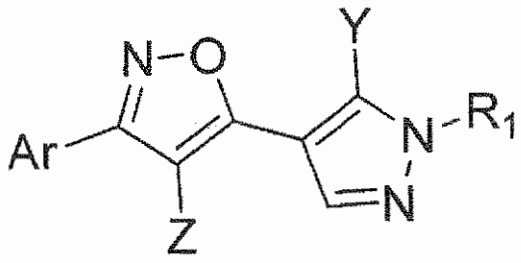
CLAIMS:

1. A compound of the general formula (I)



formula (I)

and pharmaceutically acceptable salts or solvates thereof,
wherein

R^1 is aryl, heteroaryl, cycloalkyl, heterocyclyl or alkyl, which can be substituted by one or more substituents R';

Ar is aryl, cycloalkyl, heterocyclyl or heteroaryl, which can be substituted by one or more substituents R';

Z is aryl, heteroaryl, cycloalkyl or heterocyclyl, which can be substituted by one or more substituents R';

Y is H, halogen, haloalkyl, alkyl or an alkylester, which can be substituted by one or more substituents R';

R' independently represents H, $-CO_2R''$, -CONHR'', -CR''O, $-SO_2N(R'')_2$, $-SO_2NHR''$, -NR''-CO-haloalkyl, $-NO_2$, -NR''-SO_2-haloalkyl, -NR''-SO_2-alkyl, $-SO_2$-alkyl, -NR''-CO-alkyl, -CN, alkyl, cycloalkyl, aminoalkyl, alkylamino, alkoxy, -OH, -SH, alkylthio, hydroxyalkyl, hydroxyalkylamino, halogen, haloalkyl, haloalkoxy, amino, heterocyclyl, aryl, haloaryl, haloarylalkyl, arylalkyl or heteroaryl;

R'' independently represents H, haloalkyl, hydroxyalkyl, amino, alkoxy, -N=C(R')$_2$, -NR'-CO-R', -CR'O, -CO_2R', alkyl, cycloalkyl, aryl, haloaryl, haloarylalkyl, heteroaryl, heterocyclyl, arylalkyl or aminoalkyl, which are optionally substituted by one or more substituents R';

2. A compound according to claim 1, wherein

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

R^1, Z, Y, R' and R'' are as defined in claim 1.

3. A compound according to claim 1, wherein

R^1 is aryl or heteroaryl, which can be substituted by one or more substituents R'

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

Z, Y, R' and R'' are as defined in claim 1.

4. A compound according to claim 1, wherein

R^1 is aryl or heteroaryl, which can be substituted by one or more substituents R'

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

Z is aryl, cycloalkyl, heterocyclyl or heteroaryl, which can be substituted by one or more substituents R';

Y is H, halogen, haloalkyl, or alkyl, which can be substituted by one or more substituents R';

R' independently represents H, -CO_2R'', -CONHR'', -CR''O, -SO_2N(R'')$_2$, -SO_2NHR'', -CN, alkyl, cycloalkyl, aminoalkyl, alkoxy, -OH, , hydroxyalkyl,

halogen, haloalkyl, haloalkoxy, amino, heterocyclyl, aryl, haloaryl, haloarylalkyl, arylalkyl or heteroaryl;

R'' independently represents H, haloalkyl, hydroxyalkyl, amino, alkoxy, -N=C(R')$_2$, -NR'-CO-R', -CR'O, -CO_2R', alkyl, cycloalkyl, aryl, haloaryl, haloarylalkyl, heteroaryl, heterocyclyl, arylalkyl or aminoalkyl, which are optionally substituted by one or more substituents R'

5. A compound according to claim 1, wherein

R^1 is aryl or heteroaryl, which can be substituted by one or more substituents R';

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

Z is aryl, cycloalkyl, heterocyclyl or heteroaryl, which can be substituted by one or more substituents R';

Y is H, halogen, haloalkyl, or alkyl, which can be substituted by one or more substituents R';

R' independently represents H, -CO_2R'', -SO_2N(R'')$_2$, -SO_2NHR'', -CN, alkyl, alkoxy, -OH, hydroxyalkyl, halogen, haloalkyl, haloalkoxy, heterocyclyl, aryl, haloaryl, haloarylalkyl, arylalkyl or heteroaryl;

