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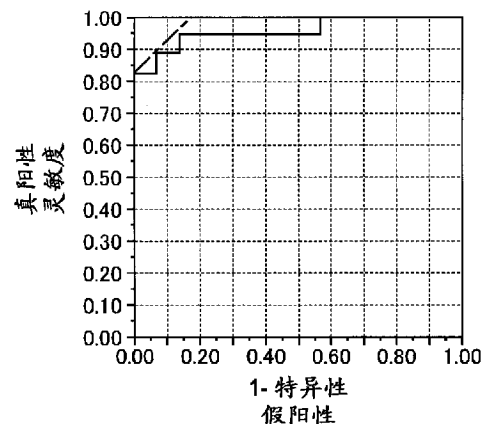
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(54) 发明名称  
 类风湿性关节炎的检查方法及类风湿性关节炎检查用试剂盒

(57) 摘要  
 本发明涉及新型的类风湿性关节炎的检查方法、及用于该检查方法的类风湿性关节炎检查用试剂盒。本发明涉及的类风湿性关节炎的检查方法的特征在于,包括测定受试动物的血浆中或血清中的踝蛋白量的步骤。该测定例如可利用使用了与踝蛋白结合的抗体的免疫学手段进行。本发明涉及的类风湿性关节炎检查用试剂盒用于在上述检查方法中使用,例如,包含固定有与踝蛋白结合的抗体的固相载体。



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1. 与踝蛋白结合的抗体在制造类风湿性关节炎检查用试剂盒中的用途,所述试剂盒用于对类风湿性关节炎的检查,所述检查中包括测定受试动物的血浆中或血清中的踝蛋白量的步骤,其中,所述受试动物是人。

2. 如权利要求 1 所述的用途,其中,进一步包括从由所述受试动物采集的血液中得到血浆或血清的步骤。

3. 如权利要求 1 所述的用途,是为了诊断类风湿性关节炎或判断类风湿性关节炎治疗药的治疗效果而进行的。

4. 如权利要求 1 ~ 3 中任一项所述的用途,其中所述类风湿性关节炎检查用试剂盒包含固定有与踝蛋白结合的抗体的固相载体。

## 类风湿性关节炎的检查方法及类风湿性关节炎检查用试剂盒

### 技术领域

[0001] 本发明涉及类风湿性关节炎的检查方法、及用于该检查方法的类风湿性关节炎检查用试剂盒。

### 背景技术

[0002] 类风湿性关节炎 (Rheumatoid Arthritis:RA) 是以关节的滑膜组织为病变主要部位的慢性炎症性疾病,患病率占人口的约1%。类风湿性关节炎,在其初期导致滑膜炎,随后侵袭软骨/骨,恶化时关节被破坏发生变形。另外,症状的经过而言,有关节炎反复缓解/复发、痊愈例或急速恶化例等多种情况。

[0003] 虽然类风湿性关节炎的诊断主要通过症状来进行,但近年来,以患者血清中所含的自身抗体为标志物的诊断方法受到关注。作为这样的自身抗体,已知有类风湿因子 (rheumatoid factor) (针对变性 IgG 的自身抗体)、抗环状瓜氨酸化肽抗体 (抗 CCP 抗体) 等 (参见非专利文献 1)。

[0004] 然而,在迄今为止的报道中,类风湿因子的灵敏度为 75 ~ 80%、特异性为 50 ~ 70%,抗 CCP 抗体的灵敏度为 50 ~ 75%、特异性为 85 ~ 95%,不一定能满足需要 (参见非专利文献 2、3)。

[0005] 非专利文献 1 :Martinus A. M. 等, Arthritis Res. Ther. ,4 :87-93,2002

[0006] 非专利文献 2 :Avouac J. 等, Ann. Rheum. Dis. 65 :845-851,2006

[0007] 非专利文献 3 :van Venrooij WJ. 等 Ann. N. Y. Acad. Sci. 1143 :268-285,2008

### 发明内容

[0008] 因此,本发明的目的在于,基于探索迄今未知的新型标志物从而发现的新型标志物,提供类风湿性关节炎的检查方法及类风湿性关节炎检查用试剂盒。

[0009] 已知在类风湿性关节炎患者中,血中的淋巴细胞被活化、与血管内皮细胞的细胞附着亢进,同时淋巴细胞游走也发生亢进,结果导致淋巴细胞浸润至血管外,该浸润淋巴细胞引起多种炎症。因此,本发明人在探索新型标志物时,着眼于主要集中在细胞与基质附着区域、尤其是淋巴细胞内细胞附着区域表达的作为高分子细胞骨架蛋白质的踝蛋白 (Talin)。

[0010] 踝蛋白是由包含 FERM 区域的分子量 47kDa 的 N 末端区域、和由一束  $\alpha$  螺旋形成的分子量 190kDa 的 C 末端区域构成的蛋白质。FERM 区域从 N 末端侧进一步分为 F1 结构域 (domain)、F2 结构域、F3 结构域三个亚区域。已知:在生物体内被钙蛋白酶 (Calpain) 剪切下的 N 末端区域的多肽其中 F3 结构域与整联蛋白  $\beta$  亚基结合,增强整联蛋白的从细胞内向细胞外的信号传递,使细胞附着或细胞游走亢进。

[0011] 本发明人对类风湿性关节炎患者的血浆或血清中踝蛋白的存在进行了研究。结果意外地发现,类风湿性关节炎患者中,踝蛋白在血浆或血清中以优势地位存在。而且发现,

当通过类风湿性关节炎治疗药实现类风湿性关节炎的低疾病活动性或缓解时, 踝蛋白量显著降低。

[0012] 本发明是基于上述见解完成的, 具体如下所述。

[0013] (1) 一种类风湿性关节炎的检查方法, 其中, 包括测定受试动物的血浆中或血清中的踝蛋白量的步骤。

[0014] (2) 上述 (1) 所述的类风湿性关节炎的检查方法, 其中, 进一步包括从由上述受试动物采集的血液中得到血浆或血清的步骤。

[0015] (3) 上述 (1) 或 (2) 所述的类风湿性关节炎的检查方法, 其中, 使用与踝蛋白结合的抗体, 测定上述血浆中或血清中的踝蛋白量。

[0016] (4) 上述 (1) ~ (3) 中任一项所述的类风湿性关节炎的检查方法, 其中, 上述受试动物是人。

[0017] (5) 上述 (1) ~ (4) 中任一项所述的类风湿性关节炎的检查方法, 是为了诊断类风湿性关节炎或判断类风湿性关节炎治疗药的治疗效果而进行的。

[0018] (6) 一种类风湿性关节炎检查用试剂盒, 用于在上述 (1) ~ (5) 中任一项所述的类风湿性关节炎的检查方法中使用。

[0019] (7) 上述 (6) 所述的类风湿性关节炎检查用试剂盒, 包含固定有与踝蛋白结合的抗体的固相载体。

[0020] 通过本发明, 可提供新型的类风湿性关节炎的检查方法、及该检查方法中使用的类风湿性关节炎检查用试剂盒。

#### 附图说明

[0021] [图 1] 为表示利用使用了 H-18 抗体及 H-300 抗体的夹心 ELISA 法进行的类风湿性关节炎诊断 (实施例 1) 的 ROC 曲线的图。

[0022] [图 2] 为表示利用使用了 H-18 抗体及 M54246M 抗体的夹心 ELISA 法进行的类风湿性关节炎诊断 (实施例 2) 的 ROC 曲线的图。

