, P1599, P1899,
P2106, P2107, P2193,
P2292, P2311, P2312,
P2334 | Johnstone, BR, | P1130 |
| Jesmok, G, | P0173, P1370, P1844 | Jolicoeur, ME, | P1042 |
| Jespersen, J, | OR235, P1965 | Jones, C, | P0858, P1342 |
| Jessup, W, | OR231 | Jones, CD, | P0049 |
| Ji, Q, | P0930, P1527, P2123 | Jones, CI, | P0474, P1070, P1553 |
| Ji, S, | P0391, P2136 | Jones, DW, | P0027, P0720, P0768
P0769, P1339, P1897 |
| Ji, X, | P1508 | Jones, G, | P0200 |
| Jialan, S, | P1320 | Jones, JE, | P0240 |
| Jiang, F, | P0571 | Jones, RM, | P1607 |
| Jiang, L, | P2137 | Jones, S, | OR017 |
| Jiang, S, | P1662 | Jones, T, | P0478 |
| Jiang, X, | P1508 | Jong, G-P, | P0501 |
| Jiang, X-C, | P2308 | Jonson, B, | P1664 |
| Jie, AFH, | P1391 | Joø, GB, | P1879 |
| Jie, QZ, | OR181 | Jood, K, | OR198, P0178, P2151 |
| Jilma, B, | P0001, P0287, P0722,
P1874, P2142 | Joosen, AMCP, | P1483 |
| Jiménez, JJ, | P1763 | Jordan, R, | P0198 |
| Jimenez, V, | P0088, P1422 | Jordi, F, | P0072 |
| Jin, J, | P0905, P1581, P1501
P1507 | Jorfeldt, L, | P1067 |
| Jin, W, | P0369 | Jorge, DP, | P0978 |
| Jin, YR, | P0596, P1709 | Jörg, I, | P2208 |
| Jin, Z, | P1320 | Jorgensen, LN, | P2187, P2213 |
| Jinga, V, | P1196 | Jorquera, JI, | P1246 |
| Jinoch, P, | P1505 | Joseph, J, | P0617, P2099, P2137,
P2232 |
| Jobe, SM, | OR346 | Joseph, JE, | OR084, P0113, P0122
P0945, P1194 |
| Joerg, S, | OR182 | Joshi, AV, | P2035 |
| Johansen, P, | P0095, P0810 | Joshi, S, | P2270 |
| Johanson, S, | P1659, P1660 | Joshua, D, | P1883 |
| Johansson, AM, | P0274, P1330, P1466,
P1919 | Journeycake, JM, | P1000 |
| Johansson, H, | P0158 | Joutsi-Korhonen, L, | OR070, P2018 |
| Johansson, M-C, | P0355, P0436, P1527
P2147, P2226 | Jover, S, | P2045 |
| Johansson, MC, | P0604 | Joyal, F, | P1634 |
| Johansson, P, | P0145 | Jr, Wojtukiewicz, MZ, | P0718 |
| Johansson, PI, | P0683 | Jr, Bajaj, SP, | OR190 |
| | | Ju, MY, | P1221 |
| | | Jude, B, | OR155, OR317 |
| | | Judge, H, | P2163 |
| | | Judge, HM, | P0378, P2148
P2162 |

- Juhan-Vague, I, OR317, P0169, P0411, P1161, P1382, P2369, SYM41
- Juin, F, P0662
- Julian, J, OR088, OR147, P1020, P2237
- Jun, M, P0798
- Jun, T, P1549
- Jung, IS, P2290
- Jung, SM, P0343, P2087, P2088
- Jungbauer, A, P0139
- Junji, N, P1054
- Junqueira-de-Azevedo, ILM, P0714, P0883
- Jünschke, A, P1299
- Jurk, K, OR016, P0495, P0925, P1486, P1876
- Jürs, M, P0005
- Just, M, P0115, P0601
- Just, O, P0695
- Just, S, P0091, P0689, P1989, P2156, P2210
- Justice, MJ, OR167
- Juszynski, A, P1419
- Jy, W, P0919, P0999, P1763
