



(19)

Europäisches Patentamt  
European Patent Office  
Office européen des brevets



(11)

EP 1 267 901 B9

(12)

## CORRECTED EUROPEAN PATENT SPECIFICATION

Note: Bibliography reflects the latest situation

(15) Correction information:  
**Corrected version no 1 (W1 B1)**  
 Corrections, see page(s) 11

(51) Int Cl.:  
**A61K 36/00 (2006.01)** **A61K 45/06 (2006.01)**  
**A61K 31/352 (2006.01)** **A61P 9/10 (2006.01)**  
**A61K 36/00 (2006.01)** **A61K 31/35 (2006.01)**

(48) Corrigendum issued on:  
**28.06.2006 Bulletin 2006/26**

(86) International application number:  
**PCT/NL2001/000233**

(45) Date of publication and mention  
 of the grant of the patent:  
**25.01.2006 Bulletin 2006/04**

(87) International publication number:  
**WO 2001/070248 (27.09.2001 Gazette 2001/39)**

(21) Application number: **01915929.2**

(22) Date of filing: **22.03.2001**

### (54) COMPOSITIONS SUITABLE FOR THE TREATMENT OF DAMAGE CAUSED BY ISCHEMIA/ REPERFUSION OR OXIDATIVE STRESS

ZUSAMMENSETZUNGEN ZUR BEHANDLUNG VON SCHÄDEN VERURSACHT DURCH ISCHÄMIE / REPERFUSION ODER OXIDATIVEM STRESS

COMPOSITIONS CONCUES POUR LE TRAITEMENT DE LESIONS PROVOQUEES PAR L'ISCHEMIE/REPERFUSION OU LE STRESS OXYDATIF

(84) Designated Contracting States:  
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU**  
**MC NL PT SE TR**

(56) References cited:  
**WO-A-00/07578** **WO-A-00/25795**  
**WO-A1-00/64277** **WO-A1-97/32947**  
**WO-A1-99/49851** **DE-U- 29 908 634**  
**US-A- 5 804 168** **US-A- 5 868 082**  
**US-A- 5 886 029** **US-A- 5 976 568**

(30) Priority: **22.03.2000 NL 0201051**

- **SHOSKES ET AL.:** "Synergy if mycophenolate mofetil and bioflavonoids in prevention of immune and ischemic injury"  
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(43) Date of publication of application:  
**02.01.2003 Bulletin 2003/01**

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- DATABASE BIOSIS [Online] BIOSCIENCES INFORMATION SERVICE, PHILADELPHIA, PA, US; AN 2000:303443, YAGI, NOBUAKI ET AL: "Protective effect of green tea extract against reperfusion injury in rats: Antioxidant and anti-inflammatory properties." retrieved from STN XP002145979 & JOURNAL OF CLINICAL BIOCHEMISTRY AND NUTRITION, (1999) VOL. 27, NO. 2, PP. 89-101. PRINT.,

- Am J Clin Nutr 1998; 1210-1218

Remarks:

The file contains technical information submitted after the application was filed and not included in this specification

