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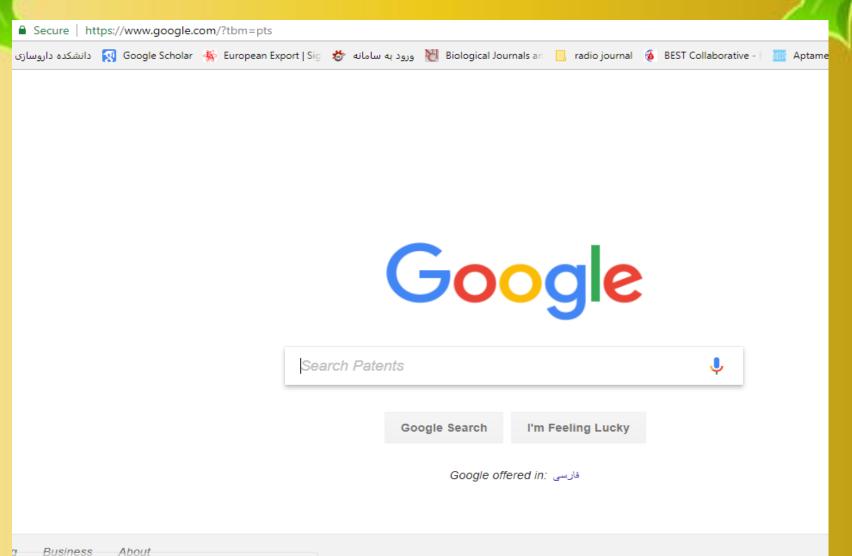
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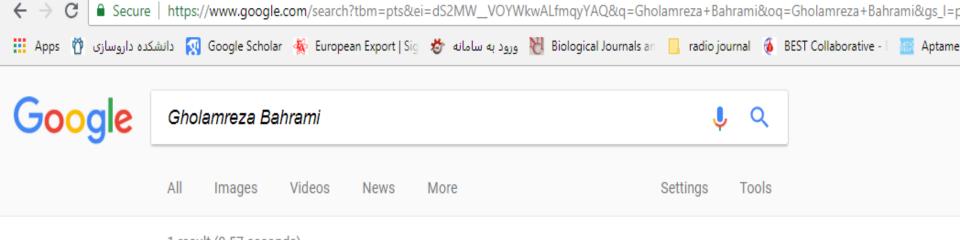


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Inventor: Gholamreza Bahrami; Original Assignee: Gholamreza Bahrami; Priority ...