[0023] [图 3] 为表示使用了抗 CCP 抗体的类风湿性关节炎诊断 (比较例 1) 的 ROC 曲线的图。

#### 具体实施方式

[0024] < 类风湿性关节炎的检查方法 >

[0025] 本发明涉及的类风湿性关节炎的检查方法包括测定受试动物的血浆中或血清中的踝蛋白量的步骤。该检查方法还可进一步包括从由受试动物采集的血液中得到血浆或血清的步骤。

[0026] 作为受试动物, 只要是可患有类风湿性关节炎的动物就没有特别限制, 可根据目的选择。例如, 可举出人、大鼠、小鼠、狗、牛、猫、兔、豚鼠等, 其中优选人。

[0027] 另外, 对于取得血浆及血清的方法没有特别限制, 可按照现有公知的方法、例如作为临床检查用样品从血液取得的血浆、血清的分离方法进行。例如, 可通过用 EDTA 管或肝素管等采集血液、并对其进行离心分离来得到血浆。另外, 可通过将血液采集至试管等并进行离心分离来得到血清。

[0028] 本发明涉及的类风湿性关节炎的检查方法中,测定如上所述得到的血浆中或血清中的踝蛋白量。此处,“踝蛋白量”是指踝蛋白的蛋白质的量。当踝蛋白具有多种同工型(isoform)时,可测定其中任一种。例如,在人的情况下,存在踝蛋白1、踝蛋白2两种同工型。踝蛋白1的mRNA序列、氨基酸序列分别如序列号1、2所示。另外,踝蛋白2的mRNA序列、氨基酸序列分别如序列号3、4所示。

[0029] 受试动物的血浆中或血清中的踝蛋白量可利用免疫化学方法、使用与踝蛋白结合的抗体进行测定。

[0030] 与踝蛋白结合的抗体可以是多克隆抗体也可以是单克隆抗体,根据情况,可使用抗体的片段,例如Fab'、Fab、F(ab')<sub>2</sub>。这些抗体可利用现有公知的方法制备。

[0031] 作为市售品,可举出H-18抗体(Santa Cruz·Biotechnology公司)、H-300抗体(Santa Cruz·Biotechnology公司)、TA205抗体(Abcam公司)、M54246M抗体(Biodesign公司)等。

[0032] 踝蛋白量的测定可采用公知的酶免疫分析(EIA)、化学发光免疫分析、放射免疫分析(RIA)、荧光免疫分析、胶乳凝集法等方法来实施。具体而言,可举出例如使用抗体及标记抗原的竞争法、组合使用针对抗原的识别部位不同的两种单克隆抗体或多克隆抗体(或单克隆抗体及多克隆抗体)的夹心EIA法、使用固定有抗体的胶乳粒子的胶乳凝集法等。

[0033] 在这些测定法中,根据需要,可将抗原或抗体固定在适当的固相载体上。作为固相载体,可举出例如聚苯乙烯、聚乙烯、聚丙烯、聚氯乙烯、聚酯、聚丙烯酸酯、尼龙、聚缩醛、含氟树脂等合成树脂,纤维素、琼脂糖等多糖类,玻璃、金属等。上述固相载体可形成微孔板状、球状、纤维状、棒状、盘状、容器状、盒状(cell)、试管等多种形状。

[0034] 上述那样的免疫化学方法中,抗体或抗原根据需要使用经标记的物质。作为所述标记,可举出酶(过氧化物酶、碱性磷酸酶等)、发光物质(吖啶酯、异氨基苯二酰肼、荧光素等),此外,可举出放射性同位素(<sup>124</sup>I、<sup>14</sup>C、<sup>3</sup>H)、荧光物质(异硫氰酸荧光素等)等。此外,还可采用组合生物素标记和链亲和素来使用的方法。

[0035] 通过如上所述对受试动物血浆中或血清中的踝蛋白量进行测定、定量,可简便地诊断是否罹患类风湿性关节炎。即,血浆中或血清中的踝蛋白量高于规定的阈值时,可判断为罹患类风湿性关节炎。所述规定的阈值例如可基于未患有类风湿性关节炎的对照动物的血浆或血清中的平均值等设定。

[0036] 另外,通过对给予类风湿性关节炎治疗药前后的踝蛋白量进行测定、定量,可简便地判断该治疗药的治疗效果。即,如果给予类风湿性关节炎治疗药之后的踝蛋白量与给药之前的踝蛋白量相比显著降低,则判断该治疗药是有效的。

[0037] 此处,作为类风湿性关节炎治疗药,可包括现有公知的治疗药以及今后开发出的所有治疗药。作为现有公知的类风湿性关节炎治疗药,可举出例如生物学制剂、非甾体性抗炎药(消炎镇痛药)、甾体药、免疫抑制剂等。

[0038] 作为生物学制剂,可举出例如嵌合型抗TNF- $\alpha$ 抗体制剂、可溶性TNF受体、或完全人型抗TNF- $\alpha$ 抗体制剂、抗IL-6受体抗体制剂等。作为非甾体性抗炎药,可举出前列腺素产生抑制剂,其被认为虽可减轻关节疼痛或肿胀,但难以抑制疾病自身的恶化和骨、关节的破坏。甾体药由于其优异的抗炎效果而作为类风湿性关节炎的特效药被利用,但其副作用也是问题。免疫抑制剂通过改善类风湿性关节炎患者的免疫异常来抑制类风湿性关节炎的

炎症,其目的是诱导缓解,存在遏制类风湿性关节炎恶化的可能性,因此也被称为疾病修饰性抗风湿药。由于效果表现需要时间,所以也被称为迟效性抗风湿药。

[0039] 虽然如上所述存在多种类风湿性关节炎治疗药,但为了判定治疗药的效果的程度、选择最有效的治疗药,本发明涉及的检查方法是有用的。

[0040] < 类风湿性关节炎检查用试剂盒 >

[0041] 本发明涉及的类风湿性关节炎检查用试剂盒,用于在本发明涉及的类风湿性关节炎的检查方法中使用。该诊断用试剂盒中,例如,包含固定有与踝蛋白结合的抗体的固相载体。另外,还可包含经标记的二抗和显色基质等。

[0042] 实施例

[0043] 以下,利用实施例详细地说明本发明,但如下记载不作任何限制本发明的解释。需要说明的是,以下的实施例 1,2、比较例 1 中,以类风湿性关节炎患者(RA 患者)17 例、对照 14 例(变形性关节症患者 8 例、全身性红斑狼疮患者 1 例、糖尿病患者 1 例、健康人 4 例)为受试者。另外,以下的实施例 3 中,以 RA 患者 5 例为受试者。

[0044] < 实施例 1 >

[0045] 将各受试者的血液采集至 EDTA 管中,在室温下以 2500rpm 的转速进行 10 分钟的离心分离,由此得到血浆。

[0046] 利用夹心 ELISA 法测定上述血浆中的踝蛋白量。

[0047] 首先,用磷酸缓冲液(PBS)将识别踝蛋白的 N 末端的 H-18 抗体(Santa Cruz · Biotechnology 公司)稀释为 1  $\mu$ g/mL,然后按照 100  $\mu$ L/孔的量加入到 96 孔微孔板中,在 4 $^{\circ}$ C 下孵育一夜,然后用 200  $\mu$ L/孔的洗涤液洗涤 3 次。接下来,将各受试者的血浆按照 100  $\mu$ L/孔的量加入到 96 孔微孔板中,在 25 $^{\circ}$ C 下孵育 1 小时,然后用 200  $\mu$ L/孔的洗涤液洗涤 3 次。接下来,用 PBS 稀释作为一抗的识别踝蛋白的 N 末端的 H-300 抗体(Santa Cruz · Biotechnology 公司),使其为 2  $\mu$ g/mL,然后按照 100  $\mu$ L/孔的量加入到 96 孔微孔板中,在 25 $^{\circ}$ C 下孵育 1 小时,用 200  $\mu$ L/孔的洗涤液洗涤 3 次。接下来,用 PBS 稀释作为二抗的 HRP 标记抗山羊 IgG 抗体(KPL 公司),使其为 2  $\mu$ g/mL,按照 100  $\mu$ L/孔的量加入到 96 孔微孔板中,在 25 $^{\circ}$ C 下孵育 1 小时,然后用 200  $\mu$ L/孔的洗涤液洗涤 3 次。