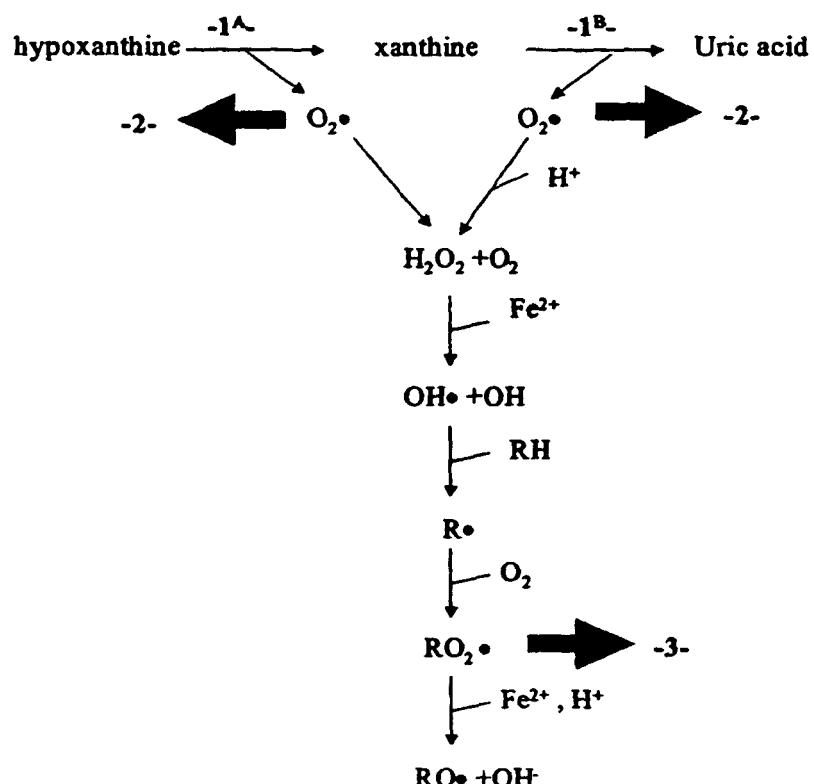
**Description**

[0001] The present invention relates to compositions containing substances which are able to repair or prevent damage caused by ischemia/reperfusion or oxidative stress. More in particular the present invention relates to a composition containing different substances each interfering at a different stage of the radical cascade caused by the above mentioned disorders.

[0002] Ischemia/reperfusion and oxidative stress related diseases and injuries are a widespread problem world wide. These processes can take place all over the body and are instantly occurring, very fast and cascade like. An approach of braking the cascade chain at only one side, carries the danger of highly reactive components slipping through, and because each component of the chain can start a reaction chain on its own, this implicates an inefficient way of approaching the problem.

[0003] The invention is based on the approach via a mixture of components, that together can interfere at three different stages of the radical cascade. In this way providing a more accurate and effective defence against ischemia/reperfusion and oxidative stress mediated injuries and diseases.

[0004] The cascade can be illustrated as follows:



[0005] According to the present invention, there is provided a preparation containing

- at least one substance having xanthine oxidase inhibiting properties;
- at least one substance having superoxide radical scavenging properties;
- at least one substance having peroxyl radical scavenging properties;

[0006] The properties of these substances a), b) and c) can be further defined:

Substance a) preferably has an IC-50 value of less than 10  $\mu M$  as determined by the xanthine oxidase assay. In the xanthine oxidase assay the reduction of xanthine oxidase activity to convert xanthine to uric acid is measured. Substance b) preferably has an IC-50 value of less than 10  $\mu M$  as determined by the superoxide scavenging assay. In the superoxide scavenging assay the reduction of the concentration superoxide radicals resulting from conversion of hypoxanthine by xanthine oxidase is measured. Moreover, the IC-50 value of a substance b) in the superoxide scavenging assay is less than the IC-50 value of said substance in the xanthine oxidase assay in order to discriminate

the anti-oxidant from the enzyme inhibitory activities and to mark the component as a superoxide radical scavenging component.

5 Substance c) has a value greater than 1.0 in relation to the value for trolox on a molar basis, as determined in the  $\beta$ -phycoerythrin fluorescence based assay. In the  $\beta$ -phycoerythrin fluorescence based assay the reduction of  $\beta$ -phycoerythrin decomposition by radicals resulting from the thermal decomposition of V50 is measured. The value obtained for the substance to be tested is compared, on a molar basis, to the value for Trolox.

[0007] The above mentioned tests are further illustrated in the examples.

10 [0008] According to the present invention substances a), b) and c) can be any substance, natural or synthetic, as long as they fulfil the above-described criteria. Examples of substance a) having xanthine oxidase inhibiting properties are quercetin, myricetin, luteolin, chrysin, apigenin and allopurinol. In particular substance a) is selected from apigenin, and luteolin.