R'' independently represents H, haloalkyl, hydroxyalkyl, amino, alkoxy, -N=C(R')$_2$, -NR'-CO-R', -CR'O, -CO_2R', alkyl, cycloalkyl, aryl, haloaryl, haloarylalkyl, heteroaryl, heterocyclyl, arylalkyl or aminoalkyl, which are optionally substituted by one or more substituents R'

6. A compound according to claim 1, wherein

R^1 is aryl or heteroaryl, which can be substituted by one or more substituents R';

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

Z is aryl, cycloalkyl, heterocyclyl or heteroaryl, which can be substituted by one or more substituents R';

Y is H, halogen, haloalkyl, or alkyl, which can be substituted by one or more substituents R';

R' independently represents H, $-CO_2R''$, -CONHR'', -CR''O, $-SO_2N(R'')_2$, $-SO_2NHR''$, -CN, alkyl, alkoxy, -OH, halogen, haloalkyl or haloalkoxy,;

R'' independently represents H, haloalkyl, hydroxyalkyl, amino, alkoxy, $-N=C(R')_2$, -NR'-CO-R', -CR'O, $-CO_2R'$, alkyl, cycloalkyl, aryl, haloaryl, haloarylalkyl, heteroaryl, heterocyclyl, arylalkyl or aminoalkyl, which are optionally substituted by one or more substituents R'

7. A compound according to any of claims 1 to 6, wherein

R^1 is selected from the group comprising phenyl which is optionally substituted by one or more substituents individually selected from trifluoromethyl, fluorine, chlorine, bromine, nitro, NH_2, -CN, $-NHCO-C_{1-4}$-alkyl, methoxy, C_{1-4}-alkyl, $-SO_2NH_2$, or $-SO_2NH-C_{1-4}$-alkyl; pyridyl which is optionally substituted by one or more of the aforementioned substituents for phenyl; pyrimidyl which is optionally substituted by one or more of the aforementioned substituents for phenyl; thienyl which is optionally substituted by one substituent $-COO-C_{1-4}$alkyl; 1,1-dioxo-tetrahydrothienyl, 2,2,2-trifluoroethyl, isopropyl, isobutyl, 2-methoxyethyl, tetrahydropyran-4-ylmethyl, 2-morpholinoethyl, pyridin-2-ylmethyl and tetrahydrofuran-2-ylmethyl;

Ar is selected from the group comprising phenyl and pyridyl, which can be substituted by one or more substituents independently selected from fluorine methoxy or chlorine;

Z is selected from the group comprising phenyl or heteroaryl, which can be substituted by one or more substituents selected from the group comprising C_{1-4}alkyl, C_{1-4}haloalkyl, hydrogen, hydroxyl, C_{1-4}alkoxy, and C_{1-4}alkoxycarbonyl;

Y is selected from the group comprising H, trifluoromethyl and methoxycarbonyl.

8. A compound according to any of claims 1 to 6, wherein

R^1 is selected from the group comprising phenyl which is optionally substituted by one or more substituents individually selected from fluorine, chlorine, bromine, nitro, NH_2, -CN, -NHCO-C_{1-4}-alkyl, methoxy, t-butyl, $-SO_2NH_2$, or $-SO_2NH$-isopropyl; pyridyl; pyrimidyl which is optionally substituted by one or more substituents selected from methyl or trifluoromethyl; thienyl which is optionally substituted by one substituent–COO-methyl, 1,1-dioxo-tetrahydrothienyl, 2,2,2-trifluoroethyl, isopropyl, isobutyl, 2-methoxyethyl, tetrahydropyran-4-ylmethyl, 2-morpholinoethyl, pyridin-2-ylmethyl and tetrahydrofuran-2-ylmethyl;

Ar is selected from the group comprising phenyl, 2,6-difluorophenyl, 2-chloro-6-fluorophenyl, 2-chloro-6-methoxyphenyl, 2-fluorophenyl, 2-chlorophenyl, 2,6-dichlorophenyl, 2,4-dichlorophenyl, 3-fluoropyridin-4-yl, 3,5-dichloropyridin-4-yl and 3,5-difluoropyridin-4-yl;

Z is selected from the group comprising phenyl, tetrazolyl, thiazolyl, 1,3,4-oxadiazolyl, oxazolyl, 1,3,4-thiadiazolyl, furanyl, and thiophenyl, which can be substituted by one or more substituents selected from the group comprising methyl, trifluoromethyl, hydrogen, hydroxyl, methoxy, and methoxycarbonyl and ethoxyoxycarbonyl;

Y is selected from the group comprising H, trifluoromethyl and methoxycarbonyl.