[0048] 接下来,按照 100  $\mu$ L/孔的量将基质加入到 96 孔微孔板中,在 25 $^{\circ}$ C 下孵育 15 分钟,然后使用酶标仪测定波长 630nm 的 OD 值。

[0049] 实施例 1 的 ROC 曲线如图 1 所示。ROC 分析的结果,图 1 的 ROC 曲线下面积(AUC)为 0.954。另外,以 OD = 0.20 为界限值时的阳性、阴性分别如下表 1 所示。

[0050] [表 1]

[0051]

H-300 抗体		RA 患者	对照	合计
	阳性	14	0	14
	阴性	3	14	17
	合计	17	14	31

[0052] 由该结果可知,利用使用了 H-18 抗体及 H-300 抗体的夹心 ELISA 法进行的类风湿性关节炎诊断的灵敏度为  $14/17 \times 100 = 82.4\%$ ,特异性为  $14/14 \times 100 = 100\%$ 。

[0053] < 实施例 2 >

[0054] 除了使用识别踝蛋白的 C 末端的 M54246M 抗体 (Biodesign 公司) 作为一抗之外, 与实施例 1 同样地操作, 利用夹心 ELISA 法测定血浆中的踝蛋白量。

[0055] 实施例 2 的 ROC 曲线如图 2 所示。ROC 分析的结果, 图 2 的 ROC 曲线下面积 (AUC) 为 0.819。另外, 以 OD = 0.05 为界限值时的阳性、阴性分别如下表 2 所示。

[0056] [表 2]

[0057]

M54246M 抗体		RA 患者	对照	合计
	阳性	15	3	18
	阴性	2	11	13
	合计	17	11	31

[0058] 由该结果可知, 利用使用了 H-18 抗体及 M54246M 抗体的夹心 ELISA 法进行的类风湿性关节炎诊断的灵敏度为  $15/17 \times 100 = 88.2\%$ 、特异性为  $11/14 \times 100 = 78.6\%$ 。

[0059] <比较例 1>

[0060] 将各受试者的血液采集至血清用采血管中, 在室温下以 2,500rpm 的转速进行 10 分钟离心分离, 由此得到血清。使用市售试剂盒 (MESACUP CCP、MBL 公司) 测定该血清中的抗 CCP 抗体效价。

[0061] 比较例 1 的 ROC 曲线如图 3 所示。ROC 分析的结果, 图 3 的 ROC 曲线下面积 (AUC) 为 0.838。另外, 以抗体效价 = 6.60 为界限值时的阳性、阴性分别如下表 3 所示。

[0062] [表 3]

[0063]

抗 CCP 抗体		RA 患者	对照	合计
	阳性	11	1	12
	阴性	6	13	19
	合计	17	14	31

[0064] 由该结果可知, 使用了抗 CCP 抗体的类风湿性关节炎诊断的灵敏度为  $11/17 \times 100 = 64.7\%$ 、特异性为  $13/14 \times 100 = 92.9\%$ 。

[0065] 由以上结果可知, RA 患者的血中, 踝蛋白以优势地位存在, 因此, 可通过测定血中的踝蛋白量, 简便地检查是否患有类风湿性关节炎。而且, 该检查方法比使用了抗 CCP 抗体的现有方法灵敏度优异。

[0066] <实施例 3>

[0067] 利用测定踝蛋白量确认了类风湿性关节炎治疗药对 5 名 RA 患者的治疗效果。与实施例 1 同样, 通过利用使用了 H-18 抗体及 H-300 抗体的夹心 ELISA 法测定 OD 值来进行踝蛋白量的测定。另外, 按照与通常的临床检查相同的方法也进行了 CRP 量、MMP-3 量的测定。进一步地, 还计算了欧洲风湿病防治联合会 (EULAR) 推荐的 DAS (Disease Activity Score) 28 的分数。就 DAS28 的分数而言, 5.1 以上判断为高疾病活动性, 3.2 以上且小于 5.1 判断为中等疾病活动性, 小于 3.2 判断为低疾病活动性。结果如下表 4 所示。

[0068] [表 4]

[0069]

病例 1·男性 (ADA 显著效果例)		前 (MTX 治疗)	后 (MTX+ADA 治疗)
	踝蛋白 (OD 值)	0.568	0.139
	CRP (mg/dL)	1.74	0.08
	MMP-3 (mg/mL)	60.6	57.4
		5.43	2.62
病例 2·女性 (SASP 无效例)		前 (未治疗)	后 (SAP 治疗)
	踝蛋白 (OD 值)	0.258	0.294
	CRP (mg/dL)	0.36	0.19
	MMP-3 (mg/mL)	70.6	106.6
		5.12	4.21
病例 3·女性 (IFX 无效例)		前 (MTX 治疗)	后 (MTX+IFX 治疗)
	踝蛋白 (OD 值)	0.205	0.223
	CRP (mg/dL)	2.82	2.68
	MMP-3 (mg/mL)	962.1	523.8
		4.87	3.95
病例 4·女性 (ADA 显著效果例)		前 (BUC 治疗)	后 (BUC+ADA 治疗)
	踝蛋白 (OD 值)	0.258	0.164
	CRP (mg/dL)	5.01	0.16
	MMP-3 (mg/mL)	323.2	52.2
		4.55	2.45
病例 5·女性 (TCZ 无效例)		前 (未治疗)	后 (TCZ 治疗)
	踝蛋白 (OD 值)	2.093	1.787
	CRP (mg/dL)	3.83	2.78
	MMP-3 (mg/mL)	117.8	93.1
		5.09	4.50

[0070] 病例 1 是通过在使用了 MTX(氨甲喋呤)的治疗中联用 ADA(阿达木单抗),当适用 EULAR 的反应性标准时为良好应答者 (good responder) (DAS28 :5.43 → 2.62) 的一个例子。踝蛋白量在仅为 MTX 的情况下为高值 (OD 值 :0.568),但通过联用 ADA 而变为正常值 (OD 值 :0.139)。另一方面, MMP-3 量未由于联用 ADA 而表现显著降低,未反映类风湿性关节炎的病情。

[0071] 病例 2 是即使利用使用了 SASP(柳氮磺吡啶)的治疗,当适用 EULAR 的反应性标准时也为无应答者 (none responder) (DAS28 :5.12 → 4.21) 的一个例子。踝蛋白量在利用 SASP 治疗后仍保持为高值 (OD 值 :0.258 → 0.294)。另一方面, CRP 量在 SASP 治疗后变为正常值 (0.19mg/dL),未反映类风湿性关节炎的病情。

[0072] 病例 3 是尽管在使用了 MTX(氨甲喋呤)的治疗中联用了 IFX(英夫利昔单抗),但仍为无应答者 (DAS28 :4.87 → 3.95) 的一个例子。踝蛋白量在联用 IFX 后仍保持为高值 (OD 值 :0.205 → 0.223)。