[0009] Substance b) having superoxide radical scavenging properties is selected from epicatechin and taxifolin.

15 [0010] Substance c) having peroxy radical scavenging properties is selected from quercetin, epicatechin, luteolin, apigenin, curcumin, chrysin, fustin and myricetin.

[0011] 20 All the above values in the assays are based on pure substances. However, mixtures of substances that comply with any of the three criteria can also be used. Most preferred mixtures are extracts of one or more natural sources which comprise flavonoids. In these extracts flavonoids are often present in glycosilated forms. In the body these forms can be converted into aglycons. As a substance a) extracts of one or more of the following plants appear to be effective: onion, hypericum perforatum, chamomillae, linden flowers, thyme, parsley, peanut (shells), Artocarpus heterophyllus, Morus alba, Cynara scolymus, Apium graveolens, Malus sylvestris, Dimorphandra mollis, Vaccinium macrocarpon, Oenothera biennis, Podophyllum species and Azadirachta indica. As a substance b) for instance green tea extract, Aerola or Hamamelidis extract can be used. As a substance c) for instance green tea extract or grape skin extract can be used.

25 [0012] In the preparation of the present invention the xanthine oxidase inhibiting components should be present in an amount such that at least x mg/day is administered, wherein:

$$\Sigma x / (IC-50 * D) = 49$$

30 wherein the IC-50 is determined according to the xanthine oxidase assay ( $\mu$ g/ml). D = 1000 ml/day. D is a correction factor for the formula, it is not related to the administration of the components. In case of luteolin this means that at least 10 mg/day luteolin is administered, when this is the only xanthine oxidase inhibitor present. For quercetin this is 25 mg/day.

35 [0013] The superoxide radical scavengers should be present in an amount such that at least x mg/day is administered, wherein:

$$\Sigma x / (IC-50 * D) = 14$$

40 wherein the IC-50 is determined according to the superoxide assay ( $\mu$ g/ml). D = 1000 ml/day. D is a correction factor for the formula, it is not related to the administration of the components. In case of epicatechin this means that at least 9.8 mg/day epicatechin is administered, when this is the only superoxide scavenger present.

45 [0014] The peroxide scavenger components should be present in an amount such that at least x mg/day is administered, wherein:

$$x * \text{trolox equivalent} = 18.$$

50 [0015] The trolox equivalent is determined according to the  $\beta$ -phycoerythrin fluorescence based assay (mg). In case of apigenin this means that at least 10.9 mg/day apigenin is administered, when this is the only lipid peroxide scavenger present.

55 [0016] The maximum amount of the substance to be administered is not limited. In general at a certain level no additional effect will be observed because the transport of the substance into the blood of the patient will be limited. For economical reasons the maximum amount of the substance will be the amount at which no additional effect occurs. This can be determined by determining plasma concentrations and testing these concentrations in in vitro assays.

[0017] The present invention also relates to the use of the preparation according to the invention for the treatment of damage caused by ischemia/reperfusion. An ischemia/reperfusion mediated condition is herein defined as a condition

caused by a reduced blood flow, or a reduced maximal blood flow to a certain organ, tissue or part of an organ or tissue; followed by a period of re-established blood flow. It is also possible that the concentration of oxygen is not matching the demands for certain time periods. This is for example the case in patients suffering from breathing problems as for example COPD, ARDS, and cystic fibrosis.

5 [0018] The preparation can also be used for the treatment of oxidative stress. Because xanthine oxidase can start the described cascade, events in which oxidative stress is associated with high levels of xanthine oxidase can be treated or prevented with the described preparation. In this case the prevention or treatment is more effective than with xanthine oxidase inhibitors alone because the invention warrants the stop of the cascade at three levels.