9. A compound according to any of claims 1 to 6, wherein

R^1 is selected from the group comprising phenyl, 2-fluorophenyl, 2-methoxyphenyl, 2-chlorophenyl, 2-bromophenyl, 2-nitrophenyl, 2-aminophenyl, 4-fluorophenyl, 4-tertbutylphenyl, 3-fluorophenyl, 3-chlorophenyl, 3-cyanophenyl, 3-acetamido-phenyl, 2-acetamido-phenyl, 3-aminosulfonyl-phenyl, 3-(isopropylamino)sulfonyl-phenyl, 3-nitrophenyl, 3-aminophenyl, 2,4-difluorophenyl, 3,5-difluorophenyl, 3,5-dichlorophenyl, 2,3,5,6-tetrafluorophenyl, 2-pyridyl; 3-pyridyl; 4-pyridyl; 4- trifluoromethyl-pyrimid-2-yl, 2,6-dimethyl-pyrimid-4-yl, 2-methoxycarbonyl-thien-3-yl, 1,1-dioxo-tetrahydrothienyl, 2,2,2-trifluoroethyl, isopropyl, isobutyl, 2-methoxyethyl, tetrahydropyran-4-yl-methyl, 2-(morpholin-4-yl)-ethyl, and tetrahydrofuran-2-yl-methyl;

Ar is selected from the group comprising phenyl, 2-chloro-6-fluorophenyl, 2-fluorophenyl, 2-chlorophenyl, 2,6-dichlorophenyl, 2,4-dichlorophenyl, 3-fluoropyridin-4-yl, and 3,5-difluoropyridin-4-yl;

Z is selected from the group comprising 1H-tetrazol-5-yl, 4-methylthiazol-2-yl, thiazol-2-yl, 5-(trifluoromethyl)-1,3,4-oxadiazol-2-yl, 5-methyl-1,3,4-oxadiazol-2-yl, 1,3,4-oxadiazol-2-yl, oxazol-5-yl, 4-hydroxy-thiazol-2-yl, 4-methoxy-thiazol-2-yl, 5-methyl-1,3,4-thiadiazol-2-yl, 1,3,4-thiadiazol-2-yl, furan-3-yl, furan-2-yl, thiophen-3-yl, phenyl, 4-methoxycarbonyl-thiazol-2-yl, and 4-ethoxycarbonyl-thiazol-2-yl;

Y is selected from the group comprising H, trifluoromethyl and methoxycarbonyl.

10. A compound according to any of claims 1 to 10, wherein the compound is selected from the group comprising the compounds of examples 4, 5, 6, 7, 8, 10, 11, 12, 14, 16, 17, B-2, B-3, B-4, B-5, B-6, B-7, B-8, B-9, B-10, B-21, B-22, B-24, B-25, B-26, B-27, and B-30 of the present invention.

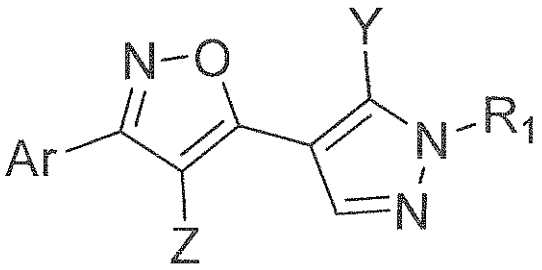
11. A compound according to any of claims 1 to 10 for use as a medicament.