[0073] 病例 4 是通过在使用了 BUC(布西拉明)的治疗中联用 ADA(阿达木单抗)而为良好应答者 (DAS28 :4.55 → 2.45) 的一个例子。踝蛋白量在仅使用 BUC 时为高值 (OD 值 :0.258),但通过联用 ADA 而变为正常值 (OD 值 :0.164)。

[0074] 病例 5 是即使利用使用了 TCZ(托珠单抗 (tocilizumab)) 的治疗,也为无应答者 (DAS28 :5.09 → 4.50) 的一个例子。踝蛋白量在利用 TCZ 治疗后仍为高值 (OD 值 :

2.093 → 1.787)。

[0075] 由以上结果可知,血中踝蛋白量与类风湿性关节炎的病情相关,而且,比 CRP、MMP-3 等其他因子更准确地反映类风湿性关节炎的病情。因此,通过测定血中踝蛋白量,可简便且准确地判断类风湿性关节炎治疗药的治疗效果。

[0001]

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Met Gln Phe Glu Pro Ser Thr Met Val Tyr Asp Ala Cys Arg Ile Ile
20 25 30
Arg Glu Arg Ile Pro Glu Ala Pro Ala Gly Pro Pro Ser Asp Phe Gly
35 40 45
Leu Phe Leu Ser Asp Asp Asp Pro Lys Lys Gly Ile Trp Leu Glu Ala
50 55 60
Gly Lys Ala Leu Asp Tyr Tyr Met Leu Arg Asn Gly Asp Thr Met Glu
65 70 75 80
Tyr Arg Lys Lys Gln Arg Pro Leu Lys Ile Arg Met Leu Asp Gly Thr
85 90 95
Val Lys Thr Ile Met Val Asp Asp Ser Lys Thr Val Thr Asp Met Leu
100 105 110
Met Thr Ile Cys Ala Arg Ile Gly Ile Thr Asn His Asp Glu Tyr Ser
115 120 125
Leu Val Arg Glu Leu Met Glu Glu Lys Lys Glu Glu Ile Thr Gly Thr
130 135 140
Leu Arg Lys Asp Lys Thr Leu Leu Arg Asp Glu Lys Lys Met Glu Lys
145 150 155 160
Leu Lys Gln Lys Leu His Thr Asp Asp Glu Leu Asn Trp Leu Asp His
165 170 175
Gly Arg Thr Leu Arg Glu Gln Gly Val Glu Glu His Glu Thr Leu Leu
180 185 190
Leu Arg Arg Lys Phe Phe Tyr Ser Asp Gln Asn Val Asp Ser Arg Asp
195 200 205
Pro Val Gln Leu Asn Leu Leu Tyr Val Gln Ala Arg Asp Asp Ile Leu
210 215 220

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[0006]

Asn 225	Gly	Ser	His	Pro	Val 230	Ser	Phe	Asp	Lys	Ala 235	Cys	Glu	Phe	Ala	Gly 240
Phe	Gln	Cys	Gln 245	Ile	Gln	Phe	Gly	Pro	His 250	Asn	Glu	Gln	Lys	His 255	Lys
Ala	Gly	Phe	Leu 260	Asp	Leu	Lys	Asp	Phe	Leu 265	Pro	Lys	Glu	Tyr 270	Val	Lys
Gln	Lys	Gly 275	Glu	Arg	Lys	Ile	Phe 280	Gln	Ala	His	Lys	Asn 285	Cys	Gly	Gln
Met	Ser 290	Glu	Ile	Glu	Ala	Lys 295	Val	Arg	Tyr	Val	Lys 300	Leu	Ala	Arg	Ser
Leu	Lys 305	Thr	Tyr	Gly	Val 310	Ser	Phe	Phe	Leu	Val 315	Lys	Glu	Lys	Met	Lys 320
Gly	Lys	Asn	Lys 325	Leu	Val	Pro	Arg	Leu	Leu 330	Gly	Ile	Thr	Lys	Glu	Cys 335
Val	Met	Arg	Val 340	Asp	Glu	Lys	Thr	Lys	Glu 345	Val	Ile	Gln	Glu	Trp	Asn 350
Leu	Thr 355	Asn	Ile	Lys	Arg	Trp	Ala 360	Ala	Ser	Pro	Lys	Ser	Phe	Thr	Leu
Asp	Phe 370	Gly	Asp	Tyr	Gln	Asp 375	Gly	Tyr	Tyr	Ser	Val 380	Gln	Thr	Thr	Glu
Gly 385	Glu	Gln	Ile	Ala	Gln 390	Leu	Ile	Ala	Gly	Tyr 395	Ile	Asp	Ile	Ile	Leu 400
Lys	Lys	Lys	Lys 405	Ser	Lys	Asp	His	Phe	Gly 410	Leu	Glu	Gly	Asp	Glu	Glu 415
Ser	Thr	Met	Leu 420	Glu	Asp	Ser	Val	Ser	Pro 425	Lys	Lys	Ser	Thr	Val	Leu
Gln	Gln 435	Gln	Tyr	Asn	Arg	Val	Gly 440	Lys	Val	Glu	His	Gly 445	Ser	Val	Ala
Leu	Pro 450	Ala	Ile	Met	Arg	Ser 455	Gly	Ala	Ser	Gly	Pro 460	Glu	Asn	Phe	Gln
Val 465	Gly	Ser	Met	Pro	Pro 470	Ala	Gln	Gln	Gln	Ile 475	Thr	Ser	Gly	Gln	Met 480
His	Arg	Gly	His 485	Met	Pro	Pro	Leu	Thr	Ser 490	Ala	Gln	Gln	Ala	Leu	Thr 495
Gly	Thr	Ile	Asn 500	Ser	Ser	Met	Gln	Ala	Val 505	Gln	Ala	Ala	Gln	Ala	Thr
Leu	Asp	Asp 515	Phe	Asp	Thr	Leu	Pro 520	Pro	Leu	Gly	Gln	Asp 525	Ala	Ala	Ser
Lys	Ala 530	Trp	Arg	Lys	Asn	Lys 535	Met	Asp	Glu	Ser	Lys 540	His	Glu	Ile	His
Ser 545	Gln	Val	Asp	Ala	Ile 550	Thr	Ala	Gly	Thr	Ala 555	Ser	Val	Val	Asn	Leu 560
Thr	Ala	Gly	Asp 565	Pro	Ala	Glu	Thr	Asp	Tyr 570	Thr	Ala	Val	Gly	Cys	Ala 575
Val	Thr	Thr	Ile 580	Ser	Ser	Asn	Leu	Thr 585	Glu	Met	Ser	Arg	Gly	Val	Lys 590

[0007]

Leu Leu Ala Ala Leu Leu Glu Asp Glu Gly Gly Ser Gly Arg Pro Leu  
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Leu Gln Ala Ala Lys Gly Leu Ala Gly Ala Val Ser Glu Leu Leu Arg  
 610 615 620

Ser Ala Gln Pro Ala Ser Ala Glu Pro Arg Gln Asn Leu Leu Gln Ala  
 625 630 635 640

Ala Gly Asn Val Gly Gln Ala Ser Gly Glu Leu Leu Gln Gln Ile Gly  
 645 650 655

Glu Ser Asp Thr Asp Pro His Phe Gln Asp Ala Leu Met Gln Leu Ala  
 660 665 670

Lys Ala Val Ala Ser Ala Ala Ala Leu Val Leu Lys Ala Lys Ser  
 675 680 685

Val Ala Gln Arg Thr Glu Asp Ser Gly Leu Gln Thr Gln Val Ile Ala  
 690 695 700

Ala Ala Thr Gln Cys Ala Leu Ser Thr Ser Gln Leu Val Ala Cys Thr  
 705 710 715 720