- Kaabache, T, P1914
- Kaabachi, N, P1097
- Kaaja, R, P0489, P1684, P2214
- Kabakchieva, R, P0207, P0908, P1424
- Kabile, F, OR105
- Kabilyanskaya, V, P0136
- Kadar, J, P0256, P2336
- Kadar, JG, P0830
- Kadinskaya, M, P0528
- Kadir, R, P0225
- Kadir, RA, SYM74
- Käehne, M, P0003
- Kafsi, N, P1097
- Kagawa, K, P0800
- Kahan, T, P0355
- Kahn, M, OR067, P0894
- Kahn, ML, OR284
- Kahn, S, P0710
- Kahn, SR, P0440, P0533, P1634, P2237
- Kahner, BN, OR072
- Kahr, W, P1132
- Kahr, WHA, OR164, P1519
- Kahrizi, K, P0067, P1730
- Kaila, N, OR135, OR390, P2355
- Kaiser, B, P0014
- Kakkar, AK, OR059, OR062, P1078, P1590, P1671
- Kakkar, V, P0549, P0570, P1754, P2304
- Kakkar, VV, P0977, P1159, P1179, P1536, P1629, P1691, P1775, P2316
- Kaku, S, P1587
- Kalafatis, M, P1923
- Kalashnik, O, P1976
- Kalayanee, A, P1084
- Kale, S, P0913
- Kälebo, P, OR062
- Kalev, I, P0207, P0827, P0908, P1424
- Kalfa, M, P1468
- Kalk, S, P0221
- Kalkkinen, N, P1158
- Kallam, S, P0522, P2166
- Kalloway, S, P0402
- Kalume, DE, OR189
- Kalus, U, P1439, P1453
- Kam, A, P2231
- Kamae, T, OR285, P0897
- Kamata, T, P0335
- Kambayashi, J, P2098, P2131
- Kambayashi, JI, P0936
- Kameneva, O, OR378
- Kaminsky, P, P0019
- Kamnar, J, P0817
- Kamocka, M, P0120
- Kamphuisen, PW, OR273, P1059, P1397, P1596, P2188
- Kanagasundaram, V, P1165
- Kanakura, Y, OR285, P0897
- Kanalikova, K, P1854
- Kanazawa, K, P0202
- Kane, M, P1197
- Kane, WH, P1248, P1826
- Kaneider, NC, OR006, OR375
- Kaneko, M, P0369, P2086
- Kaneko, T, P2371
- Kanellakis, P, H15, P1193
- Kanellopoulou, G, P1050
- Kang, MS, P1081
- Kanjilal, S, P1179, P1629, P1691, P2316
- Kannan, M, P0913, P2058
- Kannemeier, C, OR100
- Kannicht, C, P1474, P2068, P2070
- Kanogsunthornrat, N, P0732
- Kanse, SM, OR200, OR386, P0523, P1168, P1744, P2342
- Kapadia, M, P0283
- Kaplinska, K, P0120
- Kaponides, G, P1960
- Kappelmayer, J, P0323, P0926, P1156, P1556
- Kappert, G, P0403, P2276
- Kapsimali, Z, P0526, P1475, P2315
- Kaptein, AA, P0065
- Kapustin, S, P2273

Kapustyanenko, LG, P1393
 Kara, A, P0196, P0792
 Karabasova, M, P1190, P1786, P2201
 Karafoulidou, A, P1050, P1937, P2023
 Karanam, S, H14
 Karasalihoglu, S, P0590
 Kardaras, F, P2286
 Karha, J, P0974
 Karida, A, P2286
 Karimi, M, OR122, P0536, P2323
 Karimipoor, M, P0213, P1411, P2005
 Karlsson, L, P0585
 Kárpáti, L, P0774
 Karpinski, G, P0177, P1610
 Karpman, D, P0849, P1739
 Karunakaran, D, P0894
 Kasak, L, P1158
 Kasal, E, P0682
 Kasamatsu, M, P0784
 Kasanuki, H, P2224
 Kasanuki, K, P2372
 Kashem, S, P2182
 Kashiwagi, H, P0897
