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План

- Что такое Mendeley?
- Создание своей библиотеки
- Управление статьями и информацией о них
- Вставка цитат и создание библиографий
- Обмен информацией через группы

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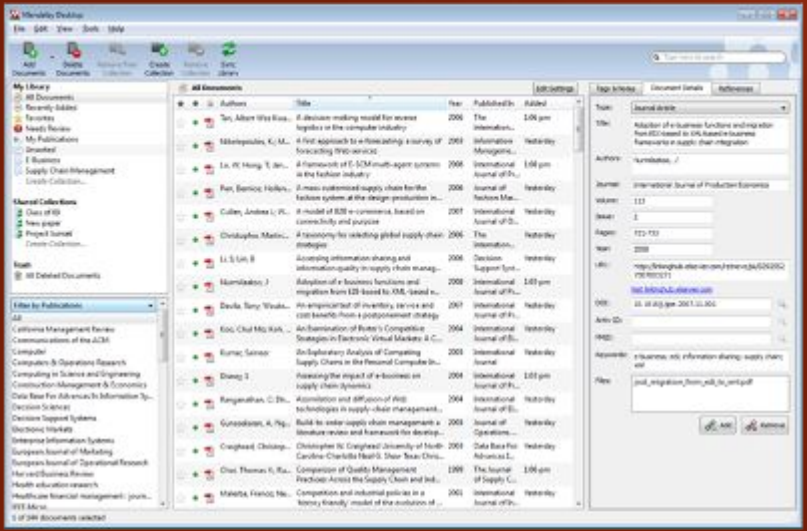

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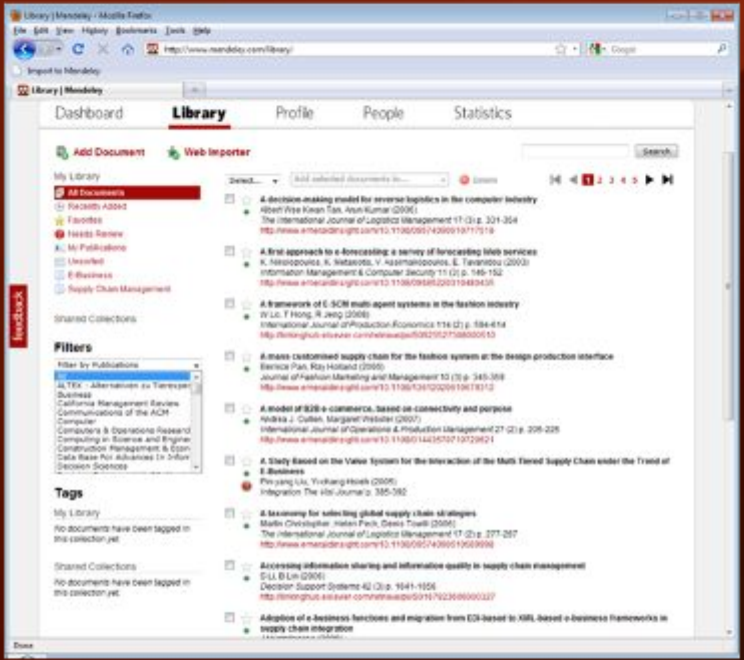
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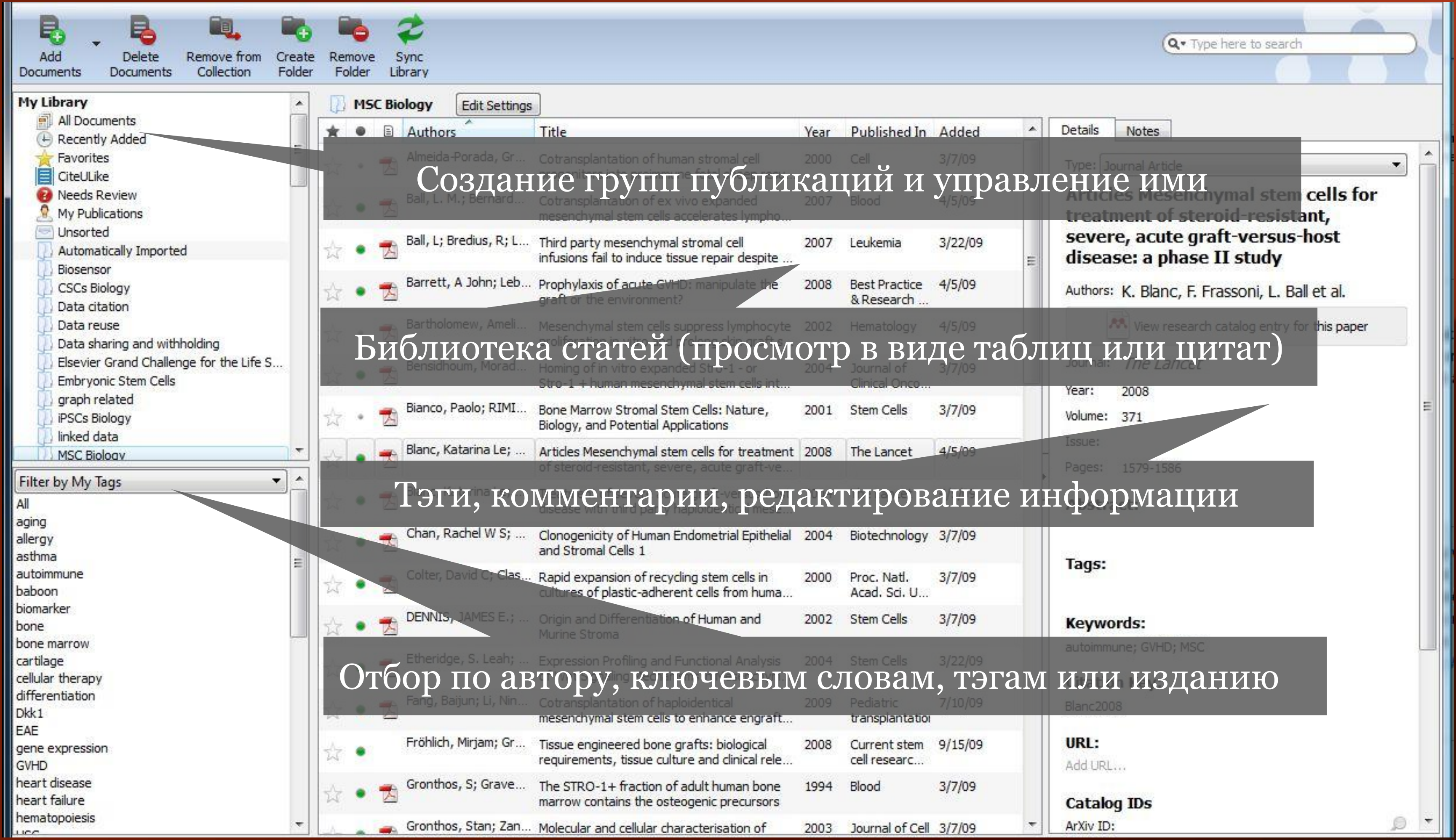
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Тэги, комментарии, редактирование информации

Отбор по автору, ключевым словам, тэгам или изданию

Authors	Title	Year	Published In	Added
Almeida-Porada, Gr...	Cotransplantation of human stromal cell	2000	Cell	3/7/09
Ball, L. M.; Bernard...	Cotransplantation of ex vivo expanded mesenchymal stem cells accelerates lympho...	2007	Blood	4/5/09
Ball, L; Bredius, R; L...	Third party mesenchymal stromal cell infusions fail to induce tissue repair despite ...	2007	Leukemia	3/22/09
Barrett, A John; Leb...	Prophylaxis of acute GVHD: manipulate the graft or the environment?	2008	Best Practice & Research ...	4/5/09
Bartholomew, Ameli...	Mesenchymal stem cells suppress lymphocyte	2002	Hematology	4/5/09
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Bianco, Paolo; RIMI...	Bone Marrow Stromal Stem Cells: Nature, Biology, and Potential Applications	2001	Stem Cells	3/7/09
Blanc, Katarina Le; ...	Articles Mesenchymal stem cells for treatment of steroid-resistant, severe, acute graft-ve...	2008	The Lancet	4/5/09
Chan, Rachel W S; ...	Clonogenicity of Human Endometrial Epithelial and Stromal Cells 1	2004	Biotechnology	3/7/09
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Fröhlich, Mirjam; Gr...	Tissue engineered bone grafts: biological requirements, tissue culture and clinical rele...	2008	Current stem cell researc...	9/15/09
Gronthos, S; Grave...	The STRO-1+ fraction of adult human bone marrow contains the osteogenic precursors	1994	Blood	3/7/09
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Articles Mesenchymal stem cells for treatment of steroid-resistant, severe, acute graft-versus-host disease: a phase II study

Authors: K. Blanc, F. Frassoni, L. Ball et al.