Lys Val Val Ala Pro Thr Ile Ser Ser Pro Val Cys Gln Glu Gln Leu  
 725 730 735

Val Glu Ala Gly Arg Leu Val Ala Lys Ala Val Glu Gly Cys Val Ser  
 740 745 750

Ala Ser Gln Ala Ala Thr Glu Asp Gly Gln Leu Leu Arg Gly Val Gly  
 755 760 765

Ala Ala Ala Thr Ala Val Thr Gln Ala Leu Asn Glu Leu Leu Gln His  
 770 775 780

Val Lys Ala His Ala Thr Gly Ala Gly Pro Ala Gly Arg Tyr Asp Gln  
 785 790 795 800

Ala Thr Asp Thr Ile Leu Thr Val Thr Glu Asn Ile Phe Ser Ser Met  
 805 810 815

Gly Asp Ala Gly Glu Met Val Gly Gln Ala Arg Ile Leu Ala Gln Ala  
 820 825 830

Thr Ser Asp Leu Val Asn Ala Ile Lys Ala Asp Ala Glu Gly Glu Ser  
 835 840 845

Asp Leu Glu Asn Ser Arg Lys Leu Leu Ser Ala Ala Lys Ile Leu Ala  
 850 855 860

Asp Ala Thr Ala Lys Met Val Glu Ala Ala Lys Gly Ala Ala Ala His  
 865 870 875 880

Pro Asp Ser Glu Glu Gln Gln Gln Arg Leu Arg Glu Ala Ala Glu Gly  
 885 890 895

Leu Arg Met Ala Thr Asn Ala Ala Ala Gln Asn Ala Ile Lys Lys Lys  
 900 905 910

Leu Val Gln Arg Leu Glu His Ala Ala Lys Gln Ala Ala Ala Ser Ala  
 915 920 925

Thr Gln Thr Ile Ala Ala Ala Gln His Ala Ala Ser Thr Pro Lys Ala  
 930 935 940

Ser Ala Gly Pro Gln Pro Leu Leu Val Gln Ser Cys Lys Ala Val Ala  
 945 950 955 960

[0008]

Glu Gln Ile Pro Leu Leu Val Gln Gly Val Arg Gly Ser Gln Ala Gln  
 965 970 975  
 Pro Asp Ser Pro Ser Ala Gln Leu Ala Leu Ile Ala Ala Ser Gln Ser  
 980 985 990  
 Phe Leu Gln Pro Gly Gly Lys Met Val Ala Ala Ala Lys Ala Ser Val  
 995 1000 1005  
 Pro Thr Ile Gln Asp Gln Ala Ser Ala Met Gln Leu Ser Gln Cys Ala  
 1010 1015 1020  
 Lys Asn Leu Gly Thr Ala Leu Ala Glu Leu Arg Thr Ala Ala Gln Lys  
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 Ala Gln Glu Ala Cys Gly Pro Leu Glu Met Asp Ser Ala Leu Ser Val  
 1045 1050 1055  
 Val Gln Asn Leu Glu Lys Asp Leu Gln Glu Val Lys Ala Ala Ala Arg  
 1060 1065 1070  
 Asp Gly Lys Leu Lys Pro Leu Pro Gly Glu Thr Met Glu Lys Cys Thr  
 1075 1080 1085  
 Gln Asp Leu Gly Asn Ser Thr Lys Ala Val Ser Ser Ala Ile Ala Gln  
 1090 1095 1100  
 Leu Leu Gly Glu Val Ala Gln Gly Asn Glu Asn Tyr Ala Gly Ile Ala  
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 Ala Arg Asp Val Ala Gly Gly Leu Arg Ser Leu Ala Gln Ala Ala Arg  
 1125 1130 1135  
 Gly Val Ala Ala Leu Thr Ser Asp Pro Ala Val Gln Ala Ile Val Leu  
 1140 1145 1150  
 Asp Thr Ala Ser Asp Val Leu Asp Lys Ala Ser Ser Leu Ile Glu Glu  
 1155 1160 1165  
 Ala Lys Lys Ala Ala Gly His Pro Gly Asp Pro Glu Ser Gln Gln Arg  
 1170 1175 1180  
 Leu Ala Gln Val Ala Lys Ala Val Thr Gln Ala Leu Asn Arg Cys Val  
 1185 1190 1195 1200  
 Ser Cys Leu Pro Gly Gln Arg Asp Val Asp Asn Ala Leu Arg Ala Val  
 1205 1210 1215  
 Gly Asp Ala Ser Lys Arg Leu Leu Ser Asp Ser Leu Pro Pro Ser Thr  
 1220 1225 1230  
 Gly Thr Phe Gln Glu Ala Gln Ser Arg Leu Asn Glu Ala Ala Ala Gly  
 1235 1240 1245  
 Leu Asn Gln Ala Ala Thr Glu Leu Val Gln Ala Ser Arg Gly Thr Pro  
 1250 1255 1260  
 Gln Asp Leu Ala Arg Ala Ser Gly Arg Phe Gly Gln Asp Phe Ser Thr  
 1265 1270 1275 1280  
 Phe Leu Glu Ala Gly Val Glu Met Ala Gly Gln Ala Pro Ser Gln Glu  
 1285 1290 1295  
 Asp Arg Ala Gln Val Val Ser Asn Leu Lys Gly Ile Ser Met Ser Ser  
 1300 1305 1310  
 Ser Lys Leu Leu Leu Ala Ala Lys Ala Leu Ser Thr Asp Pro Ala Ala  
 1315 1320 1325

[0009]

Pro Asn Leu Lys Ser Gln Leu Ala Ala Ala Ala Arg Ala Val Thr Asp  
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 Ser Ile Asn Gln Leu Ile Thr Met Cys Thr Gln Gln Ala Pro Gly Gln  
 1345 1350 1355 1360  
 Lys Glu Cys Asp Asn Ala Leu Arg Glu Leu Glu Thr Val Arg Glu Leu  
 1365 1370 1375  
 Leu Glu Asn Pro Val Gln Pro Ile Asn Asp Met Ser Tyr Phe Gly Cys  
 1380 1385 1390  
 Leu Asp Ser Val Met Glu Asn Ser Lys Val Leu Gly Glu Ala Met Thr  
 1395 1400 1405  
 Gly Ile Ser Gln Asn Ala Lys Asn Gly Asn Leu Pro Glu Phe Gly Asp  
 1410 1415 1420  
 Ala Ile Ser Thr Ala Ser Lys Ala Leu Cys Gly Phe Thr Glu Ala Ala  
 1425 1430 1435 1440  
 Ala Gln Ala Ala Tyr Leu Val Gly Val Ser Asp Pro Asn Ser Gln Ala  
 1445 1450 1455  
 Gly Gln Gln Gly Leu Val Glu Pro Thr Gln Phe Ala Arg Ala Asn Gln  
 1460 1465 1470  
 Ala Ile Gln Met Ala Cys Gln Ser Leu Gly Glu Pro Gly Cys Thr Gln  
 1475 1480 1485  
 Ala Gln Val Leu Ser Ala Ala Thr Ile Val Ala Lys His Thr Ser Ala  
 1490 1495 1500  
 Leu Cys Asn Ser Cys Arg Leu Ala Ser Ala Arg Thr Thr Asn Pro Thr  
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 Ala Lys Arg Gln Phe Val Gln Ser Ala Lys Glu Val Ala Asn Ser Thr  
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 Glu Asn Arg Ala Gln Cys Arg Ala Ala Thr Ala Pro Leu Leu Glu Ala  
 1555 1560 1565  
 Val Asp Asn Leu Ser Ala Phe Ala Ser Asn Pro Glu Phe Ser Ser Ile  
 1570 1575 1580  
 Pro Ala Gln Ile Ser Pro Glu Gly Arg Ala Ala Met Glu Pro Ile Val  
 1585 1590 1595 1600  
 Ile Ser Ala Lys Thr Met Leu Glu Ser Ala Gly Gly Leu Ile Gln Thr  
 1605 1610 1615  
 Ala Arg Ala Leu Ala Val Asn Pro Arg Asp Pro Pro Ser Trp Ser Val  
 1620 1625 1630  
 Leu Ala Gly His Ser Arg Thr Val Ser Asp Ser Ile Lys Lys Leu Ile  
 1635 1640 1645  
 Thr Ser Met Arg Asp Lys Ala Pro Gly Gln Leu Glu Cys Glu Thr Ala  
 1650 1655 1660  
 Ile Ala Ala Leu Asn Ser Cys Leu Arg Asp Leu Asp Gln Ala Ser Leu  
 1665 1670 1675 1680  
 Ala Ala Val Ser Gln Gln Leu Ala Pro Arg Glu Gly Ile Ser Gln Glu  
 1685 1690 1695