 Kashiwagim, H, OR285
 Kashiwakura, Y, OR183
 Kasirer-Friede, A, SYM16
 Kassis, J, P1634
 Kastelein, JJP, OR050
 Kasthuri, RS, P0287
 Kasza, I, OR121
 Kataoka, N, P1932
 Kates, R, P0773
 Katherine High, SOA04
 Katherine, H, OR182
 Kato, H, OR026, OR285, P0897
 Katona, E, OR035, P1556
 Katona, É, P0774
 Katsarou, O, P1937, P2023
 Katsouras, CS, P0351
 Katsuaki, I, P0888
 Katsumi, A, P1488
 Katsuo, S, P1191
 Katz, B, OR139, P1537, P1890
 Katz, E, P0547
 Katz, N, P0885
 Kauffenstein, G, OR369
 Kauffman, J, P0260
 Kaufman, RJ, P0559
 Kaufmann, R, P1591
 Kaufmann, V, P1160, P1885
 Kaun, C, P0561, P0580
 Kaushal, V, OR041, P2374
 Kavakli, K, P1909, P2024
 Kaveri, SV, P1828
 Kavitha, ML, P2004
 Kawada, S, P0789
 Kawahara, K, P2382
 Kawaja, M, P1404

Kawakami, A, P0513
 Kawamoto, R, P1586, P2287
 Kawamoto, T, OR166, P1096
 Kawanishi, T, P0767
 Kawao, N, P0789
 Kawasaki, T, P0556
 Kawasugi, K, P0799
 Kayed, R, OR327
 Kazarian, T, P0013
 Kazi, Z, P1570
 Kazumi, H, P1549
 Kearon, C, OR088, OR147, OR312, P0534, P1020, P2237, SYM39
 Keeling, D, OR092, P0874
 Keeling, DM, P0224
 Keen, JN, P0286, P2079
 Keenan, J, P1098
 Keenan, R, P0873, P1435, P2066
 Keeney, S, P0874
 Kehrel, BE, OR016, P0495, P0925, P1486, P1876
 Keidan, I, P1526
 Keiji, M, P1549
 Keijzer, M, P0480
 Keiper, T, OR261
 Keith, JC, Jr, P2355, OR390, P1287
 Keiya, O, P0798
 Kekomaki, R, OR222
 Kelleher, AA, P1306
 Keller, F, OR211
 Keller, TT, P0150, P0685, P0687, P1878
 Kellerman, DJ, P2206
 Kelly, R, P0940, P1772
 Kemball-Cook, G, OR260, OR389, P0223, P0761
 Kemkes-Matthes, B, OR059, OR211, OR386, P0099, P0523
 Kemp, B, OR196
 Kempfer, A, P0276, P1003, P1473
 Kempfer, AC, P0388, P0863
 Kempton, CL, SOA03
 Ken, F, P2075
 Ken, I, P1054
 Ken, S, P0798
 Kenderova, V, P0908, P1424
 Kenet, G, P0637, P1312, P2024, P2051
 Kenis, H, P0557, P1350
 Kennedy, A, P0428, P1614, P1115, P1116, P1714, P2305
 Kennedy, G, P0091, P2156
 Kenny, D, P1544
 Kerachian, MA, P2361
 Keren, G, OR139
 Kerényi, A, OR035, P0323
 Kerjaschki, D, P1751

Kerkhofs, M, P0928
 Kerrigan, SW, P0328
 Kerschen, EJ, H16
 Kershaw, G, P1883, SYM86
 Kfsik, J, P1353
 Kesik, J, P1959
 Kessel, C, OR034, OR191, P0201, P0738, P1929
 Kessler, C, OR268
 Kettenmann, E, P0084
 Keuren, J, P0380
 Keuren, JFW, P0386, P1584, P2179
 Kevin Hayes, K, P2205
 Kevin, B, P1569, P1570
 Kevin, H, P1568, P1569, P1570
 Kevin, N, P1580
 Kevin, PB, P1568
 Key, NS, OR371, P0287, P0823, P0916, P1075, P1280, P1361
 Khachigian, LM, OR388, P0595, P1150, P1193
 Khair, K, P0544
 Khalid, K, P1611
 Khalil, R, P1314, P1597