10 [0019] Examples of damage caused by ischemia/reperfusion and oxidative stress are multiple organ failure, subfertility, disorders that occur in relation to surgical procedures, damage to organs after surgical procedures, damage to lungs after damage to tissue or organs in other parts of the body, for instance lung damage after burn injuries or after pancreatitis, organ transplants, heart failure or angina, COPD or ARDS, cystic fibrosis, Inflammatory Bowel Disease (1, 7, 15), circulatory shock, gout, rheumatoid arthritis (10, 12), osteoarthritis (6), protection against liver damage caused by use of medication or ethanol (3), inflammation and disorders related therewith, hypertension (13), muscle damage after 15 sports (4), acute renal failure (2), pre-enclampsia or enclampsia (8, 9), sepsis (14).

20 [0020] The application of the present preparation in surgery is important. For example, during surgery arteries can be clamped to avoid excessive blood loss. If this clamp is removed, tissues and organs are reperfused. Moreover, during surgery blood is retained from the intestinal organs. This means that surgery itself induces a mild ischemia/reperfusion cycle for the intestine. Moving and twisting of tissue and organs e.g. large intestine (for the duration of a surgery) can also lead to short and local ischemia and reperfusion events. Another example in which surgery leads to an ischemia/reperfusion event is when blood flow to an ischemic part of the body is restored by surgery, e.g. a bypass of a heart, or by pursing of atherosclerotic regions in the leg. Elevations of xanthine oxidase in combination with oxidative stress can also occur in injured human skeletal muscle (4) or hypertension (13).

25 [0021] Multiple organ failure sometimes occurs when ischemia of one organ leads to ischemia of another organ. The liver receives most of the blood from the intestine. This means that ischemia/reperfusion of the intestine leads to ischemia/reperfusion of the liver. In this way not only damage to the intestine occurs, but also damage to the liver occurs. An injured liver releases all kinds of stored enzymes. One of these enzymes is xanthine oxidase (XO). The blood as drained from the liver by the vena cava reaches the lung. The lung is oxygen rich and therefore oxygen radicals can easily be formed. The xanthine oxidase produced by the liver can induce damage in the lung. Similar cascades can happen toward 30 kidney and from lung to liver. Hereafter preliminary data are given in which components mixed according to the invention are given to rats in order to prevent this multiple organ failure cascade.

35 [0022] Another specific example where ischemia/reperfusion causes damage is organ transplantation. Organs are removed from the donor (ischemic period) and transplanted into the recipient (reperfusion period). This means that the invention can be used for preparations given to the recipient to decrease ischemia/reperfusion effects in the body, as well as for fluids to preserve the organs before being transplanted into the recipient.

[0023] The preparation can also be used in case of subfertility. Oxidative stress in combination with high xanthine oxidase levels can be observed in subfertility in men (5, 11). The present invention will reduce XO activity and enhance radical scavenging and therefore improve fertility.

[0024] Another example of restoration of blood flow takes place after a circulatory shock or after a severe blood loss.

40 [0025] Ischemia/reperfusion also takes place in rheumatoid arthritis and osteoarthritis. During the swelling of the joints reduction in blood flow to the joint due to the pressure of the swelling can occur. This leads to ischemia. Reduction of the swelling leads to perfusion (6, 10).

[0026] The invention is illustrated by the examples, which also show the assays used to determine the properties of the different substances used.

45

## Examples

### Determination of properties

50 Xanthine oxidase inhibition (xanthine oxidase assay)

[0027] The activity of xanthine oxidase is determined by means of the uric acid production. Materials used:

55 phosphate buffer: 50 mM Na<sub>2</sub>HPO<sub>4</sub>; MW = 141.96 g/mol; 7.10 g/l, pH adjusted to 7.8 with 1 M HCl

xanthine solution: 21.2 mg/250 ml phosphate buffer

(for dissolving the xanthine this mixture is agitated ultrasonically for 1,5 hours and heated to 90 °C for 5 to 10 minutes)

enzyme solution: XOD (xanthine oxidase) (grade I from buttermilk) 18.5 U/ml, 10.2 µl XOD 18.5 U/ml in 5000 µl in ice-cold phosphate buffer (189\*10<sup>-3</sup> U/5000 µl). This solution is maintained on ice during the entire experiment to

prevent changes in the activity of the enzyme.