12. A compound according to any of claims 1 to 10 for use in the treatment of a disease or indication is selected from the group consisting of psoriasis, psoriatric arthritis, autoimmune thyroiditis, Grave's disease, rheumatoid arthritis, vitiligo, Crohn's disease, ulcerative colitis, inflammatory bowel disease, ankylosing spondylitis, diabetes type I, multiple sclerosis, celiac disease, systemic lupus erythematosus, uveitis, Behcet disease, atopic dermatitis, Lichen planus, Sjögren's syndrome, spinal disc herniation, acne, Graft-versus-Host-Reaction, Host -versus-Graft-Reaction and osteoarthritis

13. Use of the compounds of the formula (I) as defined in any of claims 1 to 10, and of their pharmacologically tolerable salts in the manufacture of a medicament for use in treatment of a disease or a therapeutic indication in which the inhibition of interleukin-17 (IL-17) and/or Interferon-γ (INF-γ) is beneficial.

14. The use of claim 12 wherein the disease or indication is selected from the group consisting of psoriasis, psoriatric arthritis, autoimmune thyroiditis, Grave's disease, rheumatoid arthritis, vitiligo, Crohn's disease, ulcerative colitis, inflammatory bowel disease, ankylosing spondylitis, diabetes type I, multiple sclerosis, celiac disease, systemic lupus erythematosus, uveitis, Behcet disease, atopic dermatitis, Lichen planus, Sjögren's syndrome, spinal disc herniation, acne, Graft-versus-Host-Reaction, Host -versus-Graft-Reaction and osteoarthritis

1. A compound of the general formula (I)



formula (I)

and pharmaceutically acceptable salts or solvates thereof,
wherein

R^1 is aryl, heteroaryl, cycloalkyl, heterocyclyl or alkyl, which can be substituted by one or more substituents R'

Ar is aryl, cycloalkyl, heterocyclyl or heteroaryl, which can be substituted by one or more substituents R';

Z is aryl, heteroaryl, cycloalkyl or heterocyclyl, which can be substituted by one or more substituents R';

Y is H, halogen, haloalkyl, alkyl or an alkylester, which can be substituted by one or more substituents R';

R' independently represents H, $-CO_2R''$, -CONHR'', -CR''O, $-SO_2N(R'')_2$, $-SO_2NHR''$, -NR''-CO-haloalkyl, $-NO_2$, -NR''-SO_2-haloalkyl, -NR''-SO_2-alkyl, $-SO_2$-alkyl, -NR''-CO-alkyl, -CN, alkyl, cycloalkyl, aminoalkyl, alkylamino, alkoxy, -OH, -SH, alkylthio, hydroxyalkyl, hydroxyalkylamino, halogen, haloalkyl, haloalkoxy, amino, heterocyclyl, aryl, haloaryl, haloarylalkyl, arylalkyl or heteroaryl;

R'' independently represents H, haloalkyl, hydroxyalkyl, amino, alkoxy, -N=C(R')$_2$, -NR'-CO-R', -CR'O, -CO_2R', alkyl, cycloalkyl, aryl, haloaryl, haloarylalkyl, heteroaryl, heterocyclyl, arylalkyl or aminoalkyl, which are optionally substituted by one or more substituents R';

2. A compound according to claim 1, wherein

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

R^1, Z, Y, R' and R'' are as defined in claim 1.

3. A compound according to claim 1, wherein

R^1 is aryl or heteroaryl, which can be substituted by one or more substituents R'

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

Z, Y, R' and R'' are as defined in claim 1.

4. A compound according to claim 1, wherein

R^1 is aryl or heteroaryl, which can be substituted by one or more substituents R'

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

Z is aryl, cycloalkyl, heterocyclyl or heteroaryl, which can be substituted by one or more substituents R';

Y is H, halogen, haloalkyl, or alkyl, which can be substituted by one or more substituents R';

R' independently represents H, -CO_2R'', -CONHR'', -CR''O, -SO_2N(R'')$_2$, -SO_2NHR'', -CN, alkyl, cycloalkyl, aminoalkyl, alkoxy, -OH, , hydroxyalkyl,

halogen, haloalkyl, haloalkoxy, amino, heterocyclyl, aryl, haloaryl, haloarylalkyl, arylalkyl or heteroaryl;