 Khamis, H, P1798
 Khan, A, P0225
 Khan, M, P2182
 Khan, MM, P0163, P1184
 Khan-Ahamad, H, P2005
 Khanduri, U, P2117
 Khanin, MA, P0740, P0758, P1301, P1302, P1803
 Khaw, KT, P0150
 Khoo, KH, P1805
 Khorana, AA, OR058, P1598
 Khrenov, A, OR248, P1840
 Khurana, R, P1193
 Khwaja, H, P0587
 Kiatchoosakun, S, P1279
 Kibirev, VK, P1933
 Kiechle, M, P0773
 Kieffer, N, OR105, P0333, P2125
 Kienast, J, P0005
 Kiernan, A, P0983
 Kierulf, P, P1879
 Kiesewetter, H, P1439, P1453
 Kile, BT, OR167
 Kim, CH, P1502
 Kim, H, P0970, P2160
 Kim, HS, P1082, P1679
 Kim, H-S, P1332
 Kim, J, P1865
 Kim, JH, P0596
 Kim, M, P0970, P2160
 Kim, NK, P1081, P1082, P1679
 Kim, OJ, P1082, P1679
 Kim, PY, OR096, OR250

Kim, S, P0905, P1581
 Kim, WC, P1082, P1679
 Kimmel-Jehan, C, P1970, P2033
 Kimura, R, P0057
 King, GC, OR333
 King, J, P0318, P1955
 King, L, OR178
 King, SM, P0383
 Kini, RM, P0053, P0154, P1850, P1851, SYM72
 Kirchmaier, C, P0256
 Kirchmaier, CM, P0345, P0697, P0812
 Kireev, D, P0976
 Kirgner, I, P1890
 Kirk, G, P1569
 Kirkham, FJ, OR355
 Kiseleva, E, P2349
 Kish, M, P0931
 Kishiwada, M, P0133
 Kisiel, W, P0718, P1918
 Kiss, C, OR123, P0323
 Kiss, RG, P1093
 Kitazawa, T, OR160
 Kitchen, DP, P0107, P0884, P1299, P1305, P1898
 Kitchen, S, P0107, P0605, P0708, P0884, P1271, P1299, P1304, P1305, P1307, P1898
 Kitzmüller, E, P0259
 Kjaer, A, P1347
 Kjalke, M, P0145, P0748
 Klaeffling, C, P0098, P0365, P1087, P2217
 Kläffling, C, P1202
 Klamroth, R, P0256
 Klarmann, D, OR191, P0021, P0261
 Klement, P, P1201, P1963
 Klengel, R, P0655, P0878
 Klerk, CPW, P0397
 Klime, D s, P1407
 Kling, D, P1166
 Klinge, J, P1457
 Klingebiel, T, OR191, P0602, P1204, P1795, P1929
 Klingenberg, O, P1879
 K oczko, J I, P0031
 Kloczko, J, P0170, P1368, P1392, P2385
 Kluft, C, P1689, P2281
 Kluger, Y, OR379
 Kluijtmans, LAJ, OR129
 Kluin-Nelemans, HC, P0066, P0102, P0663, P1265, P1855, P1857, P1892
 Klukowska, A, P0044, P0641, P0833, P1231

Knabb, R, P0162, P1211, P1318, P1852
 Knapp-Clevenger, R, P0546
 Knaub, S, P1795
 Kniessel, M-J, P1743
 Knight, B, OR049
 Knight, S, P0245, P1315
 Knirsch, W, P1068
 Knobe, K, P0046
 Knobe, KE, P1257
 Knobel, M, P0085
 Knobl, P, OR270
 Knöbl, P, P0722
 Knoefel, W, P1486
 Knoefler, R, P1033
 Knöfler, R, P1601
 Knudsen, JB, P2112
 Ko, Y, P2150
 Kobayashi, H, P0914
 Kobayashi, T, OR001, OR032, P0947, P2371
 Kobayashi, Y, P1011
 Koch, C, P0181
 Koch, D, P0848
 Kochanowicz, JNM, P1605
 Kochman, J, P1610
 Kocyigit, I, P1002
 Koehler-Vajta, K, P0832
 Koelling, T, P0372
 Koenen, RR, P0729
 Koerber, JM, P1143, P1144, P1516