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Journal: The Lancet

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Volume: 371

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Tags:

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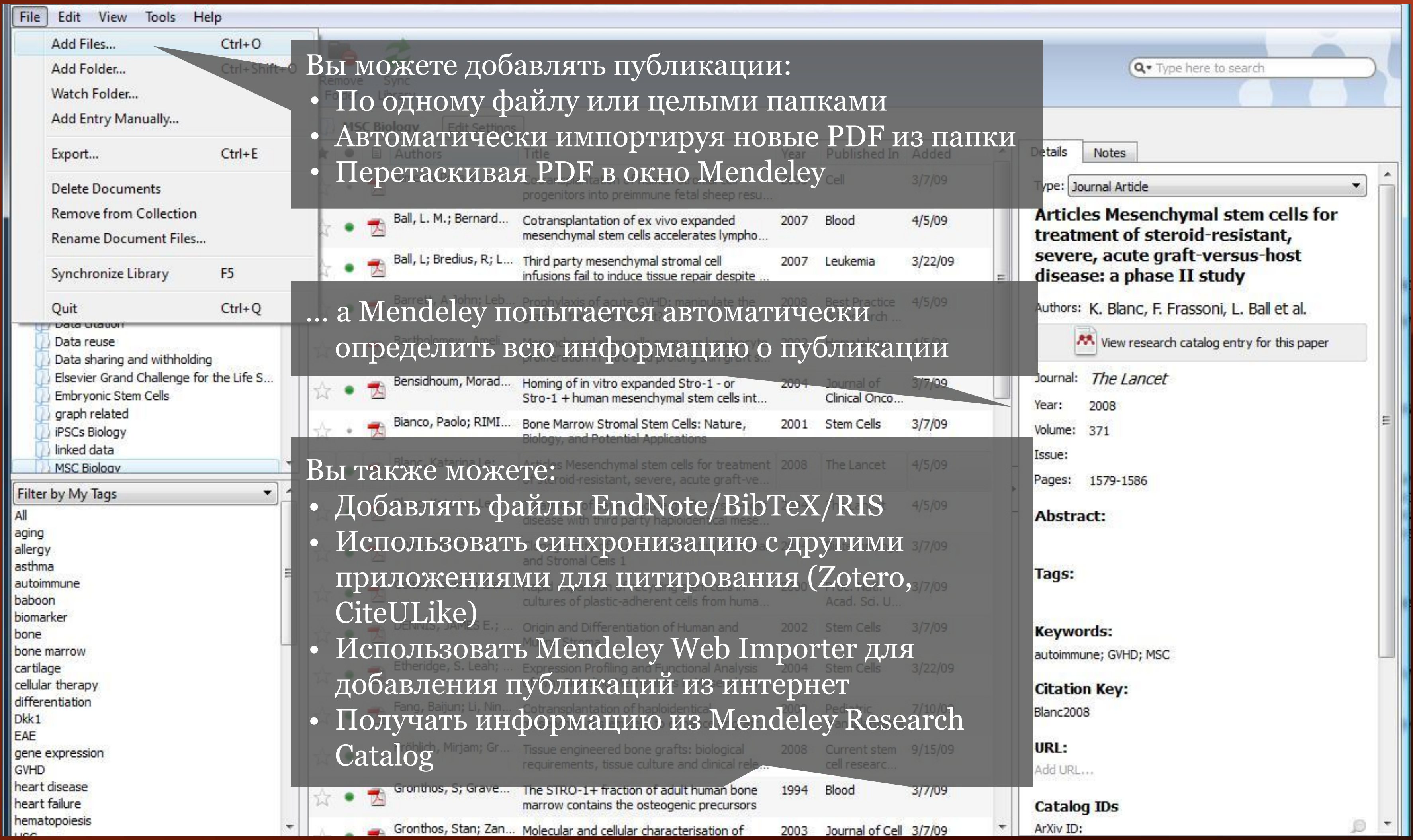
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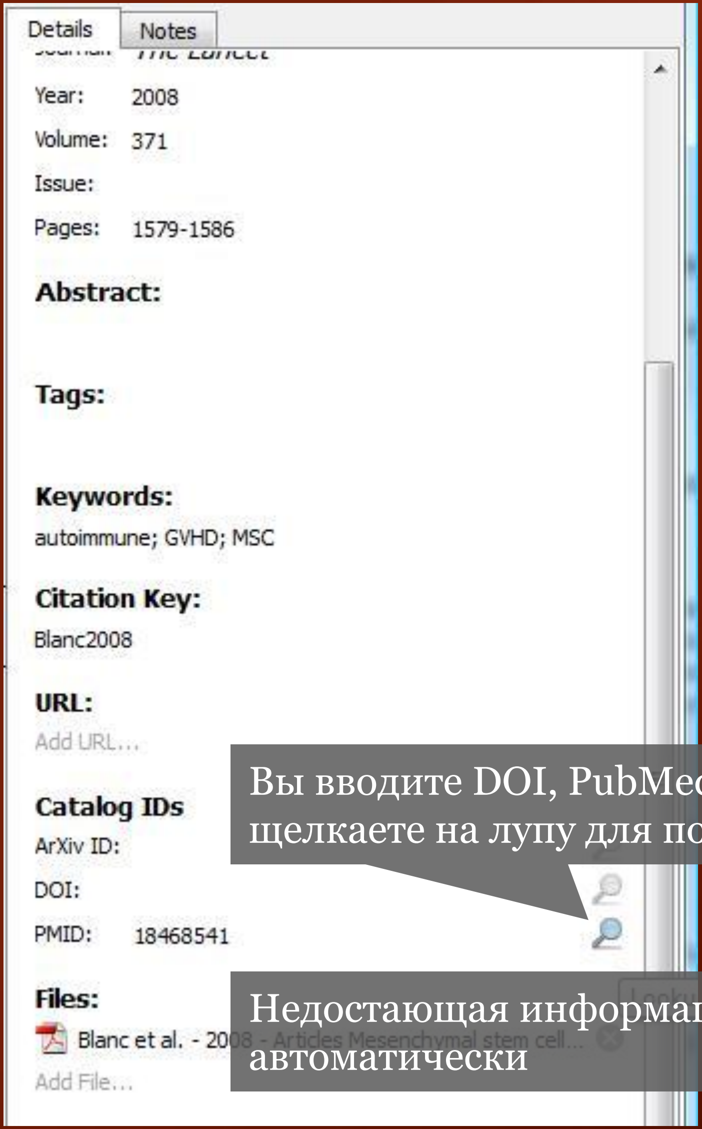

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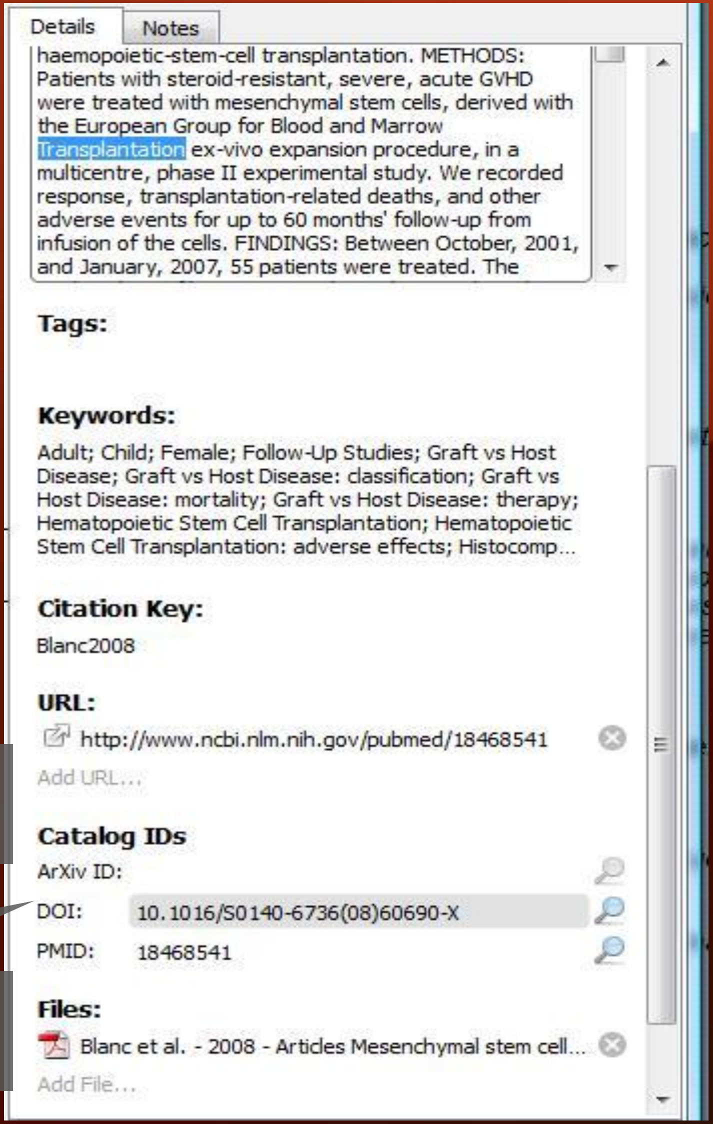

haemopoietic-stem-cell transplantation. METHODS: Patients with steroid-resistant, severe, acute GVHD were treated with mesenchymal stem cells, derived with the European Group for Blood and Marrow Transplantation ex-vivo expansion procedure, in a multicentre, phase II experimental study. We recorded response, transplantation-related deaths, and other adverse events for up to 60 months' follow-up from infusion of the cells. FINDINGS: Between October, 2001, and January, 2007, 55 patients were treated. The