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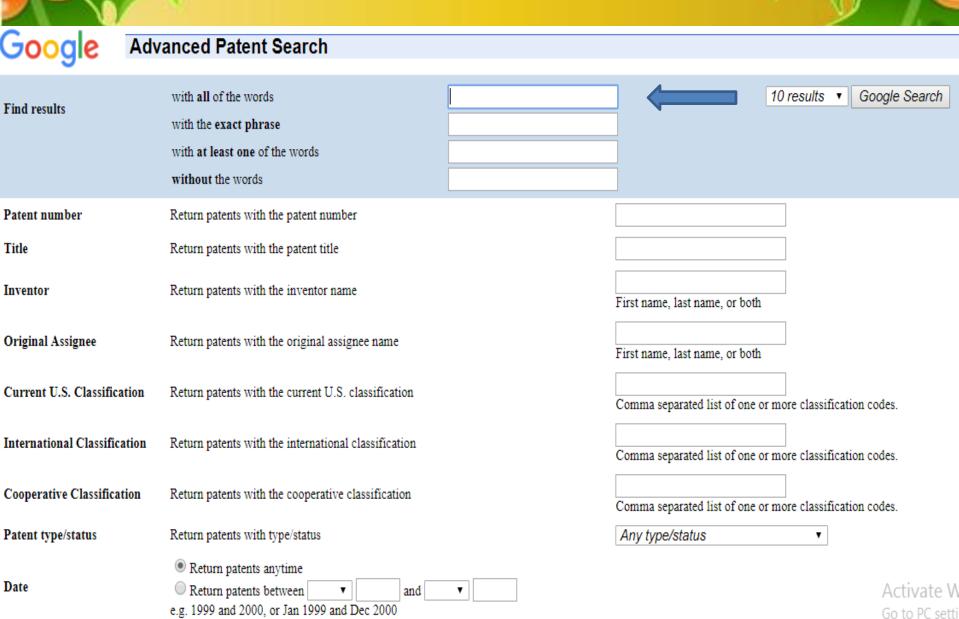
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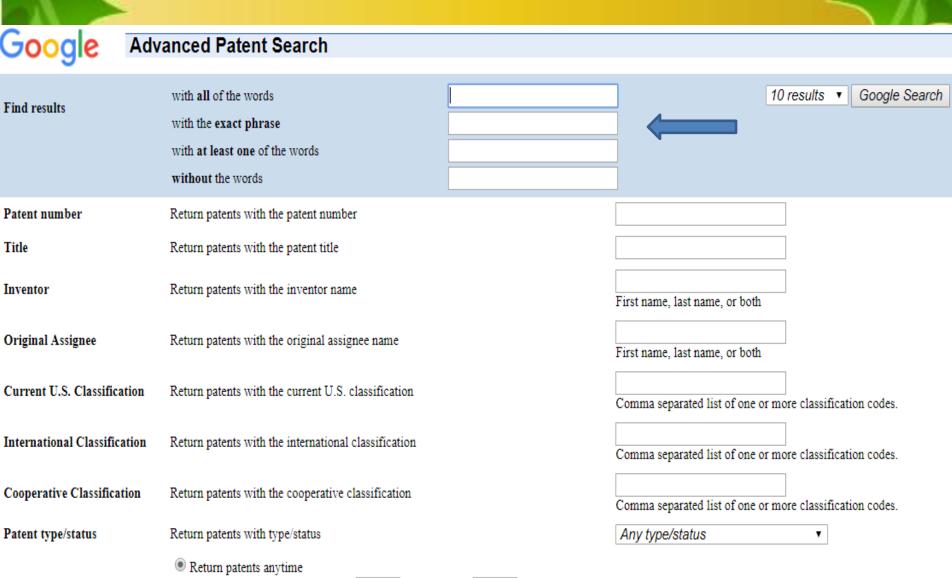
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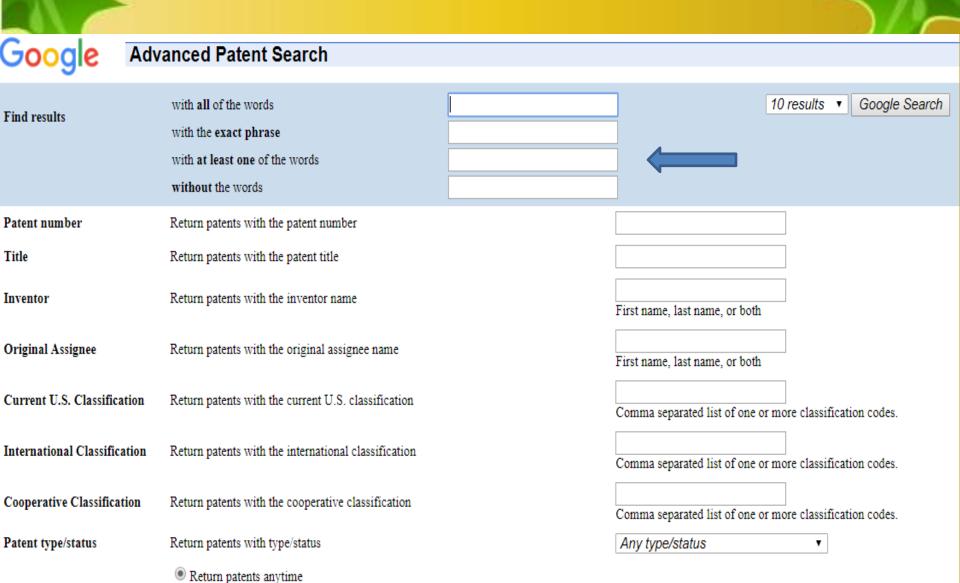
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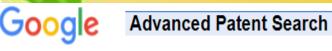
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A Sample