 Kofinas, G, P1367
 Koga, S, P0087, P1993
 Koga, Y, P1652
 Kogan, A, P0196, P0792
 Kogerman, P, P1158
 Koh, T, P0767
 Köhler, H, P2351
 Kohlerova, S, P1998
 Koide, T, OR330, P0185, P0513, P0791, P0024, P0202
 Kojima, H, OR166
 Kojima, T, OR160, P1488, P1916
 Kokaji, T, P1256
 Kokame, K, OR026, P0119, SYM87
 Kokubo, Y, P0057
 Kolaitis, N, P1234, P1642
 Kolaitis, NI, P0448
 Kolesnikova, IN, P1393, P1985
 Kolev, K, P0790
 Kollander, R, P1772
 Kollars, M, P0992, P1935, P2184
 Koltakov, A, P0929
 Komanasin, N, P1279
 Komáromi, I, OR123
 Komine, R, P0966
 Komisarenko, S, OR009
 Komisarenko, SV, P1393, P1985

Komitopoulou, A, P0526, P2315
 Komorowski, J, P0555
 Komrska, V, P1231, P1407, P1998
 Komvilaisak, P, P0393, P1214, P1425, P2185
 Kondashevskaya, MV, P2181
 Kondo, M, P0006, P0603
 Konecny, F, P1201
 Konetschny, C, P0907
 Kong, M, P1370
 Kong, MC, P1711
 Königs, C, OR034, OR191, P0738, P0201, P1929
 Konno, K, P0883
 Kono, R, P0553
 Konrad, I, OR042, OR322
 Konstantoulas, CJ, P0734
 Konupcikova, K, P1866
 Kootstra, NA, OR247
 Kopec, I, P1815
 Kopecky, E-M, P0139
 Kopecky, O, P2284
 Kopp, F, P1161
 Kopenhagen, K, P1022
 Kopylov, KG, P0834
 Koraka, P, P0203
 Korczowski, B, P0849
 Kornecki, E, P0339, P0409, P2308
 Kornek, G, P2184
 Kornhauser, D, P1318
 Kornowski, R, P2386
 Korporaal, SJA, OR013, P1578
 Korsgren, O, P0158, P1172, P1804
 Korte, W, P0118, P0458, P0469, P0776, P2240
 Korteweg, FJ, OR213, P1090
 Korzh, I, P0577, P0777, P2291
 Korzh, O, P0576, P1174, P1188, P1780, P1921
 Koschelnic, Y, P1157
 Koscielny, J, P1439, P1453
 Koshihara, K, P0800
 Kostantinidis, K, P1617, P2197
 Kostka, H, P1233
 Kotchuev, G, P1188
 Kotev-Emeth, S, P0929, P2145
 Kotitschke, R, OR045
 Kotsi, P, P1050, P1937, P2023
 Kotsia, A, P0351
 Kottke-Marchant, K, OR303, P0974, P2204
 Kouassi, D, P0169
 Kouivaskaia, D, P1841
 Kouides, PA, OR383, P0258, P0973
 Koumbaniou, CH, P0448
 Kounas, S, P2286
 Kouramba, A, P2023
 Kovac, M, P1030, P1674
 Kovacs, MJ, P2237

- | | | | |
|----------------------|--|---------------------------|---|
| Kovalchuk, O, | P1945 | Krishnaswamy, S, | OR192, OR251, P1922 |
| Kovalenko, T, | P1681 | Krisinger, MJ, | P0653 |
| Kovalenko, TF, | P1083 | Kristensen, AT, | P0230, P0748 |
| Kovaleva, V, | P1588 | Kristensen, JB, | P1347 |
| Kovárník, T, | P2212 | Kristensen, SR, | OR235, P0003 |
| Kowal, K, | P1372 | Kristin, MS, | P1326 |
| Kowalski, H, | P1751 | Kristoffersson, AC, | P1739, P2061 |
| Koyama, T, | OR100, P1917 | Kritharides,#2;,#2;,, | P1146 |
| Koyano, K, | P1053 | Kriukelyte, D, | P2288 |