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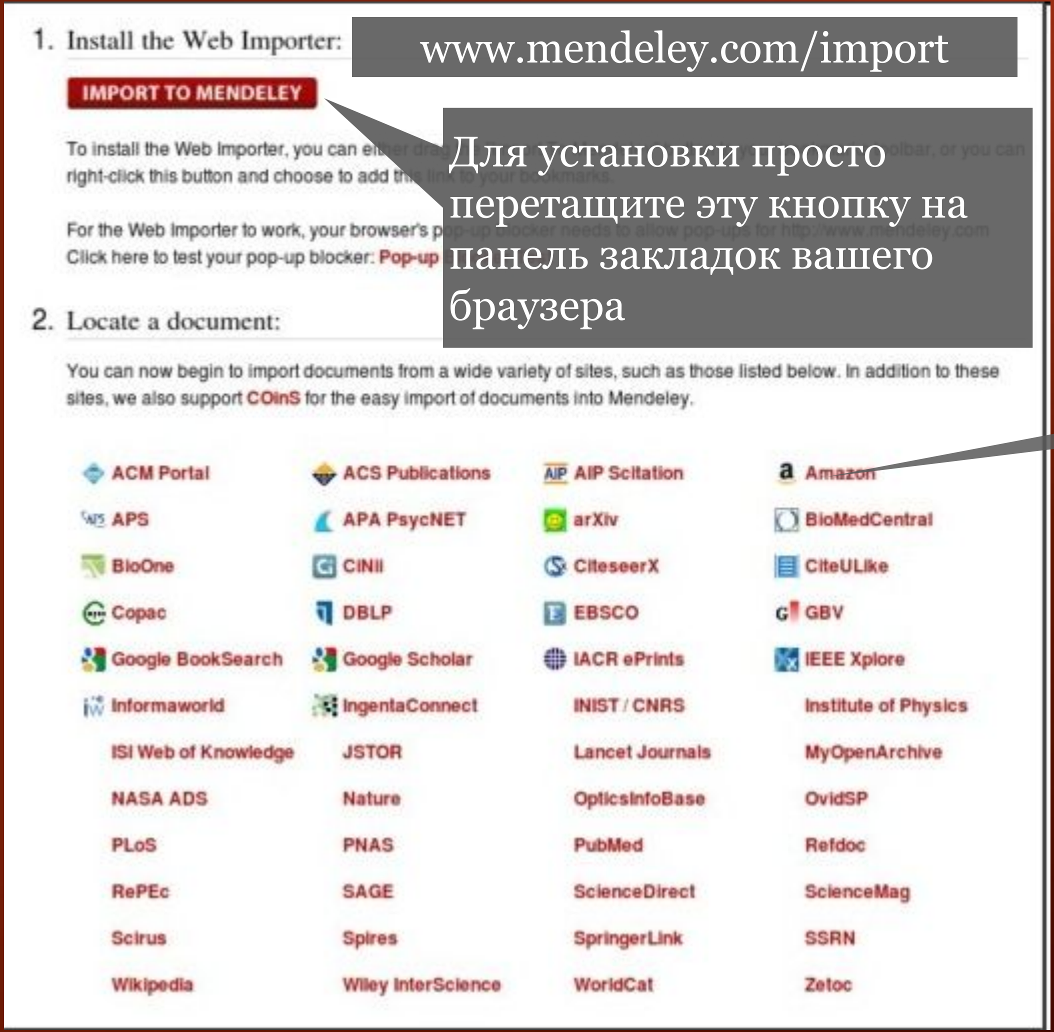

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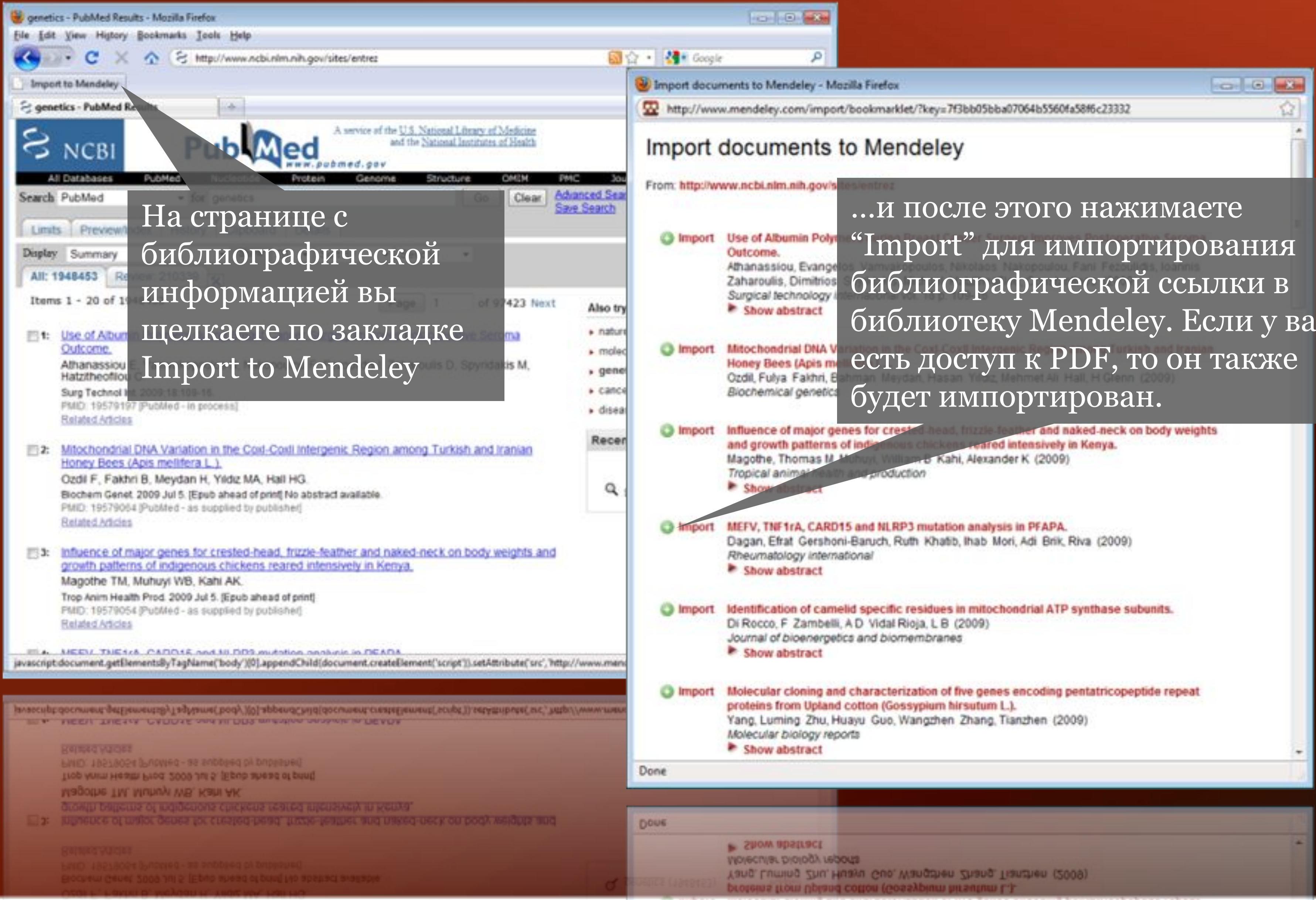