Gholamreza Bahrami





Herbal extract composition for the treatment of diabetes and a method of extracting the same

Abstract

The embodiments herein provide a method for isolation and purification of a novel oligosaccharide molecule from the fruits of Rosa arvensis for the treatment of diabetes. The fruit is dried, powdered and subjected to deionized water. The deionized water solution is subjected to silica gel column and the filtrate is subjected to ethanol-water mixture. The filtrate is subjected to column chromatography. The purified extract is structurally and chemically analyzed to find that the molecule is oligosaccharide. The purified oligosaccharide is administered to diabetic patients for 21 days. After the administration of oligosaccharide for 21 days the pancreatic tissues are collected and the blood glucose level in fasting and non fasting states is measured to find that there is significant effect in the blood glucose levels and the pancreatic tissue/cell regeneration after administration of the isolated oligosaccharide. The toxicity tests reveal that the oligosaccharide is non-toxic in nature.

Classifications

C08B37/0087 Glucomannans or galactomannans; Tara or tara gum, i.e. D-mannose and Dgalactose units, e.g. from Cesalpinia spinosa; Tamarind gum, i.e. D-galactose, D-glucose and D-xylose units, e.g. from Tamarindus indica; Gum Arabic, i.e. L-arabinose, L-rhamnose, Dgalactose and D-glucuronic acid units, e.g. from Acacia Senegal or Acacia Seyal; Derivatives thereof

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US9296831B2

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Inventor: Gholamreza Bahrami

Original Assignee: Gholamreza Bahrami

Priority date: 2014-01-26

Family: US (1)