Method:

5 [0028] Assay mixture:

724 µl phosphate buffer pH=7.8  
 200 µl xanthine solution  
 10 µl DMSO or the substance to be tested dissolved in DMSO

10 [0029] The reaction is started by addition of 66 µl of the enzyme solution to the above described assay mixture. The reaction is monitored after half a minute from the start for 6 minutes, by measuring the absorbance at 295 nm. If the reaction is linear, the result is expressed as Δabsorbance/min. The absorbance at 295 nm is a measure for the uric acid concentration.

15 [0030] The IC-50 value is determined as the concentration of the substance to be tested corresponding to 50 % inhibition as compared to the maximum activity. The maximum activity is the activity of the enzyme, when no substance is dissolved in DMSO.

[0031] Examples of substances and extracts having xanthine oxidase inhibiting properties:

Sample	IC-50 (µg/ml)	IC-50 (µM)
Luteolin	0.215	0.75
Apigenin	0.27	1.0
Myricetin	0.48	1.5
Quercetin dihydrate	0.51	1.5
Chrysin	0.645	2.5
Allopurinol	0.84	6.2
Catechin	not detectable	not detectable
Epicatechin	not detectable	not detectable
Curcumin	not detectable	not detectable
Epigalo-catechin-galate (ECCG)	not detectable	not detectable

[0032] These materials were obtained from Indofine chemicals and Sigma.

40 Superoxide radical scavenging (superoxide scavenging assay)

[0033] Literature reference: Re-evaluation of assay methods and establishment of kit for superoxide dismutase activity, Yoshihiko Oyanagui, Analytical Biochemistry 142, 290-96 (1984).

Materials used:

45 All solutions are prepared with analytical grade chemicals and milliQ water.

Buffer A: 0.2 mM hydroxylamine and 1.87 mM hypoxanthine dissolved in buffer B. For dissolving the components it can be necessary to ultrasonically agitate the mixture of the substances followed by heating the mixture to 80 °C for several minutes.

50 Buffer B: 20.8 mM K<sub>2</sub>HPO<sub>4</sub> and 15.6 mM Na<sub>2</sub>B<sub>4</sub>O<sub>7</sub>, the pH being adjusted to 7.0 with 1 M NaOH

Buffer C (colouring reagent): 300 µg/ml sulfanilic acid and 10 µg/ml N-(1-naphtylethylenediamine) dissolved in 16.7 % acetic acid.

Enzyme solution: xanthine oxidase from Buttermilk Grade I 5.25 µl of 18.5 U/ml dissolved in 7575 µl buffer B (97\*10<sup>-3</sup> U in 7575 µl).

55 Method:

[0034] 10 µl of the substances to be tested were dissolved in DMSO or milli Q water and added to and mixed with,

respectively: 390 µl buffer B and 200 µl buffer A. Next 400 µl of the enzyme solution is added and mixed and the sample is placed directly at 37 °C and incubated at this temperature for 30 minutes. Then the reaction is stopped by adding 2 ml of buffer C. The sample is then incubated at 20 °C for 20 minutes and the absorbance at 550 nm is measured. The absorbance is a result of reaction of a coloring reagent with remaining superoxide radicals and therefore a measure of the amount of superoxide radicals remaining.

5 The results were compared with a reference containing only the solvent, i.e. either DMSO or milliQ water.

The IC-50 value is the concentration of a substance or mixture of which the scavenging activity is 50% of the maximum activity. The maximum activity is the absorbance at 550 nm of the above mentioned reference.

10 [0035] The absorbance measured is corrected for background colouring. The correction factor is the absorbance at 550 nm of a sample, incubated as described above, but containing only 10 µl of DMSO and to which instead of 400 µl enzyme solution, 400 µl buffer B is added.