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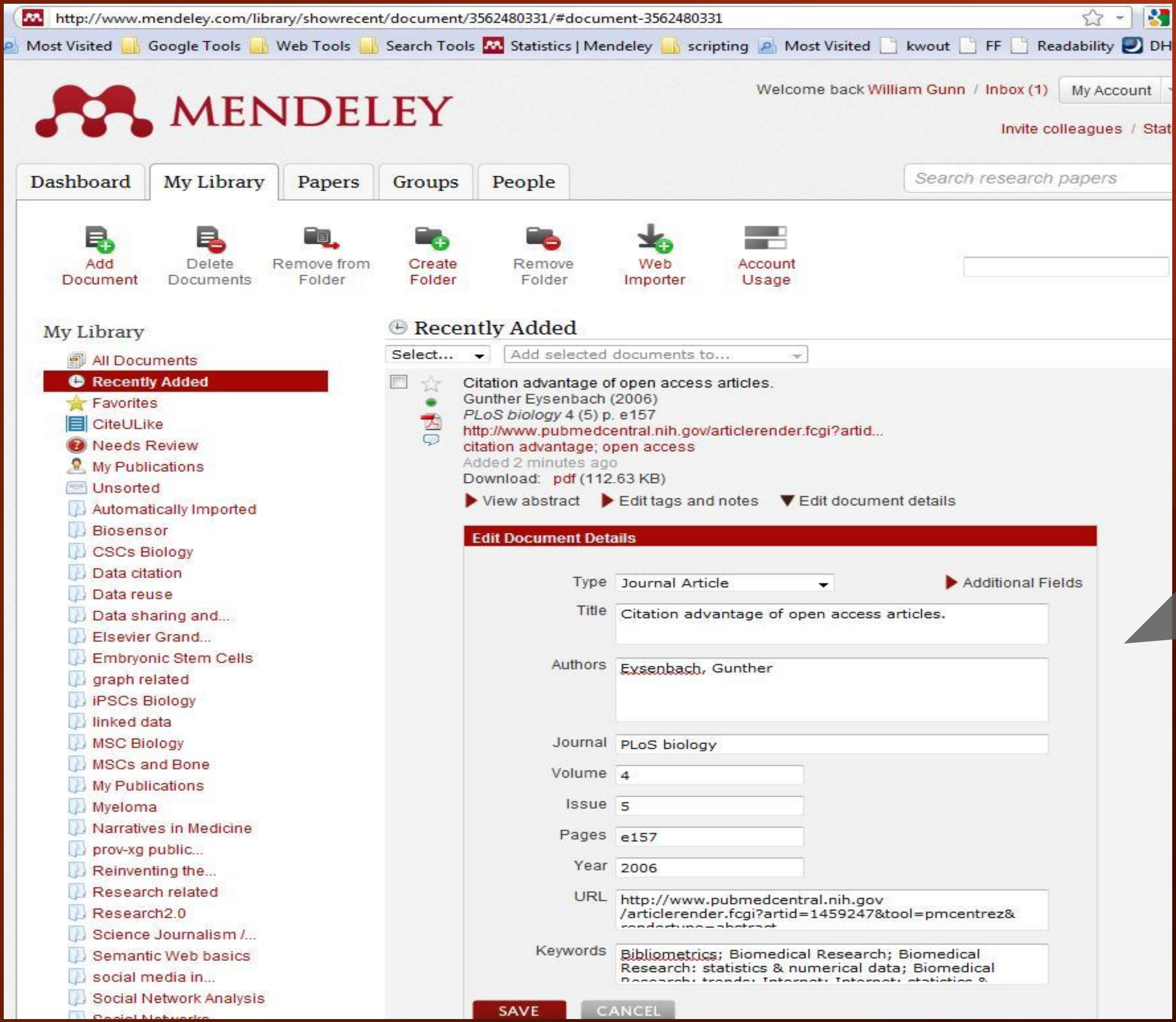
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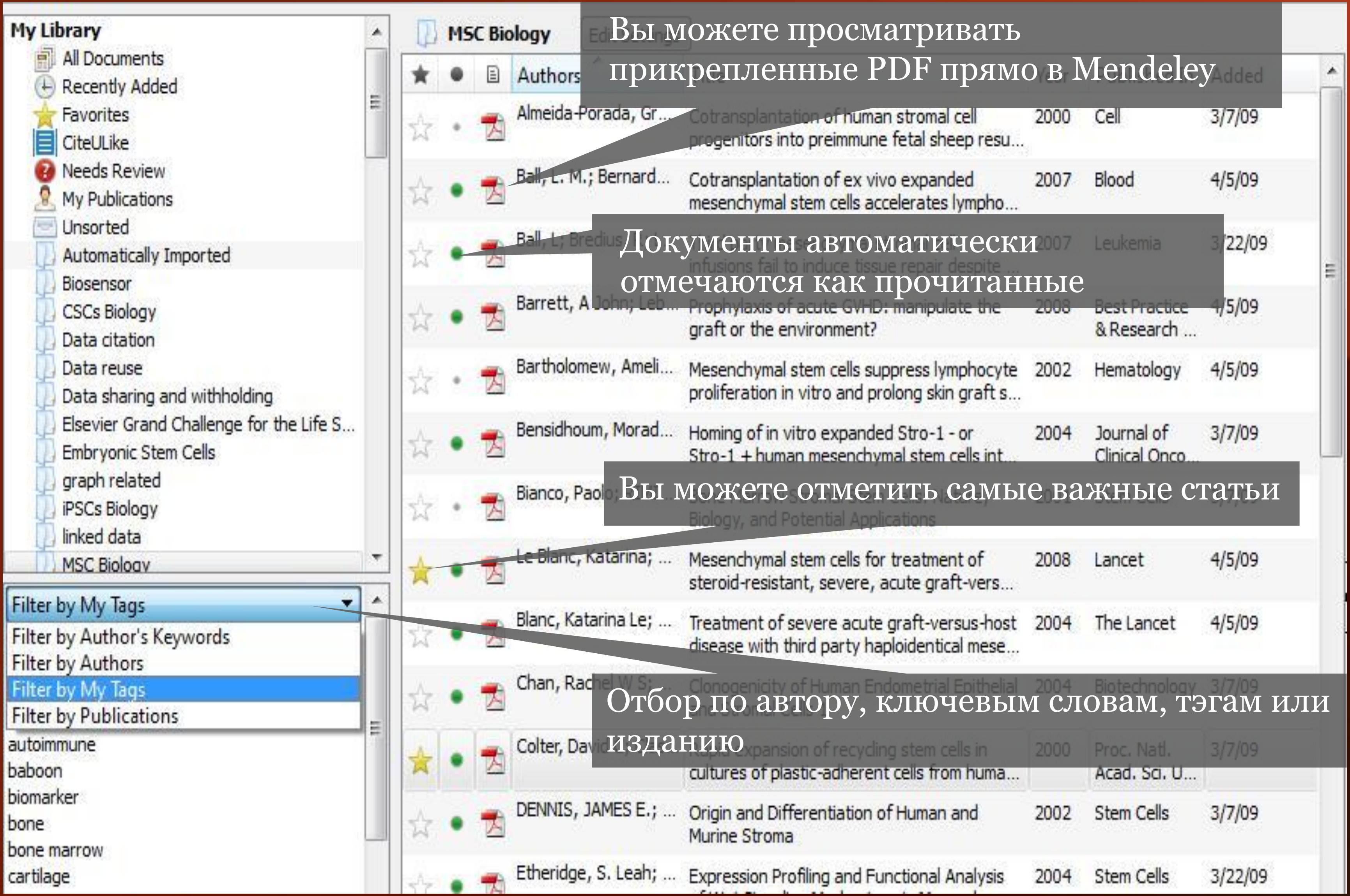
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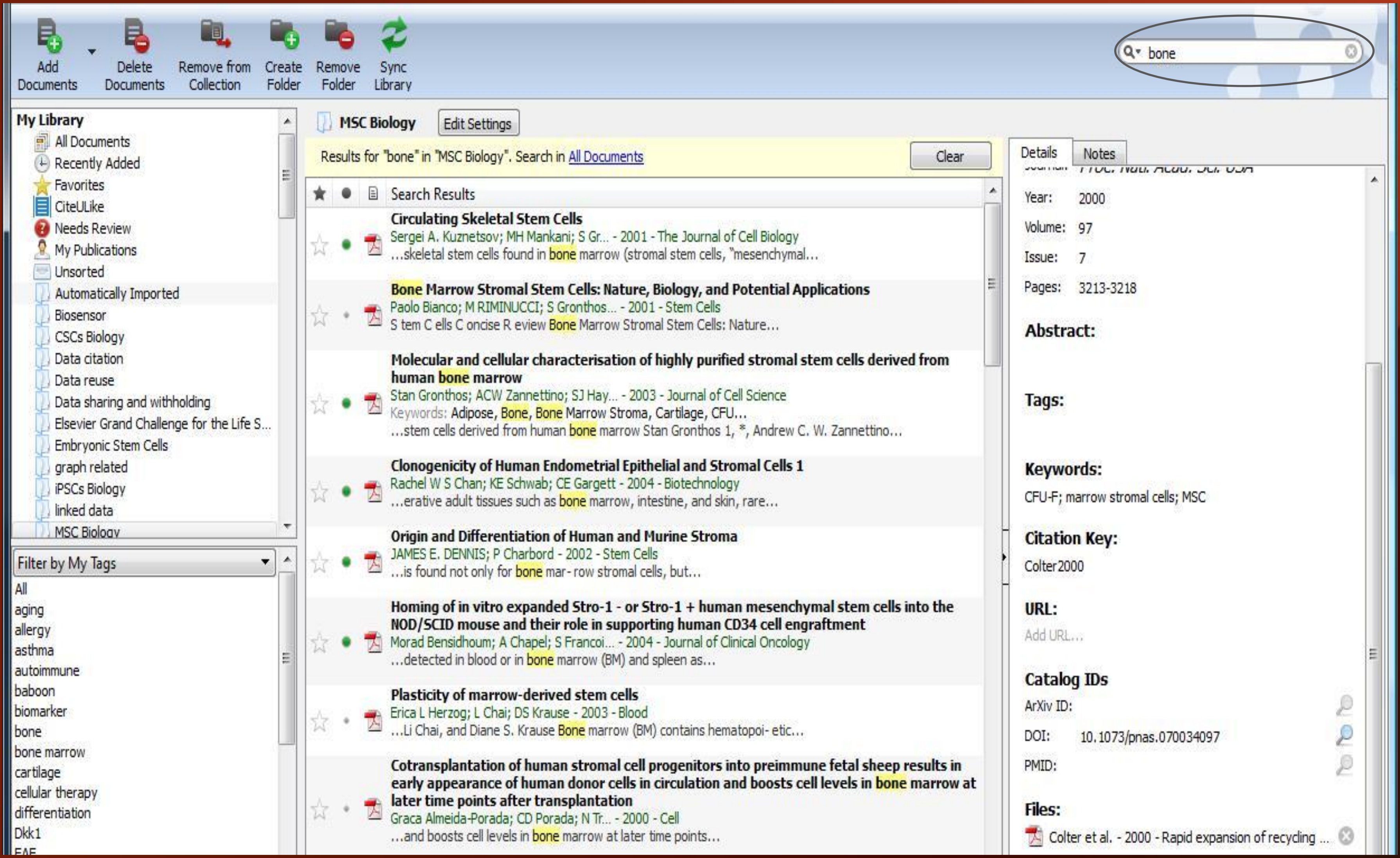
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Cotransplantation of human stromal cell progenitors into preimmune fetal sheep results in early appearance of human donor cells in circulation and boosts cell levels in bone marrow at later time points after transplantation
Graca Almeida-Porada; CD Porada; N Tr... - 2000 - Cell
...and boosts cell levels in bone marrow at later time points...