[0010]

Ala Leu His Thr Gln Met Leu Thr Ala Val Gln Glu Ile Ser His Leu  
1700 1705 1710

Ile Glu Pro Leu Ala Asn Ala Ala Arg Ala Glu Ala Ser Gln Leu Gly  
1715 1720 1725

His Lys Val Ser Gln Met Ala Gln Tyr Phe Glu Pro Leu Thr Leu Ala  
1730 1735 1740

Ala Val Gly Ala Ala Ser Lys Thr Leu Ser His Pro Gln Gln Met Ala  
1745 1750 1755 1760

Leu Leu Asp Gln Thr Lys Thr Leu Ala Glu Ser Ala Leu Gln Leu Leu  
1765 1770 1775

Tyr Thr Ala Lys Glu Ala Gly Gly Asn Pro Lys Gln Ala Ala His Thr  
1780 1785 1790

Gln Glu Ala Leu Glu Glu Ala Val Gln Met Met Thr Glu Ala Val Glu  
1795 1800 1805

Asp Leu Thr Thr Thr Leu Asn Glu Ala Ala Ser Ala Ala Gly Val Val  
1810 1815 1820

Gly Gly Met Val Asp Ser Ile Thr Gln Ala Ile Asn Gln Leu Asp Glu  
1825 1830 1835 1840

Gly Pro Met Gly Glu Pro Glu Gly Ser Phe Val Asp Tyr Gln Thr Thr  
1845 1850 1855

Met Val Arg Thr Ala Lys Ala Ile Ala Val Thr Val Gln Glu Met Val  
1860 1865 1870

Thr Lys Ser Asn Thr Ser Pro Glu Glu Leu Gly Pro Leu Ala Asn Gln  
1875 1880 1885

Leu Thr Ser Asp Tyr Gly Arg Leu Ala Ser Glu Ala Lys Pro Ala Ala  
1890 1895 1900

Val Ala Ala Glu Asn Glu Glu Ile Gly Ser His Ile Lys His Arg Val  
1905 1910 1915 1920

Gln Glu Leu Gly His Gly Cys Ala Ala Leu Val Thr Lys Ala Gly Ala  
1925 1930 1935

Leu Gln Cys Ser Pro Ser Asp Ala Tyr Thr Lys Lys Glu Leu Ile Glu  
1940 1945 1950

Cys Ala Arg Arg Val Ser Glu Lys Val Ser His Val Leu Ala Ala Leu  
1955 1960 1965

Gln Ala Gly Asn Arg Gly Thr Gln Ala Cys Ile Thr Ala Ala Ser Ala  
1970 1975 1980

Val Ser Gly Ile Ile Ala Asp Leu Asp Thr Thr Ile Met Phe Ala Thr  
1985 1990 1995 2000

Ala Gly Thr Leu Asn Arg Glu Gly Thr Glu Thr Phe Ala Asp His Arg  
2005 2010 2015

Glu Gly Ile Leu Lys Thr Ala Lys Val Leu Val Glu Asp Thr Lys Val  
2020 2025 2030

Leu Val Gln Asn Ala Ala Gly Ser Gln Glu Lys Leu Ala Gln Ala Ala  
2035 2040 2045

Gln Ser Ser Val Ala Thr Ile Thr Arg Leu Ala Asp Val Val Lys Leu  
2050 2055 2060

[0011]

Gly Ala Ala Ser Leu Gly Ala Glu Asp Pro Glu Thr Gln Val Val Leu  
 2065 2070 2075 2080  
 Ile Asn Ala Val Lys Asp Val Ala Lys Ala Leu Gly Asp Leu Ile Ser  
 2085 2090 2095  
 Ala Thr Lys Ala Ala Ala Gly Lys Val Gly Asp Asp Pro Ala Val Trp  
 2100 2105 2110  
 Gln Leu Lys Asn Ser Ala Lys Val Met Val Thr Asn Val Thr Ser Leu  
 2115 2120 2125  
 Leu Lys Thr Val Lys Ala Val Glu Asp Glu Ala Thr Lys Gly Thr Arg  
 2130 2135 2140  
 Ala Leu Glu Ala Thr Thr Glu His Ile Arg Gln Glu Leu Ala Val Phe  
 2145 2150 2155 2160  
 Cys Ser Pro Glu Pro Pro Ala Lys Thr Ser Thr Pro Glu Asp Phe Ile  
 2165 2170 2175  
 Arg Met Thr Lys Gly Ile Thr Met Ala Thr Ala Lys Ala Val Ala Ala  
 2180 2185 2190  
 Gly Asn Ser Cys Arg Gln Glu Asp Val Ile Ala Thr Ala Asn Leu Ser  
 2195 2200 2205  
 Arg Arg Ala Ile Ala Asp Met Leu Arg Ala Cys Lys Glu Ala Ala Tyr  
 2210 2215 2220  
 His Pro Glu Val Ala Pro Asp Val Arg Leu Arg Ala Leu His Tyr Gly  
 2225 2230 2235 2240  
 Arg Glu Cys Ala Asn Gly Tyr Leu Glu Leu Leu Asp His Val Leu Leu  
 2245 2250 2255  
 Thr Leu Gln Lys Pro Ser Pro Glu Leu Lys Gln Gln Leu Thr Gly His  
 2260 2265 2270  
 Ser Lys Arg Val Ala Gly Ser Val Thr Glu Leu Ile Gln Ala Ala Glu  
 2275 2280 2285  
 Ala Met Lys Gly Thr Glu Trp Val Asp Pro Glu Asp Pro Thr Val Ile  
 2290 2295 2300  
 Ala Glu Asn Glu Leu Leu Gly Ala Ala Ala Ala Ile Glu Ala Ala Ala  
 2305 2310 2315 2320  
 Lys Lys Leu Glu Gln Leu Lys Pro Arg Ala Lys Pro Lys Glu Ala Asp  
 2325 2330 2335  
 Glu Ser Leu Asn Phe Glu Glu Gln Ile Leu Glu Ala Ala Lys Ser Ile  
 2340 2345 2350  
 Ala Ala Ala Thr Ser Ala Leu Val Lys Ala Ala Ser Ala Ala Gln Arg  
 2355 2360 2365  
 Glu Leu Val Ala Gln Gly Lys Val Gly Ala Ile Pro Ala Asn Ala Leu  
 2370 2375 2380  
 Asp Asp Gly Gln Trp Ser Gln Gly Leu Ile Ser Ala Ala Arg Met Val  
 2385 2390 2395 2400  
 Ala Ala Ala Thr Asn Asn Leu Cys Glu Ala Ala Asn Ala Ala Val Gln  
 2405 2410 2415  
 Gly His Ala Ser Gln Glu Lys Leu Ile Ser Ser Ala Lys Gln Val Ala  
 2420 2425 2430