Date

App/Pub Number

Status

2014-01-26

US14164223

Active

2014-09-11

US20140256673A1

Application

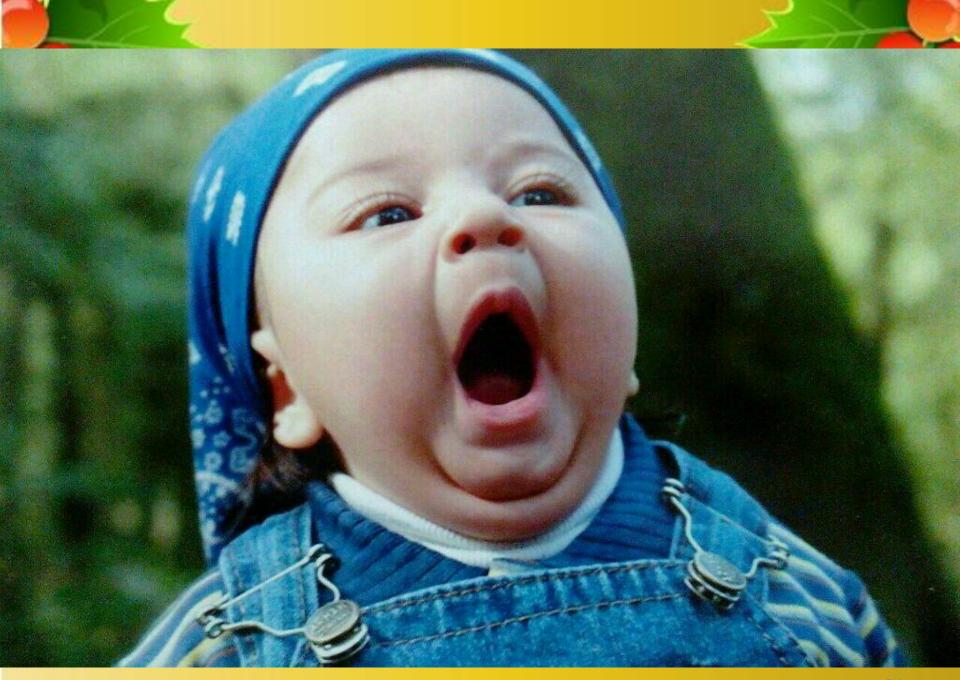
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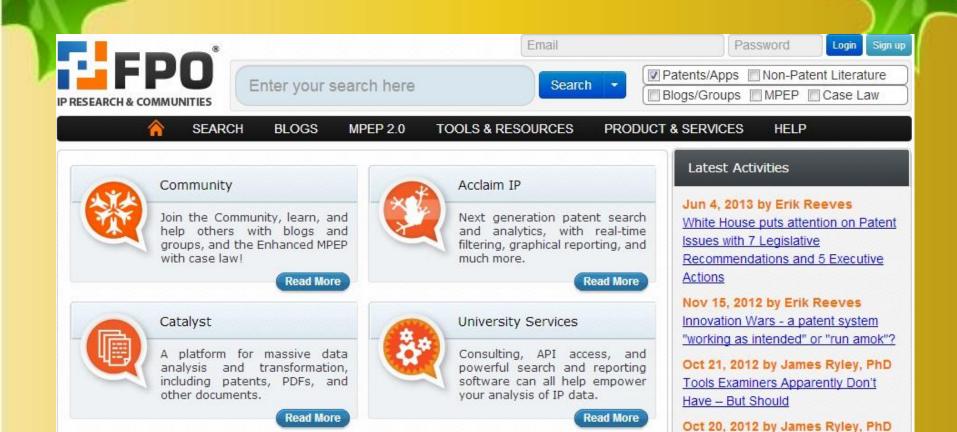
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| 2 | 9576692 | Method and device for producing 99mTc A method for producing 99mTc may include: providing a solution comprising 100Mo-molybdate-ions; providing a proton beam having an energy suitable for inducing a 100Mo(p,2n)99mTc-nuclear reaction | 965 |
| 3 | 9754694 | Method and device for producing a 99mTc reaction product A method for producing a reaction product containing 99mTC may include providing 100Mometal targets to be irradiated, irradiating the 100Mometal target with a proton stream having an energy for | 852 |
| 4 | 9125955 | 99mTc imaging agents and methods of use An embodiment of the invention comprises method of imaging a target site comprising administrating ligand of Formula I complexed to 99mTc wherein R1 and R2 are independently an alkyl or | 845 Activate W Go to PC setti |
| 5 | 9449726 | 100Mo compounds as accelerator targets for production of 99mTc | |

| Publication Date: | 07/04/2017 | | | | | |
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| Filing Date: | 10/14/2005 | | | | | |
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| Assignee: | GE Healthcare Limited (GB) | | | | | |
| International Classes: | A61K51/00; A61K51/04; A61M7 | | | | | |
| Field of Search: | 424/1.11, 424/1 | 1.49, 424/1.65, 50, 424/9.1, 424/1.77, 534/7, 534/10-16, 206/223, 206/569, 206/570 | | | | |
| View Patent Images: | Download PDI | F 9694092 PDF help | | | | |
| US Patent References: | 7052672 | Stabilized radiopharmaceutical compositions | 2006- 05-30 | Forster et | 424/1.6 | |
| | 7029653 | Method of the administration of drugs having binding affinity with plasma protein and | 2006- | Kawai et | | |
| | | preparation to be used in the method | 04-18 | al. | | |
| | 6713042 | Ascorbic acid analogs for metalloradiopharmaceuticals | 2004- 03-30 | Liu | 424/1.6 | |
| | 20040057899 | Stabilised radiopharmaceutical compositions | 2004- 03-25 | Forster et | | |
| | 6428768 | Stabilized radiopharmaceutical kits | 2002- 08-06 | DeRosch et al. | | |
| | 5300280 | Stabilized radiopharmaceutical kits | 1994- 04-05 | DeRosch et al. | 424/1.5 | |
| | 5045302 | Ligands and cationic complexes thereof with technetium-99m | 1991- 09-03 | Kelly et al | | |
| | 4489053 | Stable radioactive diagnostic agent and a non-radioactive carrier therefor | 1984- 12-18 | Azuma et | | |
| | 4439413 | Radioactive diagnostic agent for bone scanning and non-radioactive carrier therefor | 1984- | Hayashi | | |