Journal: Proc. Natl. Acad. Sci. USA
Year: 2000
Volume: 97
Issue: 7
Pages: 3213-3218

Abstract:

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Keywords:
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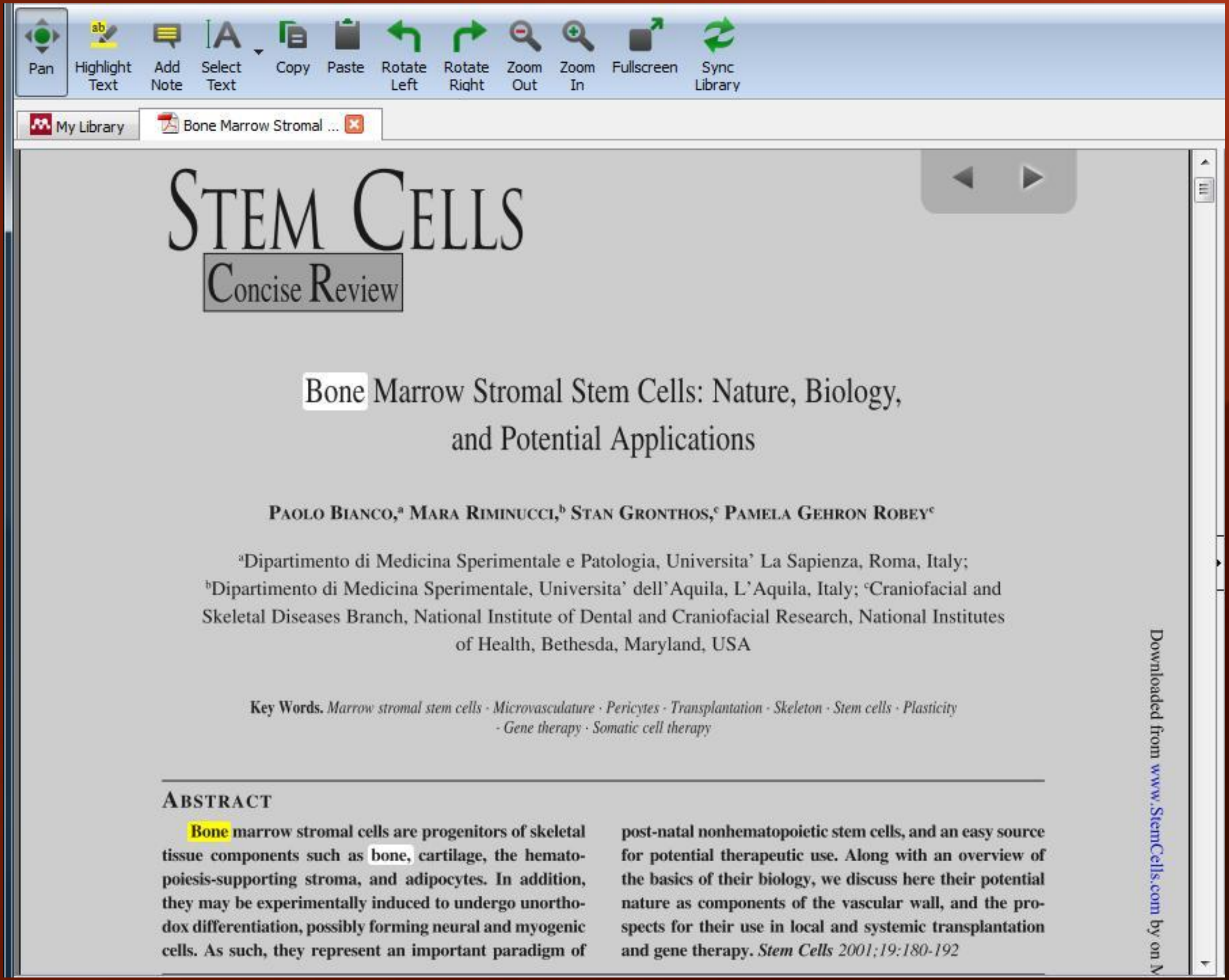

STEM CELLS

Concise Review

Bone Marrow Stromal Stem Cells: Nature, Biology, and Potential Applications

PAOLO BIANCO,[a] MARA RIMINUCCI,[b] STAN GRONTHOS,[c] PAMELA GEHRON ROBEY[c]

[a]Dipartimento di Medicina Sperimentale e Patologia, Universita' La Sapienza, Roma, Italy; [b]Dipartimento di Medicina Sperimentale, Universita' dell'Aquila, L'Aquila, Italy; [c]Craniofacial and Skeletal Diseases Branch, National Institute of Dental and Craniofacial Research, National Institutes of Health, Bethesda, Maryland, USA

Key Words. *Marrow stromal stem cells · Microvasculature · Pericytes · Transplantation · Skeleton · Stem cells · Plasticity · Gene therapy · Somatic cell therapy*

ABSTRACT

Bone marrow stromal cells are progenitors of skeletal tissue components such as bone, cartilage, the hematopoiesis-supporting stroma, and adipocytes. In addition, they may be experimentally induced to undergo unorthodox differentiation, possibly forming neural and myogenic cells. As such, they represent an important paradigm of post-natal nonhematopoietic stem cells, and an easy source for potential therapeutic use. Along with an overview of the basics of their biology, we discuss here their potential nature as components of the vascular wall, and the prospects for their use in local and systemic transplantation and gene therapy. *Stem Cells* *2001;19:180-192*

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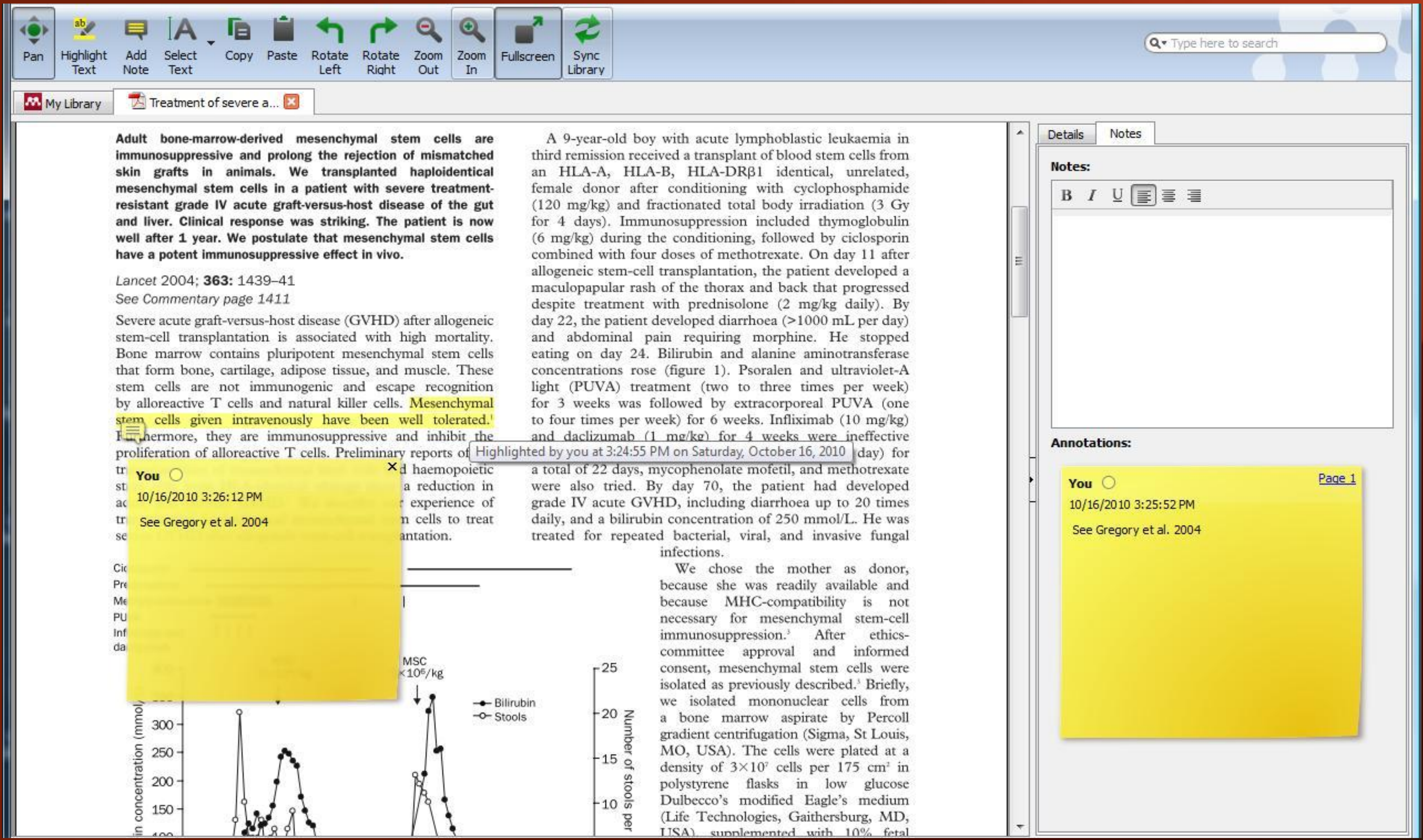

Adult bone-marrow-derived mesenchymal stem cells are immunosuppressive and prolong the rejection of mismatched skin grafts in animals. We transplanted haploidentical mesenchymal stem cells in a patient with severe treatment-resistant grade IV acute graft-versus-host disease of the gut and liver. Clinical response was striking. The patient is now well after 1 year. We postulate that mesenchymal stem cells have a potent immunosuppressive effect in vivo.

Lancet 2004; **363:** 1439–41

See Commentary page 1411

Severe acute graft-versus-host disease (GVHD) after allogeneic stem-cell transplantation is associated with high mortality. Bone marrow contains pluripotent mesenchymal stem cells that form bone, cartilage, adipose tissue, and muscle. These stem cells are not immunogenic and escape recognition by alloreactive T cells and natural killer cells. Mesenchymal stem cells given intravenously have been well tolerated.[1] Furthermore, they are immunosuppressive and inhibit the proliferation of alloreactive T cells. Preliminary reports of

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See Gregory et al. 2004

A 9-year-old boy with acute lymphoblastic leukaemia in third remission received a transplant of blood stem cells from an HLA-A, HLA-B, HLA-DRβ1 identical, unrelated, female donor after conditioning with cyclophosphamide (120 mg/kg) and fractionated total body irradiation (3 Gy for 4 days). Immunosuppression included thymoglobulin (6 mg/kg) during the conditioning, followed by ciclosporin combined with four doses of methotrexate. On day 11 after allogeneic stem-cell transplantation, the patient developed a maculopapular rash of the thorax and back that progressed despite treatment with prednisolone (2 mg/kg daily). By day 22, the patient developed diarrhoea (>1000 mL per day) and abdominal pain requiring morphine. He stopped eating on day 24. Bilirubin and alanine aminotransferase concentrations rose (figure 1). Psoralen and ultraviolet-A light (PUVA) treatment (two to three times per week) for 3 weeks was followed by extracorporeal PUVA (one to four times per week) for 6 weeks. Infliximab (10 mg/kg) and daclizumab (1 mg/kg) for 4 weeks were ineffective day) for a total of 22 days, mycophenolate mofetil, and methotrexate were also tried. By day 70, the patient had developed grade IV acute GVHD, including diarrhoea up to 20 times daily, and a bilirubin concentration of 250 mmol/L. He was treated for repeated bacterial, viral, and invasive fungal infections.