R'' independently represents H, haloalkyl, hydroxyalkyl, amino, alkoxy, $-N=C(R')_2$, -NR'-CO-R', -CR'O, $-CO_2R'$, alkyl, cycloalkyl, aryl, haloaryl, haloarylalkyl, heteroaryl, heterocyclyl, arylalkyl or aminoalkyl, which are optionally substituted by one or more substituents R'

5. A compound according to claim 1, wherein

R^1 is aryl or heteroaryl, which can be substituted by one or more substituents R';

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

Z is aryl, cycloalkyl, heterocyclyl or heteroaryl, which can be substituted by one or more substituents R';

Y is H, halogen, haloalkyl, or alkyl, which can be substituted by one or more substituents R';

R' independently represents H, $-CO_2R''$, $-SO_2N(R'')_2$, $-SO_2NHR''$, -CN, alkyl, alkoxy, -OH, hydroxyalkyl, halogen, haloalkyl, haloalkoxy, heterocyclyl, aryl, haloaryl, haloarylalkyl, arylalkyl or heteroaryl;

R'' independently represents H, haloalkyl, hydroxyalkyl, amino, alkoxy, $-N=C(R')_2$, -NR'-CO-R', -CR'O, $-CO_2R'$, alkyl, cycloalkyl, aryl, haloaryl, haloarylalkyl, heteroaryl, heterocyclyl, arylalkyl or aminoalkyl, which are optionally substituted by one or more substituents R'

6. A compound according to claim 1, wherein

R^1 is aryl or heteroaryl, which can be substituted by one or more substituents R';

Ar is aryl or heteroaryl, which can be substituted by one or more substituents R';

Z is aryl, cycloalkyl, heterocyclyl or heteroaryl, which can be substituted by one or more substituents R';

Y is H, halogen, haloalkyl, or alkyl, which can be substituted by one or more substituents R';

R' independently represents H, $-CO_2R''$, -CONHR'', -CR''O, $-SO_2N(R'')_2$, $-SO_2NHR''$, -CN, alkyl, alkoxy, -OH, halogen, haloalkyl or haloalkoxy,;

R'' independently represents H, haloalkyl, hydroxyalkyl, amino, alkoxy, $-N=C(R')_2$, -NR'-CO-R', -CR'O, $-CO_2R'$, alkyl, cycloalkyl, aryl, haloaryl, haloarylalkyl, heteroaryl, heterocyclyl, arylalkyl or aminoalkyl, which are optionally substituted by one or more substituents R'

7. A compound according to any of claims 1 to 6, wherein

R^1 is selected from the group comprising phenyl which is optionally substituted by one or more substituents individually selected from trifluoromethyl, fluorine, chlorine, bromine, nitro, NH_2, -CN, $-NHCO-C_{1-4}$-alkyl, methoxy, C_{1-4}-alkyl, $-SO_2NH_2$, or $-SO_2NH-C_{1-4}$-alkyl; pyridyl which is optionally substituted by one or more of the aforementioned substituents for phenyl; pyrimidyl which is optionally substituted by one or more of the aforementioned substituents for phenyl; thienyl which is optionally substituted by one substituent–$COO-C_{1-4}$alkyl; 1,1-dioxo-tetrahydrothienyl, 2,2,2-trifluoroethyl, isopropyl, isobutyl, 2-methoxyethyl, tetrahydropyran-4-ylmethyl, 2-morpholinoethyl, pyridin-2-ylmethyl and tetrahydrofuran-2-ylmethyl;

Ar is selected from the group comprising phenyl and pyridyl, which can be substituted by one or more substituents independently selected from fluorine methoxy or chlorine;

Z is selected from the group comprising phenyl or heteroaryl, which can be substituted by one or more substituents selected from the group comprising C_{1-4}alkyl, C_{1-4}haloalkyl, hydrogen, hydroxyl, C_{1-4}alkoxy, and C_{1-4}alkoxycarbonyl;

Y is selected from the group comprising H, trifluoromethyl and methoxycarbonyl.