[0012]

Ala Ser Thr Ala Gln Leu Leu Val Ala Cys Lys Val Lys Ala Asp Gln  
 2435 2440 2445  
 Asp Ser Glu Ala Met Lys Arg Leu Gln Ala Ala Gly Asn Ala Val Lys  
 2450 2455 2460  
 Arg Ala Ser Asp Asn Leu Val Lys Ala Ala Gln Lys Ala Ala Ala Phe  
 2465 2470 2475 2480  
 Glu Glu Gln Glu Asn Glu Thr Val Val Val Lys Glu Lys Met Val Gly  
 2485 2490 2495  
 Gly Ile Ala Gln Ile Ile Ala Ala Gln Glu Glu Met Leu Arg Lys Glu  
 2500 2505 2510  
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[0013]

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[0018]

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Leu Glu Tyr Lys Lys Lys Gln Arg Pro Gln Lys Ile Arg Met Leu Asp
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Glu Lys Leu Lys Ala Lys Leu His Thr Asp Asp Asp Leu Asn Trp Leu
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His Lys Pro Gly Phe Leu Asp Leu Lys Glu Phe Leu Pro Lys Glu Tyr
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[0019]

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[0020]

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[0021]

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[0022]

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 Val Glu Ala Leu Gln Glu Gln Leu Thr Ser Val Val Gln Glu Ile Gly  
 1700 1705 1710  
 His Leu Ile Asp Pro Ile Ala Thr Ala Ala Arg Gly Glu Ala Ala Gln  
 1715 1720 1725  
 Leu Gly His Lys Val Thr Gln Leu Ala Ser Tyr Phe Glu Pro Leu Ile  
 1730 1735 1740  
 Leu Ala Ala Val Gly Val Ala Ser Lys Ile Leu Asp His Gln Gln Gln  
 1745 1750 1755 1760  
 Met Thr Val Leu Asp Gln Thr Lys Thr Leu Ala Glu Ser Ala Leu Gln  
 1765 1770 1775  
 Met Leu Tyr Ala Ala Lys Glu Gly Gly Gly Asn Pro Lys Ala Gln His  
 1780 1785 1790

[0023]

Thr His Asp Ala Ile Thr Glu Ala Ala Gln Leu Met Lys Glu Ala Val  
 1795 1800 1805  
 Asp Asp Ile Met Val Thr Leu Asn Glu Ala Ala Ser Glu Val Gly Leu  
 1810 1815 1820  
 Val Gly Gly Met Val Asp Ala Ile Ala Glu Ala Met Ser Lys Leu Asp  
 1825 1830 1835 1840  
 Glu Gly Thr Pro Pro Glu Pro Lys Gly Thr Phe Val Asp Tyr Gln Thr  
 1845 1850  
 Thr Val Val Lys Tyr Ser Lys Ala Ile Ala Val Thr Ala Gln Glu Met  
 1860 1865 1870  
 Met Thr Lys Ser Val Thr Asn Pro Glu Glu Leu Gly Gly Leu Ala Ser  
 1875 1880 1885  
 Gln Met Thr Ser Asp Tyr Gly His Leu Ala Phe Gln Gly Gln Met Ala  
 1890 1895 1900  
 Ala Ala Thr Ala Glu Pro Glu Glu Ile Gly Phe Gln Ile Arg Thr Arg  
 1905 1910 1915 1920  
 Val Gln Asp Leu Gly His Gly Cys Ile Phe Leu Val Gln Lys Ala Gly  
 1925 1930 1935  
 Ala Leu Gln Val Cys Pro Thr Asp Ser Tyr Thr Lys Arg Glu Leu Ile  
 1940 1945 1950  
 Glu Cys Ala Arg Ala Val Thr Glu Lys Val Ser Leu Val Leu Ser Ala  
 1955 1960 1965  
 Leu Gln Ala Gly Asn Lys Gly Thr Gln Ala Cys Ile Thr Ala Ala Thr  
 1970 1975 1980  
 Ala Val Ser Gly Ile Ile Ala Asp Leu Asp Thr Thr Ile Met Phe Ala  
 1985 1990 1995 2000  
 Thr Ala Gly Thr Leu Asn Ala Glu Asn Ser Glu Thr Phe Ala Asp His  
 2005 2010 2015  
 Arg Glu Asn Ile Leu Lys Thr Ala Lys Ala Leu Val Glu Asp Thr Lys  
 2020 2025 2030  
 Leu Leu Val Ser Gly Ala Ala Ser Thr Pro Asp Lys Leu Ala Gln Ala  
 2035 2040 2045  
 Ala Gln Ser Ser Ala Ala Thr Ile Thr Gln Leu Ala Glu Val Val Lys  
 2050 2055 2060  
 Leu Gly Ala Ala Ser Leu Gly Ser Asp Asp Pro Glu Thr Gln Val Val  
 2065 2070 2075 2080  
 Leu Ile Asn Ala Ile Lys Asp Val Ala Lys Ala Leu Ser Asp Leu Ile  
 2085 2090 2095  
 Ser Ala Thr Lys Gly Ala Ala Ser Lys Pro Val Asp Asp Pro Ser Met  
 2100 2105 2110  
 Tyr Gln Leu Lys Gly Ala Ala Lys Val Met Val Thr Asn Val Thr Ser  
 2115 2120 2125  
 Leu Leu Lys Thr Val Lys Ala Val Glu Asp Glu Ala Thr Arg Gly Thr  
 2130 2135 2140  
 Arg Ala Leu Glu Ala Thr Ile Glu Cys Ile Lys Gln Glu Leu Thr Val  
 2145 2150 2155 2160

[0024]