We chose the mother as donor, because she was readily available and because MHC-compatibility is not necessary for mesenchymal stem-cell immunosuppression.[3] After ethics-committee approval and informed consent, mesenchymal stem cells were isolated as previously described.[3] Briefly, we isolated mononuclear cells from a bone marrow aspirate by Percoll gradient centrifugation (Sigma, St Louis, MO, USA). The cells were plated at a density of 3×10^7 cells per 175 cm^2 in polystyrene flasks in low glucose Dulbecco's modified Eagle's medium (Life Technologies, Gaithersburg, MD, USA) supplemented with 10% fetal

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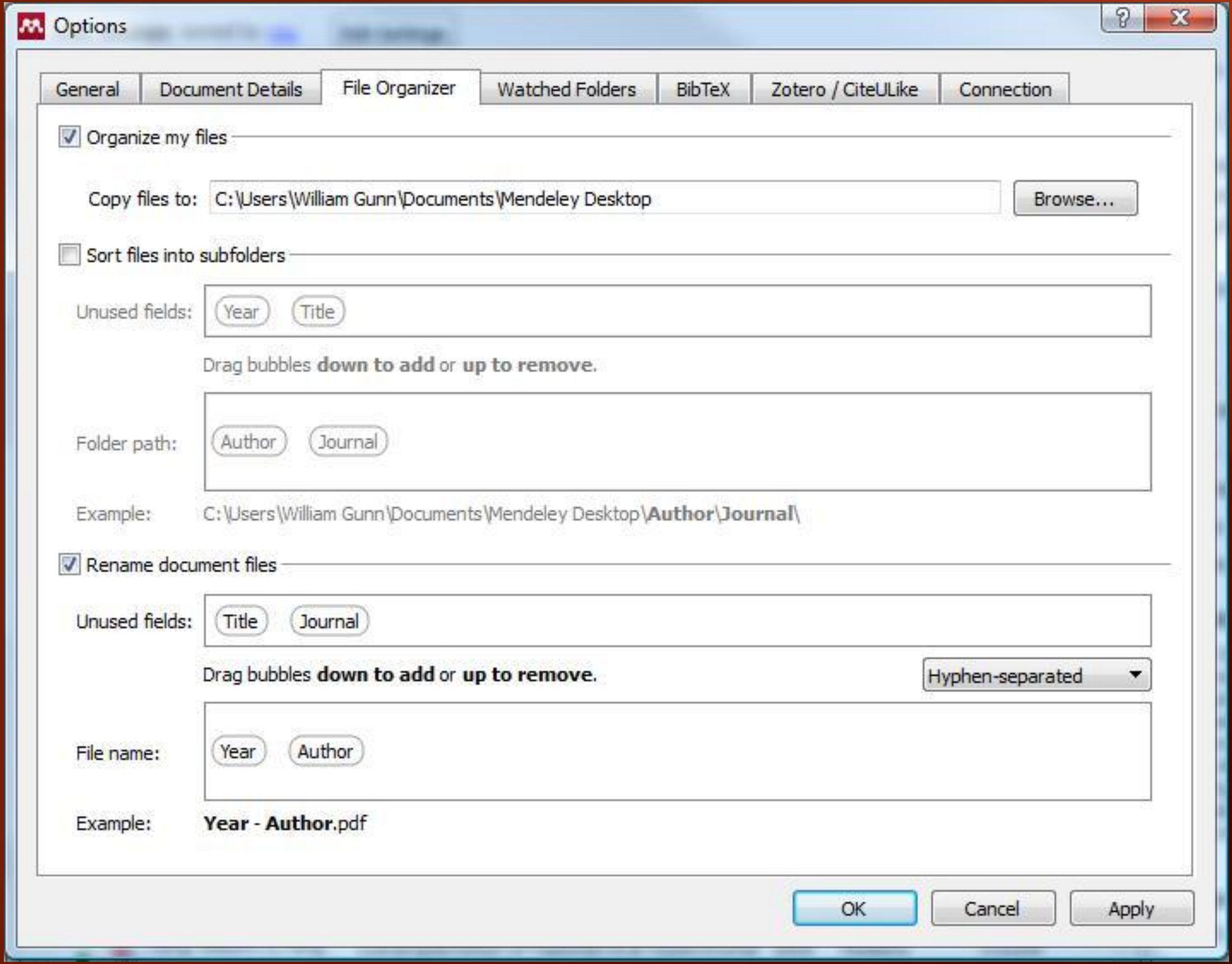
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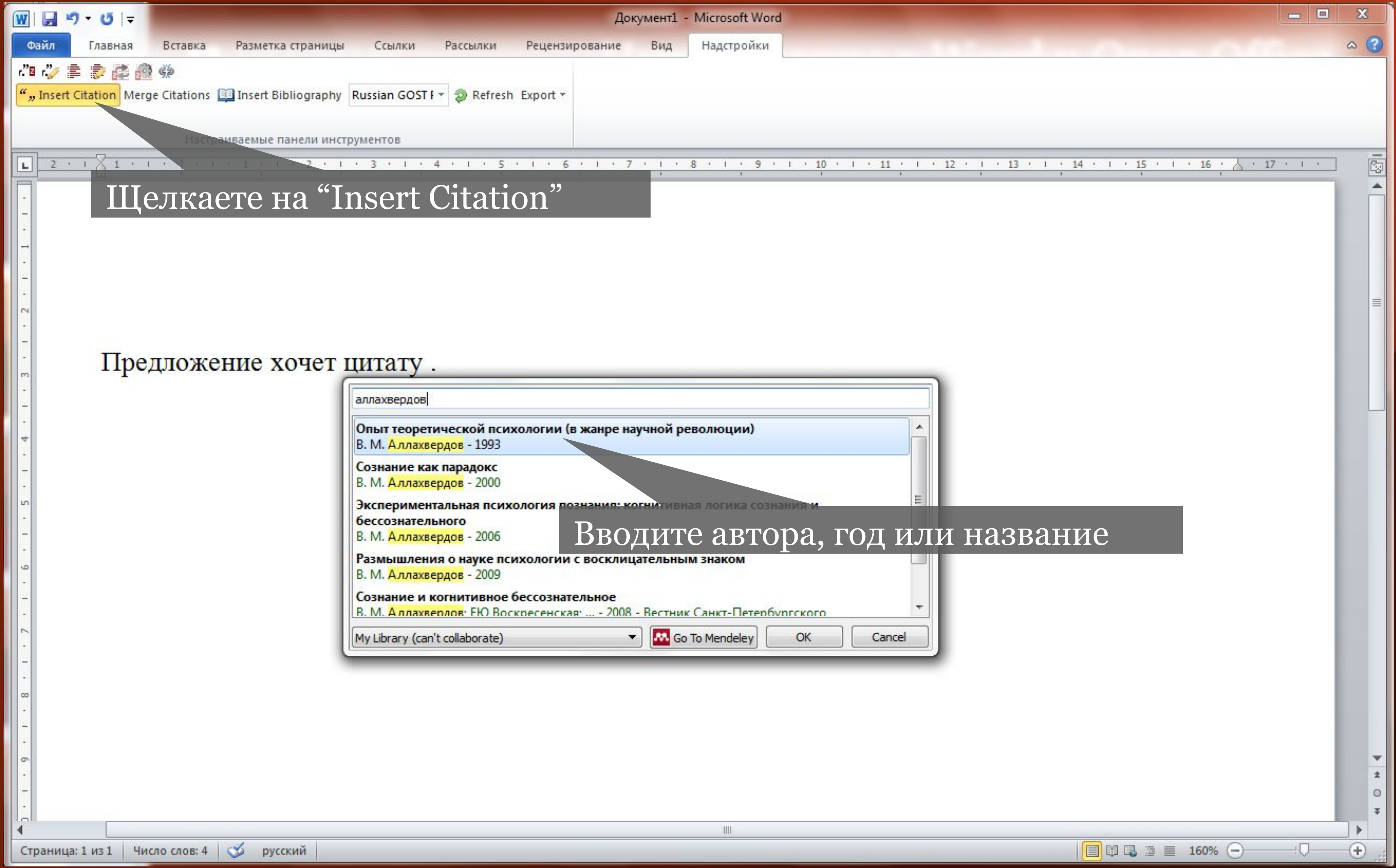
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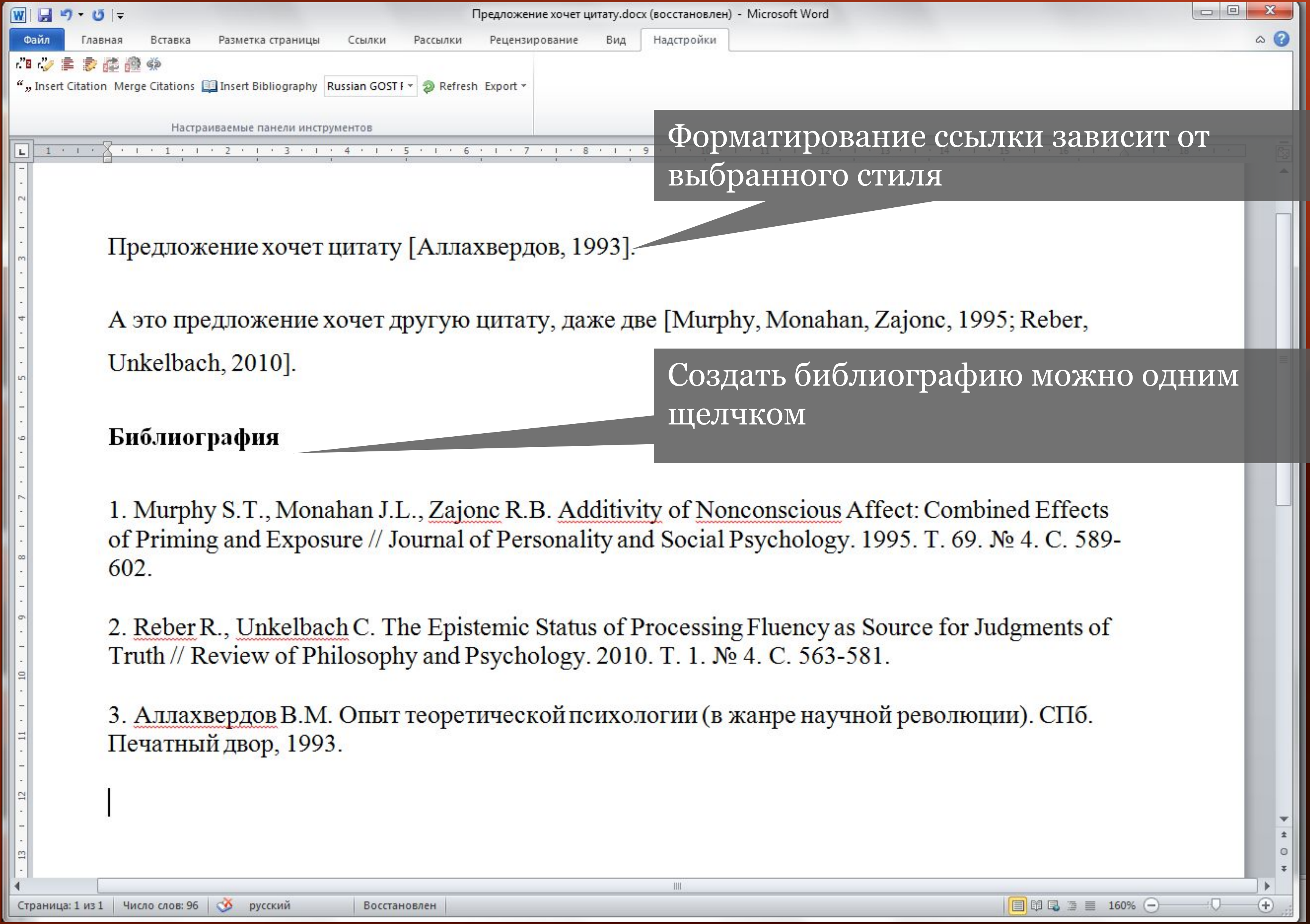
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Библиография