8. A compound according to any of claims 1 to 6, wherein

R^1 is selected from the group comprising phenyl which is optionally substituted by one or more substituents individually selected from fluorine, chlorine, bromine, nitro, NH_2, -CN, -NHCO-C_{1-4}-alkyl, methoxy, t-butyl, $-SO_2NH_2$, or $-SO_2NH$-isopropyl; pyridyl; pyrimidyl which is optionally substituted by one or more substituents selected from methyl or trifluoromethyl; thienyl which is optionally substituted by one substituent–COO-methyl, 1,1-dioxo-tetrahydrothienyl, 2,2,2-trifluoroethyl, isopropyl, isobutyl, 2-methoxyethyl, tetrahydropyran-4-ylmethyl, 2-morpholinoethyl, pyridin-2-ylmethyl and tetrahydrofuran-2-ylmethyl;

Ar is selected from the group comprising phenyl, 2,6-difluorophenyl, 2-chloro-6-fluorophenyl, 2-chloro-6-methoxyphenyl, 2-fluorophenyl, 2-chlorophenyl, 2,6-dichlorophenyl, 2,4-dichlorophenyl, 3-fluoropyridin-4-yl, 3,5-dichloropyridin-4-yl and 3,5-difluoropyridin-4-yl;

Z is selected from the group comprising phenyl, tetrazolyl, thiazolyl, 1,3,4-oxadiazolyl, oxazolyl, 1,3,4-thiadiazolyl, furanyl, and thiophenyl, which can be substituted by one or more substituents selected from the group comprising methyl, trifluoromethyl, hydrogen, hydroxyl, methoxy, and methoxycarbonyl and ethoxyoxycarbonyl;

Y is selected from the group comprising H, trifluoromethyl and methoxycarbonyl.

9. A compound according to any of claims 1 to 6, wherein

R^1 is selected from the group comprising phenyl, 2-fluorophenyl, 2-methoxyphenyl, 2-chlorophenyl, 2-bromophenyl, 2-nitrophenyl, 2-aminophenyl, 4-fluorophenyl, 4-tertbutylphenyl, 3-fluorophenyl, 3-chlorophenyl, 3-cyanophenyl, 3-acetamido-phenyl, 2-acetamido-phenyl, 3-aminosulfonyl-phenyl, 3-(isopropylamino)sulfonyl-phenyl, 3-nitrophenyl, 3-aminophenyl, 2,4-difluorophenyl, 3,5-difluorophenyl, 3,5-dichlorophenyl, 2,3,5,6-tetrafluorophenyl, 2-pyridyl; 3-pyridyl; 4-pyridyl; 4- trifluoromethyl-pyrimid-2-yl, 2,6-dimethyl-pyrimid-4-yl, 2-methoxycarbonyl-thien-3-yl, 1,1-dioxo-tetrahydrothienyl, 2,2,2-trifluoroethyl, isopropyl, isobutyl, 2-methoxyethyl, tetrahydropyran-4-yl-methyl, 2-(morpholin-4-yl)-ethyl, and tetrahydrofuran-2-yl-methyl;

Ar is selected from the group comprising phenyl, 2-chloro-6-fluorophenyl, 2-fluorophenyl, 2-chlorophenyl, 2,6-dichlorophenyl, 2,4-dichlorophenyl, 3-fluoropyridin-4-yl, and 3,5-difluoropyridin-4-yl;

Z is selected from the group comprising 1H-tetrazol-5-yl, 4-methylthiazol-2-yl, thiazol-2-yl, 5-(trifluoromethyl)-1,3,4-oxadiazol-2-yl, 5-methyl-1,3,4-oxadiazol-2-yl, 1,3,4-oxadiazol-2-yl, oxazol-5-yl, 4-hydroxy-thiazol-2-yl, 4-methoxy-thiazol-2-yl, 5-methyl-1,3,4-thiadiazol-2-yl, 1,3,4-thiadiazol-2-yl, furan-3-yl, furan-2-yl, thiophen-3-yl, phenyl, 4-methoxycarbonyl-thiazol-2-yl, and 4-ethoxycarbonyl-thiazol-2-yl;

Y is selected from the group comprising H, trifluoromethyl and methoxycarbonyl.