15 Examples of substances having superoxide scavenging properties

Sample	IC-50 (µg/ml)	IC-50 (µM)
Epicatechin	0.7	2.4
Myricetin	1.6	3.4 <sup>1)</sup>
Taxifolin	2.8	9.2
QDH	3.6	10.8 <sup>1)</sup>
Luteolin		>100
Apigenin		>100
Chrysin		>100
Catechin		>100
curcumin		>100
ECCG		>100
Vitamin C	4.4	25

20 25 30

<sup>1)</sup> These substances are excluded because the IC-50 or superoxide radical scavenging >> IC-50 xanthine oxidase inhibition.

35 Peroxyl radical scavenging ( $\beta$ -phycoerythrin fluorescence assay)

40 [0036] Literature reference:

1. Oxygen Radical Absorbance capacity assay for antioxidants, Guohua Cao et al, Free Radical Biology and medicine, vol. 14, pp. 303-311, 1993.

2. Phycoerythrin fluorescence based assay for peroxy radicals: A screen for biologically relevant protective agents; Delange, R.J., Glazer A.N., Anal. Biochem., 177; 300-306 (1989).

45 [0037] In order to avoid differences in result by small changes in measuring conditions, such as analytical instruments, sensitivity, etc. the end results are expressed in relation to the peroxy radical scavenging capacity of a known amount of Trolox (a vitamin E analogue).

50 [0038] The principle underlying the present method is that  $\beta$ -phycoerythrin is decomposed by radicals that are the result of the thermolitic decomposition of V50 at 20 °C. When a product scavenges these radicals, the decrease in fluorescence in time will occur less quickly. All measurements should be carried out at least in threefold.

Materials used:

55 [0039] Equipment: Fluorimeter for 96 wells. The temperature of the tray must be kept constant. The variation between the highest and the lowest temperature of the wells should not be more than 0.5 °C. 96 well white trays of Perkin and Elmer were used.

Chemicals:

**[0040]**

5 AAPH (2,2'-azobis(2-aminopropane)dihydrochloride) (e.g. from Wako Chemicals)= V50; MW = 271.19 g/mol; 16 mM in the assay  
 β-PE (e.g. from Sigma) MW=240,000 g /mol; 16.7 nM in the assay  
 Trolox (e.g. from Aldrich) MW = 250.29 g/mol; in the assay test in the range of 1-10 μM.

10 Conditions in the assay ORAC<sub>ROO</sub>:

**[0041]**

15 10 μl sample in ethanol

50 μl 50 mM PBS pH=7.4

50 μl 66.8 nM β-PE

20 **[0042]** Each tray should contain trolox as a reference and as a standard (2 μM or 0.5 mg/l). As a negative control a sample is taken containing only 10 μl ethanol, 50 μl 50 mM PBS, pH=7.4 and 50 μl 66.8 nM β-PE.

**[0043]** The reaction is started by adding 100 μl 32nM V-50. The time between pipetting and mixing of this last step in a well and the measurement of this well, should be constant. Further, the time between the last pipetting step in a well and the first measurement of the well should be less than 2 minutes.

25 V-50 should be stored in the dark at 4 °C until used. The reaction kinetics of the wells are monitored in a continuous monitoring process. The excitation wavelength is 540 nm, while the emission wavelength is 565 nm. The reaction is monitored until the signal of the sample differs no longer from the background signal (+/- 10%). The area underneath the curve of the negative reference is subtracted from the area underneath the substance to be measured. This difference in area is then related to a difference in area in relation to a negative reference resulting from trolox.