1. Murphy S.T., Monahan J.L., Zajonc R.B. Additivity of Nonconscious Affect: Combined Effects of Priming and Exposure // Journal of Personality and Social Psychology. 1995. T. 69. № 4. C. 589-602.

2. Reber R., Unkelbach C. The Epistemic Status of Processing Fluency as Source for Judgments of Truth // Review of Philosophy and Psychology. 2010. T. 1. № 4. C. 563-581.

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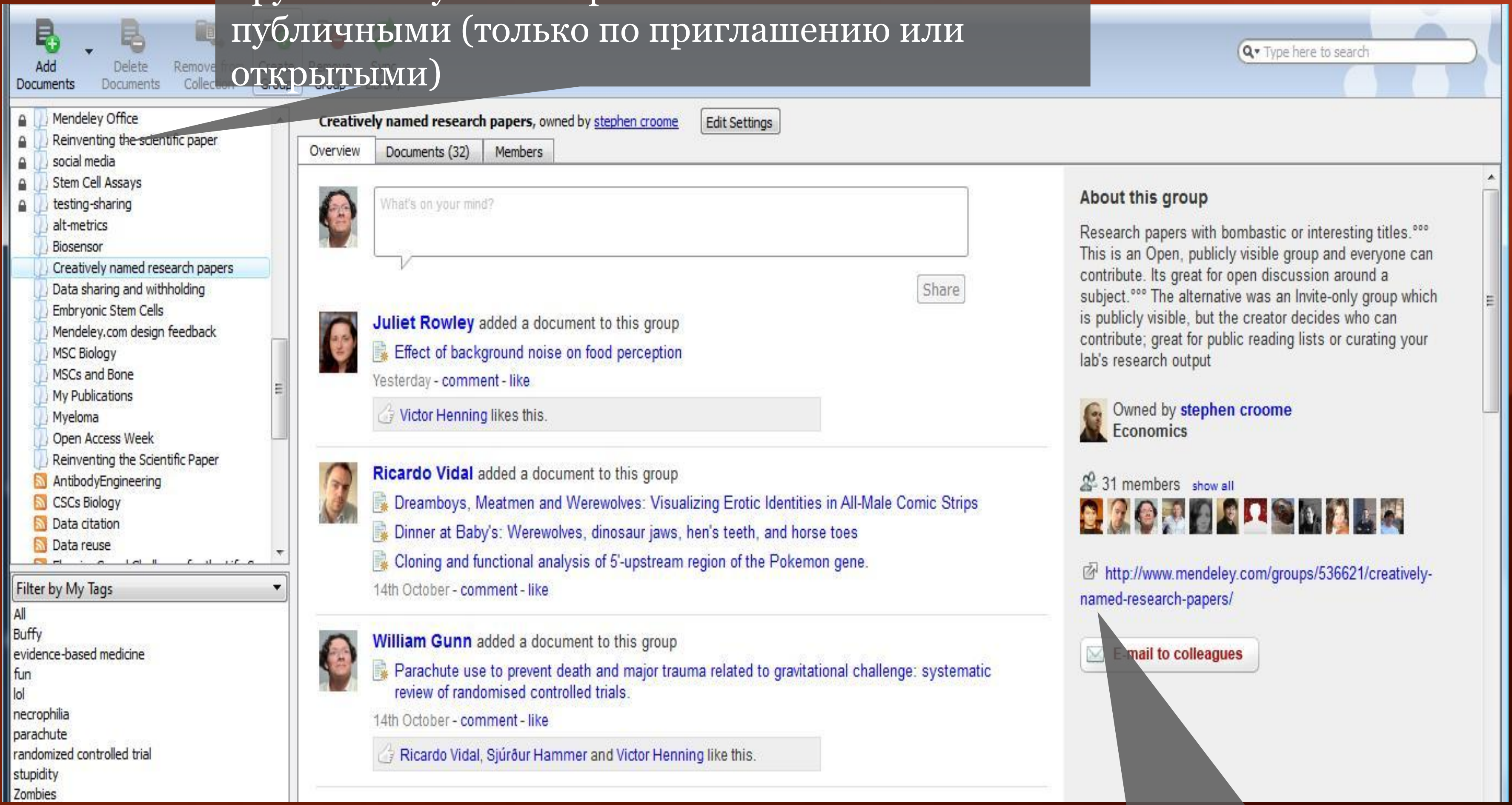