10. A compound according to any of claims 1 to 10, wherein the compound is selected from the group comprising the compounds of examples
3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(4-methylthiazol-2-yl)isoxazole,
3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(thiazol-2-yl)isoxazole,

2-(3-(2-chloro-6-fluorophenyl)-5-(1-(3-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)-5-(trifluoromethyl)-1,3,4-oxadiazole,

2-(3-(2-chloro-6-fluorophenyl)-5-(1-(3-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)-1,3,4-oxadiazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(oxazol-5-yl)isoxazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(4-methoxythiazol-2-yl)isoxazole,

2-(3-(2-chloro-6-fluorophenyl)-5-(1-(2-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)-5-methyl-1,3,4-oxadiazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(2-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(5-methyl-1,3,4-thiadiazol-2-yl)isoxazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-phenylisoxazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(thiophen-3-yl)isoxazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(furan-2-yl)isoxazole,

methyl 2-(3-(2-chloro-6-fluorophenyl)-5-(1-(3-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)thiazole-4-carboxylate,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(5-methyl-1,3,4-thiadiazol-2-yl)isoxazole,

2-(3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)-5-methyl-1,3,4-oxadiazole,

2-(3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)-1,3,4-oxadiazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(furan-2-yl)isoxazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(furan-3-yl)isoxazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(3-chlorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(1,3,4-thiadiazol-2-yl)isoxazole,

3-(2-chloro-6-fluorophenyl)-5-(1-(2-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(furan-3-yl)isoxazole,

2-(3-(2-chloro-6-fluorophenyl)-5-(1-(pyridin-3-yl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)-5-methyl-1,3,4-oxadiazole,
3-(2-chloro-6-fluorophenyl)-4-(5-methyl-1,3,4-thiadiazol-2-yl)-5-(1-(pyridin-3-yl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazole,
3-(2-chloro-6-fluorophenyl)-4-(5-methyl-1,3,4-thiadiazol-2-yl)-5-(1-(pyridin-4-yl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazole,
3-(2-chloro-6-fluorophenyl)-4-(furan-3-yl)-5-(1-(pyridin-4-yl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazole,
2-(3-(2-chloro-6-fluorophenyl)-5-(1-(pyridin-2-yl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)-1,3,4-oxadiazole,
2-(3-(2-chloro-6-fluorophenyl)-5-(1-(pyridin-2-yl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazol-4-yl)-5-methyl-1,3,4-oxadiazole,
3-(2-chloro-6-fluorophenyl)-4-(5-methyl-1,3,4-thiadiazol-2-yl)-5-(1-(pyridin-2-yl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)isoxazole and
3-(2-chloro-6-fluorophenyl)-5-(1-(2-fluorophenyl)-5-(trifluoromethyl)-1H-pyrazol-4-yl)-4-(thiazol-2-yl)isoxazole
of the present invention.

11. A compound according to any of claims 1 to 10 for use as a medicament.

12. A compound according to any of claims 1 to 10 for use in the treatment of a disease or indication is selected from the group consisting of psoriasis, psoriatric arthritis, autoimmune thyroiditis, Grave's disease, rheumatoid arthritis, vitiligo, Crohn's disease, ulcerative colitis, inflammatory bowel disease, ankylosing spondylitis, diabetes type I, multiple sclerosis, celiac disease, systemic lupus erythematosus, uveitis, Behcet disease, atopic dermatitis, Lichen planus, Sjögren's syndrome, spinal disc herniation, acne, Graft-versus-Host-Reaction, Host -versus-Graft-Reaction and osteoarthritis

13. Use of the compounds of the formula (I) as defined in any of claims 1 to 10, and of their pharmacologically tolerable salts in the manufacture of a medicament for use in treatment of a disease or a therapeutic indication in which the inhibition of interleukin-17 (IL-17) and/or Interferon-γ (INF-γ) is beneficial.

14. The use of claim 12 wherein the disease or indication is selected from the group consisting of psoriasis, psoriatric arthritis, autoimmune thyroiditis, Grave's disease, rheumatoid arthritis, vitiligo, Crohn's disease, ulcerative colitis, inflammatory bowel disease, ankylosing spondylitis, diabetes type I, multiple sclerosis, celiac disease, systemic lupus erythematosus, uveitis, Behcet disease, atopic dermatitis, Lichen planus, Sjögren's syndrome, spinal disc herniation, acne, Graft-versus-Host-Reaction, Host -versus-Graft-Reaction and osteoarthritis.