30 Examples

**[0044]**

35	Sample	TE (Trolox equivalent) (based on mass)	TE (Trolox equivalent) (based on mole weight)
	apigenin	1.56	1.69
	quercetin	1.32	1.78
40	curcumin	1.27	1.72
	epicatechin	1.24	1.44
	chrysin	1.23	1.25
45	luteolin	1.22	1.40
	fustin	1.07	1.24
	myricetin	0.89	1.13
	EGCG	0.54	0.99

50 **[0045]** A mixture having xanthine oxidase inhibiting, superoxide radical scavenging and peroxy radical scavenging properties according to the present invention was used in an in vitro experiment to show that this mixtures indeed had these properties. Next a rat model was developed in which a cascade of multiple organ failure occurs. In this model rats were fed a normal feed or a feed of a flavonoid mixture according to the present invention.

55 **[0046]** An experiment was carried out in male Wistar rats. Rats were fed for 5 days a flavonoid enriched diet containing luteolin, quercetin, apigenin, epicatechin and tea extract, or a control diet. Rats were not fasted, but could eat ad libitum until the anaesthesia. Rats were anaesthetised and a clamp on the arteria mesenterica was placed. The experiment was carried out in two groups of nine rats. Ischemia was performed for one hour, followed by two hours of reperfusion.

Biochemical parameters were measured in order to check if xanthine oxidase had been inhibited. Parameters indicating oxidative stress and other parameters indicating damage to an organ were measured.

5 1. Biochemical

[0047] Determination of uric acid and allatonin levels.

	Comparative	SE	Flavonoid mix	SE	p
Uric acid ( $\mu\text{mol/L}$ )	386	193.5	261	90.9	0.03

[0048] Xanthine oxidase was significantly inhibited by the flavonoid mix. As is shown in the above table uric acid levels decrease significantly, when flavonoids are added.

15 2. Small intestine

[0049]

	Comparative	SE	Flavonoid mix	SE	p
Apoptose	2.58	1.91	0.20	0.11	0.02

[0050] A significant reduction was found in apoptosis (spontaneous cell death) in animals fed with the flavonoid mixture.

25 3. Liver parameters

[0051]

	Comparative	SE	Flavonoid mix	SE	p
Plasma:					
ALAT (U/L)	47	4.1	45.8	7.3	0.45
Arginase ( $\mu\text{mol/min}$ )	2.35	0.55	1.83	0.71	0.28
Bilburine ( $\mu\text{mol/L}$ )	1.7	0.6	1	0	0.18
ASAT (U/L)	94.3	13.3	78	9.5	0.15
GGT (U/L)	0.4	0.2	0.1	0.1	0.12
Urea (mmol/L)	8.8	3.4	7.2	0.6	0.11
Tissue homogenates					
8-iso-PGF2 alpha (pg/ml)	304.7	36	242.2	30	0.12
MDA (nM)	263.6	12.1	236.7	13.1	0.075

[0052] All parameters described above are parameters indicating liver injury. From the table above tendency towards liver injury can be detected.

50 4. Kidney function

[0053]

	Comparative	SE	Flavonoid mix	SE	p
Creatinine ( $\mu\text{mol/l}$ )	57.4	7.1	49.4	2.9	0.16

[0054] Also kidney function and in general lactate production seem to be decreased when flavonoids are administered.

## 5. General

[0055]

	Comparative	SE	Flavonoid mix	SE	p
Lactate (mmol/L)	1.9	0.2	1.6	0.1	0.2

10 [0056] In general: intestine function is improved significantly and also other organs seem to profit from the invention.

## Formulation example 1

15 [0057] Tablet prepared according to methods known in the art containing as active ingredients

5 mg apigenin

18 mg quercetin dihydrate

10 mg (-)epicatechin

## Formulation example 2

20 [0058] Tablet prepared according to methods known in the art containing as active ingredients

Parsley extract comprising 10 mg apigenin aglycon equivalents

Apple extract comprising 20 mg quercetin aglycon equivalents

Green tea extract comprising 15 mg epicatechin aglycon equivalents

Curcuma extract comprising 20 mg curcumin aglycon equivalents

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**Claims****1. Use of a preparation containing**