Phe Gln Ser Lys Asp Val Pro Glu Lys Thr Ser Ser Pro Glu Glu Ser  
 2165 2170 2175  
 Ile Arg Met Thr Lys Gly Ile Thr Met Ala Thr Ala Lys Ala Val Ala  
 2180 2185 2190  
 Ala Gly Asn Ser Cys Arg Gln Glu Asp Val Ile Ala Thr Ala Asn Leu  
 2195 2200 2205  
 Ser Arg Lys Ala Val Ser Asp Met Leu Thr Ala Cys Lys Gln Ala Ser  
 2210 2215 2220  
 Phe His Pro Asp Val Ser Asp Glu Val Arg Thr Arg Ala Leu Arg Phe  
 2225 2230 2235 2240  
 Gly Thr Glu Cys Thr Leu Gly Tyr Leu Asp Leu Leu Glu His Val Leu  
 2245 2250 2255  
 Val Ile Leu Gln Lys Pro Thr Pro Glu Phe Lys Gln Gln Leu Ala Ala  
 2260 2265 2270  
 Phe Ser Lys Arg Val Ala Gly Ala Val Thr Glu Leu Ile Gln Ala Ala  
 2275 2280 2285  
 Glu Ala Met Lys Gly Thr Glu Trp Val Asp Pro Glu Asp Pro Thr Val  
 2290 2295 2300  
 Ile Ala Glu Thr Glu Leu Leu Gly Ala Ala Ala Ser Ile Glu Ala Ala  
 2305 2310 2315 2320  
 Ala Lys Lys Leu Glu Gln Leu Lys Pro Arg Ala Lys Pro Lys Gln Ala  
 2325 2330 2335  
 Asp Glu Thr Leu Asp Phe Glu Glu Gln Ile Leu Glu Ala Ala Lys Ser  
 2340 2345 2350  
 Ile Ala Ala Ala Thr Ser Ala Leu Val Lys Ser Ala Ser Ala Ala Gln  
 2355 2360 2365  
 Arg Glu Leu Val Ala Gln Gly Lys Val Gly Ser Ile Pro Ala Asn Ala  
 2370 2375 2380  
 Ala Asp Asp Gly Gln Trp Ser Gln Gly Leu Ile Ser Ala Ala Arg Met  
 2385 2390 2395 2400  
 Val Ala Ala Ala Thr Ser Ser Leu Cys Glu Ala Ala Asn Ala Ser Val  
 2405 2410 2415  
 Gln Gly His Ala Ser Glu Glu Lys Leu Ile Ser Ser Ala Lys Gln Val  
 2420 2425 2430  
 Ala Ala Ser Thr Ala Gln Leu Leu Val Ala Cys Lys Val Lys Ala Asp  
 2435 2440 2445  
 Gln Asp Ser Glu Ala Met Arg Arg Leu Gln Ala Ala Gly Asn Ala Val  
 2450 2455 2460  
 Lys Arg Ala Ser Asp Asn Leu Val Arg Ala Ala Gln Lys Ala Ala Phe  
 2465 2470 2475 2480  
 Gly Lys Ala Asp Asp Asp Asp Val Val Val Lys Thr Lys Phe Val Gly  
 2485 2490 2495  
 Gly Ile Ala Gln Ile Ile Ala Ala Gln Glu Glu Met Leu Lys Lys Glu  
 2500 2505 2510  
 Arg Glu Leu Glu Glu Ala Arg Lys Lys Leu Ala Gln Ile Arg Gln Gln  
 2515 2520 2525  
 Gln Tyr Lys Phe Leu Pro Thr Glu Leu Arg Glu Asp Glu Gly  
 2530 2535 2540

[0025]

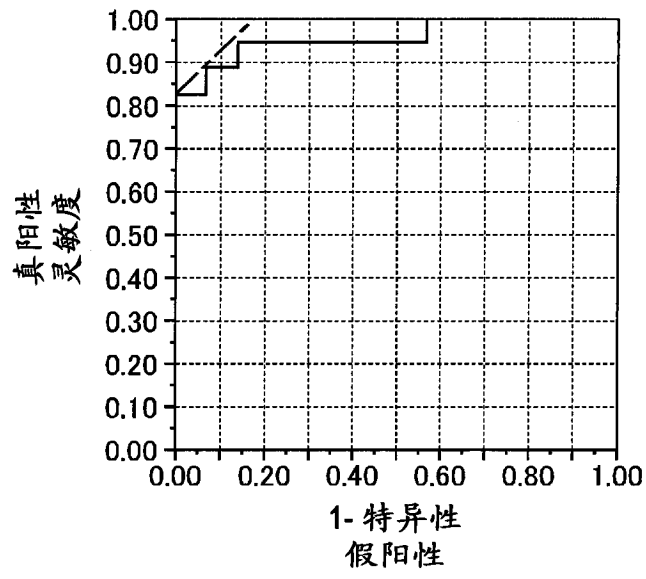


图 1

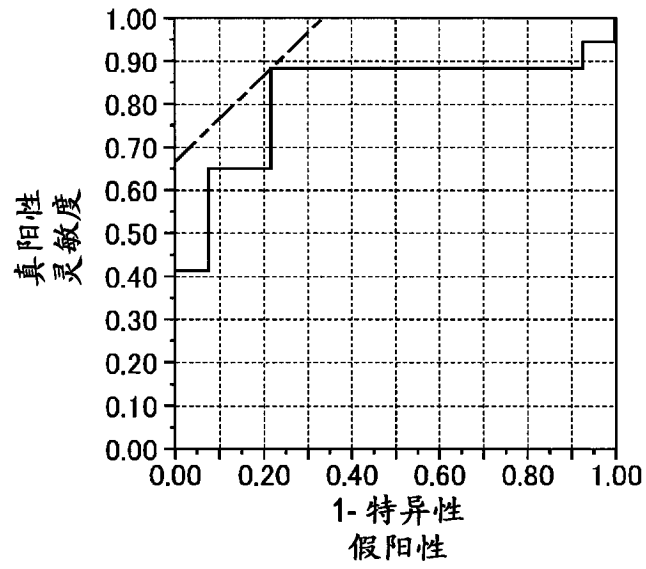


图 2

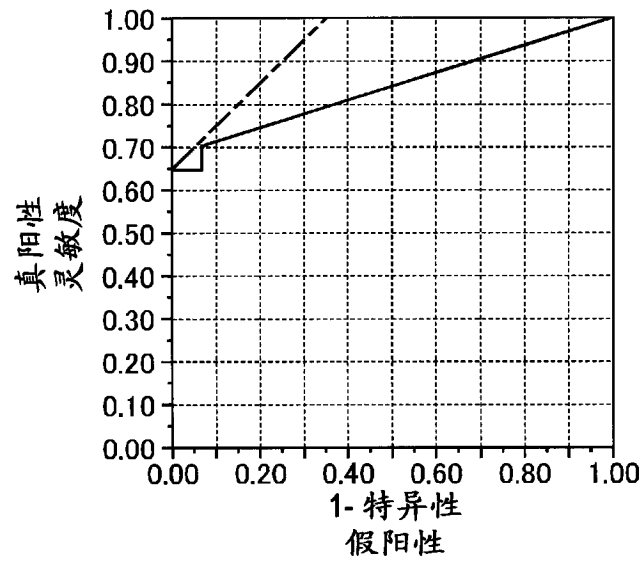


图 3

专利名称(译)	类风湿性关节炎的检查方法及类风湿性关节炎检查用试剂盒		
公开(公告)号	<a href="#">CN102971631B</a>	公开(公告)日	2015-12-16
申请号	CN201180024137.5	申请日	2011-06-14
[标]发明人	津坂宪政		
发明人	津坂宪政		
IPC分类号	G01N33/68 G01N33/53 G01N33/564		
CPC分类号	G01N33/564 G01N2333/4703 G01N2800/102		
代理人(译)	杨宏军		
优先权	2010279005 2010-12-15 JP		
其他公开文献	CN102971631A		
外部链接	<a href="#">Espacenet</a> <a href="#">SIPO</a>		

摘要(译)

本发明涉及新型的类风湿性关节炎的检查方法、及用于该检查方法的类风湿性关节炎检查用试剂盒。本发明涉及的类风湿性关节炎的检查方法的特征在于，包括测定受试动物的血浆中或血清中的踝蛋白量的步骤。该测定例如可利用使用了与踝蛋白结合的抗体的免疫学手段进行。本发明涉及的类风湿性关节炎检查用试剂盒用于在上述检查方法中使用，例如，包含固定有与踝蛋白结合的抗体的固相载体。