| Kozak, M, | P1908 | Kílová, P ř, | P0682 |
| Koziol, J, | P0313 | Kroemer, G, | OR165 |
| Kozlova, T, | P0484, P1137, P1663 | Kroese, A, | P1778 |
| Kozlova, TV, | P1083 | Krogh, V, | OR194 |
| Kozubski, W, | P0344 | Kroll, H, | OR222, P0307, P0910,
P0911, P2113 |
| Kozuma, Y, | OR166 | | |
| Kraan, MC, | P1777 | Kroon, AA, | P1384 |
| Krahn, EL, | P1727 | Kroon-Batenburg, LMJ, | P0563 |
| Krajewska, U, | P0120 | Kruchinsky, N, | P1274, P1958 |
| Kralisz, U, | P0381, P0574 | Kruchinsky, NG, | P0568 |
| Kramkowski, K, | P2180 | Krueger, B, | P0962 |
| Kranenburg, O, | OR327 | Kruglov, O, | P1511 |
| Kranendonk, SE, | P0464 | Kruip, MJHA, | P1658, P2252 |
| Krapp, T, | OR023 | Kruihof, CJ, | P1625 |
| Kraska, Y, | P0022 | Kruihof, EKO, | OR140, OR144, P1223,
P1224, P1770 |
| Krasnobryzha, I, | P0188 | | |
| Krasnokutskiy, S, | P1174, P1188, P1780 | Kruppenbacher, J, | P0256 |
| Kraus, AE, | P0599 | Krüssel, JS, | OR215, P2127 |
| Krause, M, | OR034 | Kubáková, K c̃, | P0255 |
| Krause, M, | P0201, P0232, P0311,
P0636, P0697, P0812,
P1005, P1395, P2019 | Kubatiev, A, | P0483, P1680, P2274,
P2275 |
| Kraut, EH, | P1443 | | P0360 |
| Kravets, V, | P2026 | Kubisz, P, | P1105, P1704 |
| Kr ová, V c̃, | P1407 | Kubitza, D, | P0603 |
| Krcova, V, | P0809, P2168 | Kubo, Y, | P1139 |
| Krebs, VLJ, | P0673 | Kuchler, H, | P0078 |
| Krejza, J, | P1605 | Kuczynska-Zardzewialy, A, | P1421 |
| Kremer Hovinga, J, | P0915 | Kudryk, B, | OR114 |
| Kremer Hovinga, JA, | P0849, P1139, P1740,
SOA33 | Kueter, TJ, | P1013 |
| Kremmidiotis, G, | OR298 | Kuffner, T, | P2137 |
| Kretschmer, V, | P0885 | Kuhl, P, | P1162 |
| Kretzschmar, E, | P2113 | Kuhle, S, | P0392 |
| Kreutz, R, | P0362, P0374, P0517,
P2205 | Kuhlich, E, | P1033, P1233 |
| Kreutz, Y, | P0362, P0414, P1014,
P2167 | Kuijpers, MJE, | OR254, OR343, P0290,
P0906A, P2170 |
| Kreutzenbeck, HJ, | P0508 | | |
| Kreuz, W, | OR034, OR191, P0021,
P0201, P0261, P0602,
P0738, P0812, P1204,
P1795, P1929 | Kuint, J, | P2051 |
| Kreymborg-Grosse, K, | P0995 | Kuiper, J, | P0434 |
| Kriaa, A, | P1097 | Kuipers, S, | OR289 |
| Kriegbaum, B, | P0175 | Kuipers, S, | P1657 |
| Krijanovski, Y, | P2347 | Kuipers, S, | P2256 |
| Krimer, A, | P1473 | Kuliopulos, A, | OR015 |
| Krishnamoorthy, R, | P2206 | Kulkarni, R, | OR383, P0052, P0258,
P0973, P1460, P2037 |
| | | | P1165 |
| | | Kulkarni, S, | P1263 |
| | | Kulle, B, | P0181 |
| | | Kullmer, M, | P0079, P0783, P1352 |
| | | Kulwas, A, | P0512 |
| | | Kumada, T, | P0394 |
| | | Kumar, KR, | |