5 a) at least one substance having xanthine oxidase inhibiting properties, comprising luteolin [**deletion(s)**] or apigenin  
 b) at least one substance having superoxide radical scavenging properties, comprising epicatechin or taxifolin; and  
 10 c) at least one substance having peroxy radical scavenging properties, comprising a substance selected from apigenin, quercetin, curcumin, epicatechin, chrysins, luteolin, fustin and myricetin;

15 for the preparation of a medicament for the treatment of damage caused by ischemia/reperfusion and/or oxidative stress; wherein the substances having xanthine oxidase inhibiting properties are to be administered in an amount of at least x mg/day, wherein  $\Sigma x / (IC50 \cdot D) = 49$ , wherein the IC50 is determined according to the xanthine oxidase assay ( $\mu\text{g/ml}$ ) and D = 1000 ml/day.

**2. Use according to claim 1, wherein the medicament is for the prevention of multiple organ failure.****3. Use according to claim 1, wherein the medicament is for the treatment or prevention of peri-operative complications.****Patentansprüche****1. Verwendung einer Zusammensetzung, umfassend**

25 a) mindestens eine Substanz mit xanthinoxidaseinhibierenden Eigenschaften, umfassend Luteolin oder Apigenin;  
 b) mindestens eine Substanz mit Superoxid-Radikal-Fängereigenschaften, umfassend Epicatechin oder Taxifolin; und  
 30 c) mindestens eine Substanz mit Peroxyl-Radikal-Fängereigenschaften, umfassend eine Substanz, ausgewählt aus der Gruppe von Apigenin, Quercetin, Curcumin, Epicatechin, Chrysins, Luteolin, Fustin und Myricetin;

35 zur Herstellung eines Arzneimittels zur Behandlung von durch Ischämie/ Reperfusion verursachten Schäden und/ oder oxidativen Stress; wobei die Substanzen mit xanthinoxidaseinhibierenden Eigenschaften in einer Menge von mindestens x mg/Tag verabreicht werden sollen, wobei  $\Sigma x / (IC50 \cdot D) = 49$  ist, wobei der IC50-Wert gemäß dem Xanthinoxidase-Assay ( $\mu\text{g/ml}$ ) bestimmt wird und D = 1.000 ml/Tag beträgt.

**2. Verwendung nach Anspruch 1, wobei das Arzneimittel zur Prävention von multiplem Organversagen eingesetzt wird.****40 3. Verwendung nach Anspruch 1, wobei das Arzneimittel zur Behandlung oder Prävention von perioperativen Komplikationen eingesetzt wird.****Revendications****45 1. Utilisation d'une préparation contenant**

50 a) au moins une substance possédant des propriétés d'inhibition de la xanthine oxydase, comprenant de la lutéoline ou de l'apigénine ;  
 b) au moins une substance possédant des propriétés de piégeage de radicaux superoxydes, comprenant de l'épicatéchine ou de la taxifoline ; et  
 55 c) au moins une substance possédant des propriétés de piégeage de radicaux peroxyde, comprenant une substance choisie parmi l'apigénine, la quercétine, la curcumine, l'épicatéchine, la chrysine, la lutéoline, la fustine et la myricétine ;

55 pour la préparation d'un médicament destiné au traitement de lésions provoquées par l'ischémie / reperfusion et/ou un stress oxydatif ; dans lequel les substances possédant des propriétés d'inhibition de la xanthine oxydase sont destinées à être administrées selon une quantité d'au moins x mg/jour, où  $\Sigma x / (CI50 * D) = 49$ , où la CI50 est

déterminée conformément au test de la xanthine oxydase ( $\mu\text{g/ml}$ ) et  $D = 1\ 000\ \text{ml/jour}$ .

2. Utilisation selon la revendication 1, dans laquelle le médicament est destiné à la prévention d'insuffisances d'organes multiples.

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3. Utilisation selon la revendication 1, dans laquelle le médicament est destiné au traitement ou à la prévention de complications péri-opératoires.

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