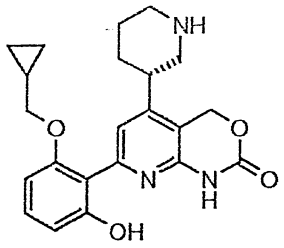
CLAIMS

1. Use of a 2,3-dihydroimidazo[1,2-c]quinazoline compound or a physiologically acceptable salt, solvate, hydrate or stereoisomer thereof as a sole active agent, or of a pharmaceutical composition containing the compound or a physiologically acceptable salt, solvate, hydrate or stereoisomer thereof,
for the preparation of a medicament for the treatment or prophylaxis of non-Hodgkin's lymphoma (NHL), particularly 1st line, 2nd line, relapsed, refractory, indolent or agressive non-Hodgkin's lymphoma (NHL), in particular follicular lymphoma (FL), chronic lymphocytic leukaemia (CLL), marginal zone lymphoma (MZL), diffuse large B-cell lymphoma (DLBCL), mantle cell lymphoma (MCL), transformed lymphoma (TL), or peripheral T-cell lymphoma (PTCL),
wherein said 2,3-dihydroimidazo[1,2-c]quinazoline compound is 2-amino-N-[7-methoxy-8-(3-morpholin-4-ylpropoxy)-2,3-dihydroimidazo[1,2-c]quinazolin-5-yl]pyrimidine-5-carboxamide.

2. Use according to claim 1, wherein said salt is 2-amino-N-[7-methoxy-8-(3-morpholin-4-ylpropoxy)-2,3-dihydroimidazo[1,2-c]quinazolin-5-yl]pyrimidine-5-carboxamide dihydrochloride.

3. A combination drug of :
a) a 2,3-dihydroimidazo[1,2-c]quinazoline compound, or a physiologically acceptable salt, solvate, hydrate or stereoisomer thereof ; or a pharmaceutical composition containing the compound or a physiologically acceptable salt, solvate, hydrate or stereoisomer thereof,
and
b) one or more further active agents selected from the group consisting of : PI3Kδ-selective inhibitor GS-1101, BTK inhibitor ibrutinib, IKK inhibitor BAY Compound B of the formula



, and REFAMETINIB (BAY 86-9766 (RDEA-119)),

for the treatment or prophylaxis of non-Hodgkin's lymphoma (NHL), particularly 1st line, 2nd line, relapsed, refractory, indolent or agressive non-Hodgkin's lymphoma (NHL), in particular follicular lymphoma (FL), chronic lymphocytic leukaemia (CLL), marginal zone lymphoma (MZL), diffuse large B-cell lymphoma (DLBCL), mantle cell lymphoma (MCL), transformed lymphoma (TL), or peripheral T-cell lymphoma (PTCL),
wherein said 2,3-dihydroimidazo[1,2-c]quinazoline compound is 2-amino-N-[7-methoxy-8-(3-morpholin-4-ylpropoxy)-2,3-dihydroimidazo[1,2-c]quinazolin-5-yl]pyrimidine-5-carboxamide.

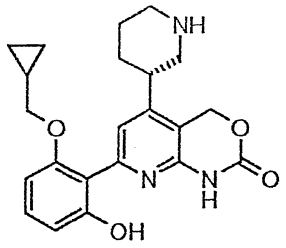
4. The combination drug according to claim 3, comprising 2-amino-N-[7-methoxy-8-(3-morpholin-4-ylpropoxy)-2,3-dihydroimidazo[1,2-c]quinazolin-5-yl]pyrimidine-5-carboxamide.

5. The combination drug according to claim 3, compriding 2-amino-N-[7-methoxy-8-(3-morpholin-4-ylpropoxy)-2,3-dihydroimidazo[1,2-c]quinazolin-5-yl]pyrimidine-5-carboxamide dihydrochloride.

6. The combination drug according to claim 4 or 5, wherein said further active agent is PI3Kδ-selective inhibitor GS-1101.

7. The combination drug according to claim 4 or 5, wherein said further active agent is BTK inhibitor ibrutinib.

8. The combination drug according to claim 4 or 5, wherein said further active agent is IKK inhibitor BAY Compound B of the formula



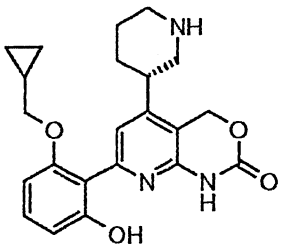
9. The combination drug according to claim 4 or 5, wherein said further active agent is REFAMETINIB (BAY 86-9766 (RDEA-119)).

10. A pharmaceutical composition which comprises a combination of :

a) a 2,3-dihydroimidazo[1,2-c]quinazoline compound, or a physiologically acceptable salt, solvate, hydrate or stereoisomer thereof ;

and

b) one or more further active agents selected from the group consisting of : PI3Kδ-selective inhibitor GS-1101, BTK inhibitor ibrutinib, IKK inhibitor BAY Compound B of the formula



, and REFAMETINIB (BAY 86-9766 (RDEA-119)),

for the treatment or prophylaxis of non-Hodgkin's lymphoma (NHL), particularly 1st line, 2nd line, relapsed, refractory, indolent or agressive non-Hodgkin's lymphoma (NHL), in particular follicular lymphoma (FL), chronic lymphocytic leukaemia (CLL), marginal zone lymphoma (MZL), diffuse large B-cell lymphoma (DLBCL), mantle cell lymphoma (MCL), transformed lymphoma (TL), or peripheral T-cell lymphoma (PTCL),

wherein said 2,3-dihydroimidazo[1,2-c]quinazoline compound is 2-amino-N-[7-methoxy-8-(3-morpholin-4-ylpropoxy)-2,3-dihydroimidazo[1,2-c]quinazolin-5-yl]pyrimidine-5-carboxamide.

11. Use of a combination or a pharmaceutical composition containing the combination for the preparation of a medicament for the treatment or prophylaxis of non-Hodgkin's lymphoma (NHL), particularly 1st line, 2nd line, relapsed, refractory, indolent or agressive non-Hodgkin's lymphoma (NHL), in particular follicular lymphoma (FL), chronic lymphocytic leukaemia (CLL), marginal zone lymphoma (MZL), diffuse large B-cell lymphoma (DLBCL), mantle cell lymphoma (MCL), transformed lymphoma (TL), or peripheral T-cell lymphoma (PTCL),

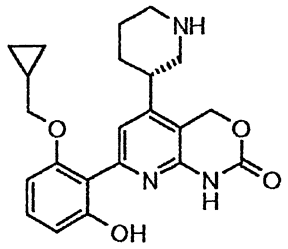
wherein the combination comprises

a) a 2,3-dihydroimidazo[1,2-c]quinazoline compound, or a physiologically acceptable salt, solvate, hydrate or stereoisomer thereof ;

or of a pharmaceutical composition containing such a compound or a physiologically acceptable salt, solvate, hydrate or stereoisomer thereof,

and

b) one or more further active agents selected from the group consisting of : PI3Kδ-selective inhibitor GS-1101, BTK inhibitor ibrutinib, IKK inhibitor BAY Compound B of the formula



, and REFAMETINIB (BAY 86-9766 (RDEA-119)) ;

wherein said 2,3-dihydroimidazo[1,2-c]quinazoline compound is 2-amino-N-[7-methoxy-8-(3-morpholin-4-ylpropoxy)-2,3-dihydroimidazo[1,2-c]quinazolin-5-yl]pyrimidine-5-carboxamide.