| 0+ | تولید دانش فنی/ اختراع یا اکتشاف منجر به تولید و تجاری سازی محصول یا فرآیند با تأیید مراجع ذی صلاح وزار تین و نیز هر نوآوری که برای حل مشکلات و معضلات کشور مؤثر باشد و یا منجر به تولید خدمت یا محصول جدیدی در کشور شود. نحوه امتیازدهی به این فعالیتها به شرح موارد ۱ تا ۱۰ خواهد بود: | | |
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| | ی سیستهها، روشها و خدمات جدید به منظور حل معضلات آموزشی، پژوهشی، بهداشتی و درمانی در سطح منطقه با گواهی صلاح در منطقه آمایشی | | |
| | سازی وسایل پزشکی و آزمایشگاهی و ساخت هر قطعه که منجر به تولید دستگاه شود؛ با گواهی اداره کل تجهیزات پزشکی یک تا ۳ هداشت و تأیید هیات ممیزه مرکزی | | |
| | تولید مواد برای ازمایشهای پزشکی و تولیدات دارویی جدید از طریق مهندسی معکوس، به ترتیب با گواهی معاونت درمان یا ونت غذا و داروی وزارت بهداشت (حسب مورد) و تأییدهیأت ممیزه مرکزی وزارت بهداشت | | |
| | € 15 +/0 | ۵. انجام فعالیتهای جدید پزشکی که به خودکفایی کشور کمک مؤثر کند مانند اجرای روشهای تشخیصی - درمانی نوین برای اولین بار در ایران با گواهی هیأت ممتحنه رشته مربوطه و تأیید هیأت ممیزه مرکزی وزارت بهداشت | |
| | ٥/+ تا ٥ | ۲. تنوین راهنمای طبابت بالینی کشور، با استناد بر پژوهش و براساس شواهد علمی با گواهی معاونت درمان وزارت بهداشت و تأیید هیأت ممیزه مرکزی وزارت بهداشت | |
| از بندهای ۷ تا ۱۰ ۳۰ امتیاز | ه/+ تا ه | ۷. اختراع، اکتشاف و تولید محصولات پژوهشی کاربردی ثبت شده در مراجع قانونی داخل کشور با تأیید معاونت پژوهش و فناوری وزارتین | |
| | 1051 | ۸ اختراع، اکتشاف و تولید محصولات پژوهشی کاربردی ثبت شده و دستاوردهای فناورانه که در چارچوب پژوهشیهای کاربردی، پایان نامهها و رسالههای دکتری با رعایت مالکیت فکری در قالب شرکت دانش بنیان یا شرکت دانشگاهی مراحل تجاری سازی آن با تأیید مراجع مربوطه به انجام رسیده باشد. تبصره: چنانچه ثبت اختراع در سطح بین المللی، به همراه چاپ مقالهای باشد به یک مورد امتیاز کامل و به مورد دیگر نصف امتیاز تعلق می گیرد. | |
| | ۹. ثبت مالکیت فکری (پتنت) دستاوردهای پژوهشی و فناوری به نام مؤسسه محل خدمت متقاضی، به شرط تا ۱۵ این که به فروش رسیده و یا به صورت تحت لیسانس از آن استفاده شده باشد. | | |
| | ٠/٠ تا ٢ | ۱۰. تعیین توالی ژن ثبت شده در مراجع قانونی یا علمی داخل یا خارج از کشور با تأیید مراجع ذی صلاح وزارتین | |



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