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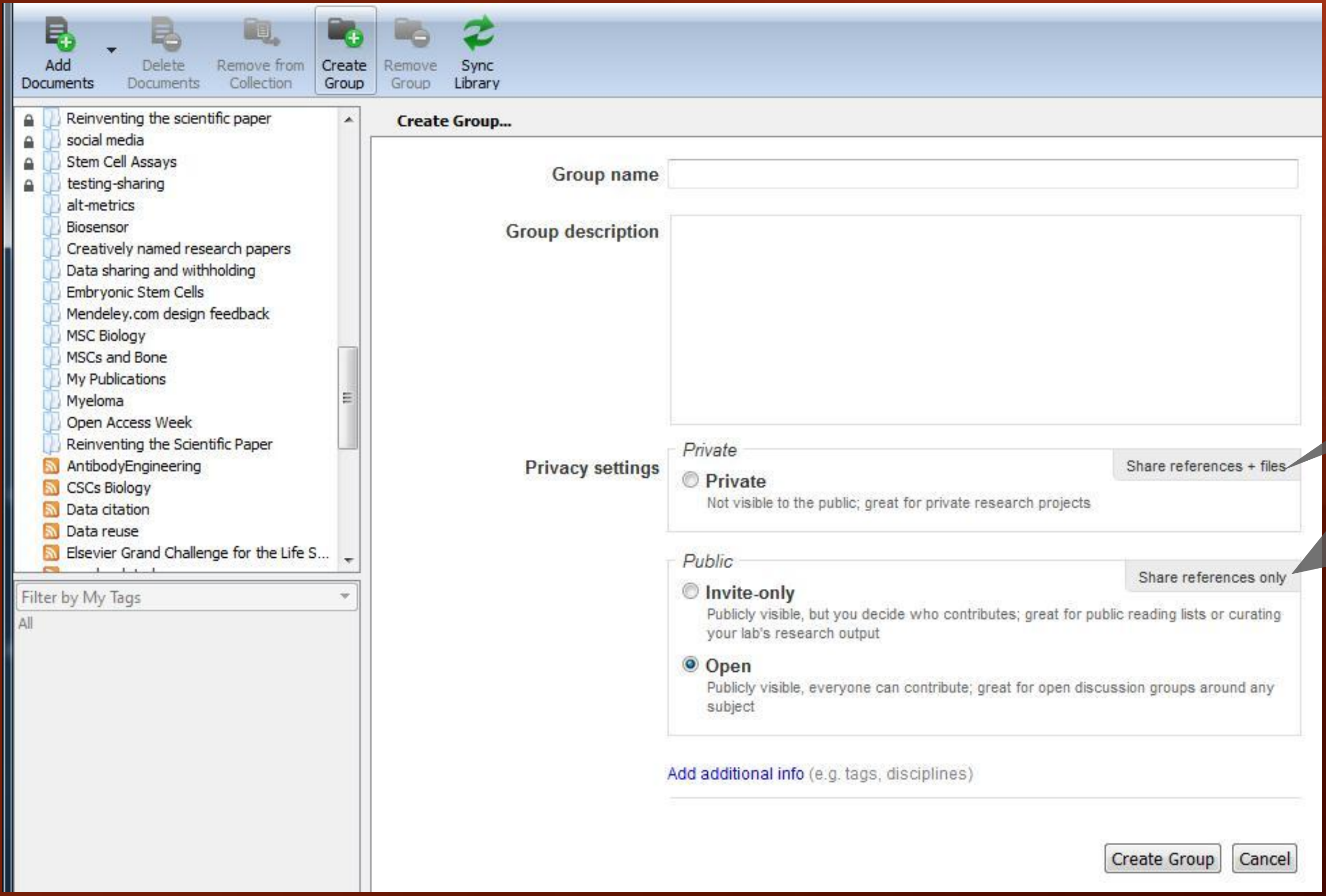
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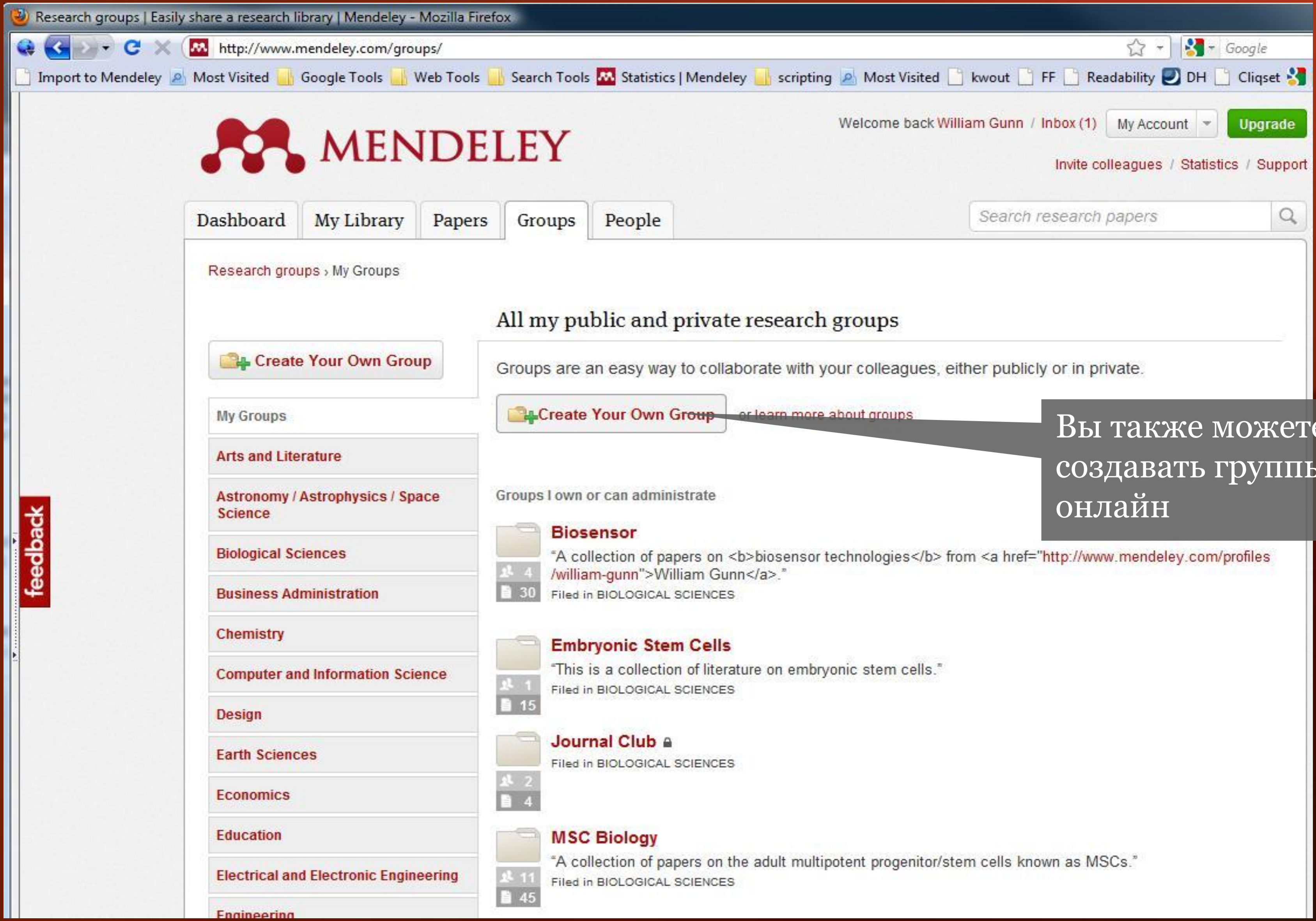
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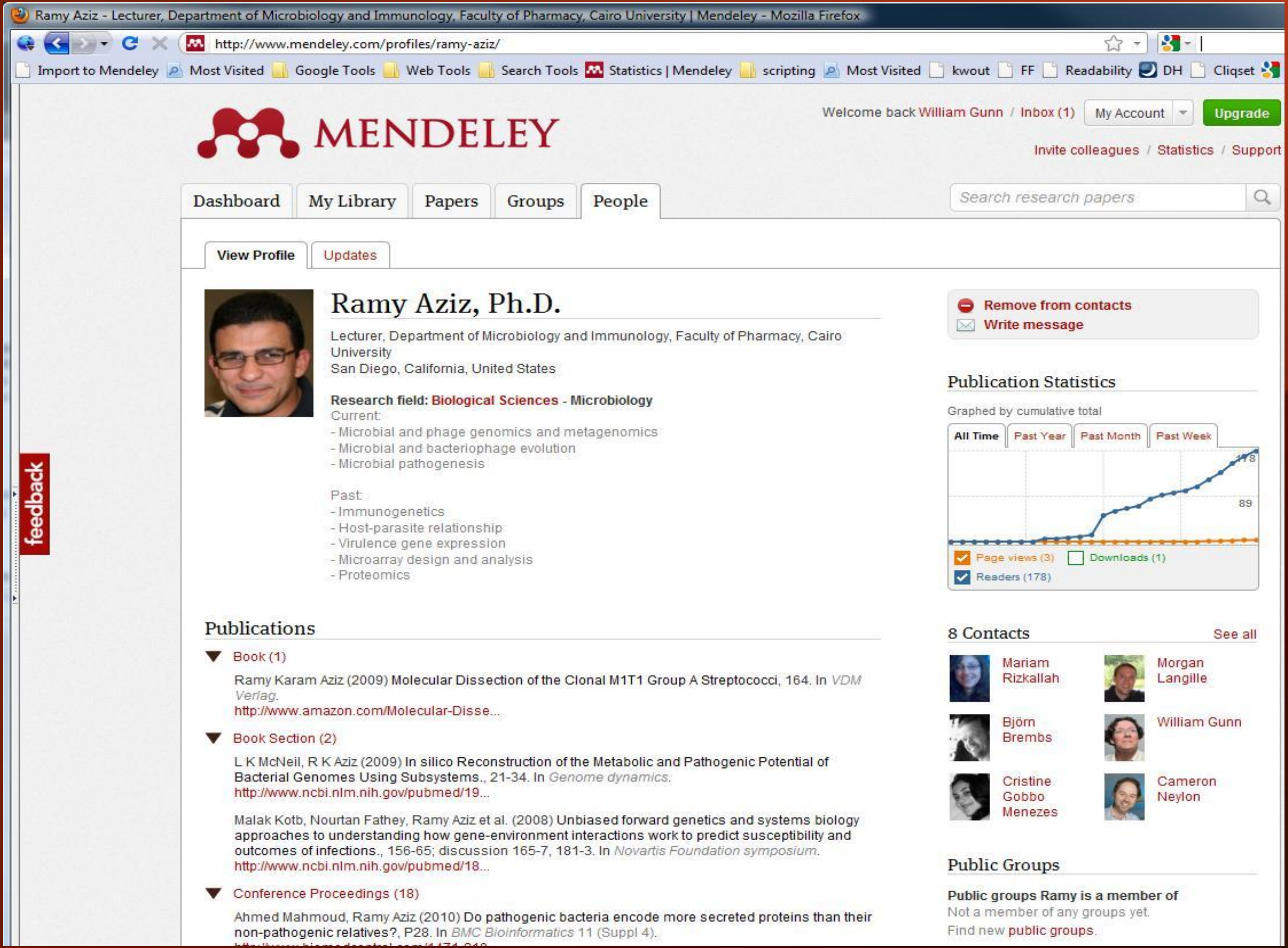
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L K McNeil, R K Aziz (2009) In silico Reconstruction of the Metabolic and Pathogenic Potential of Bacterial Genomes Using Subsystems., 21-34. In *Genome dynamics*.
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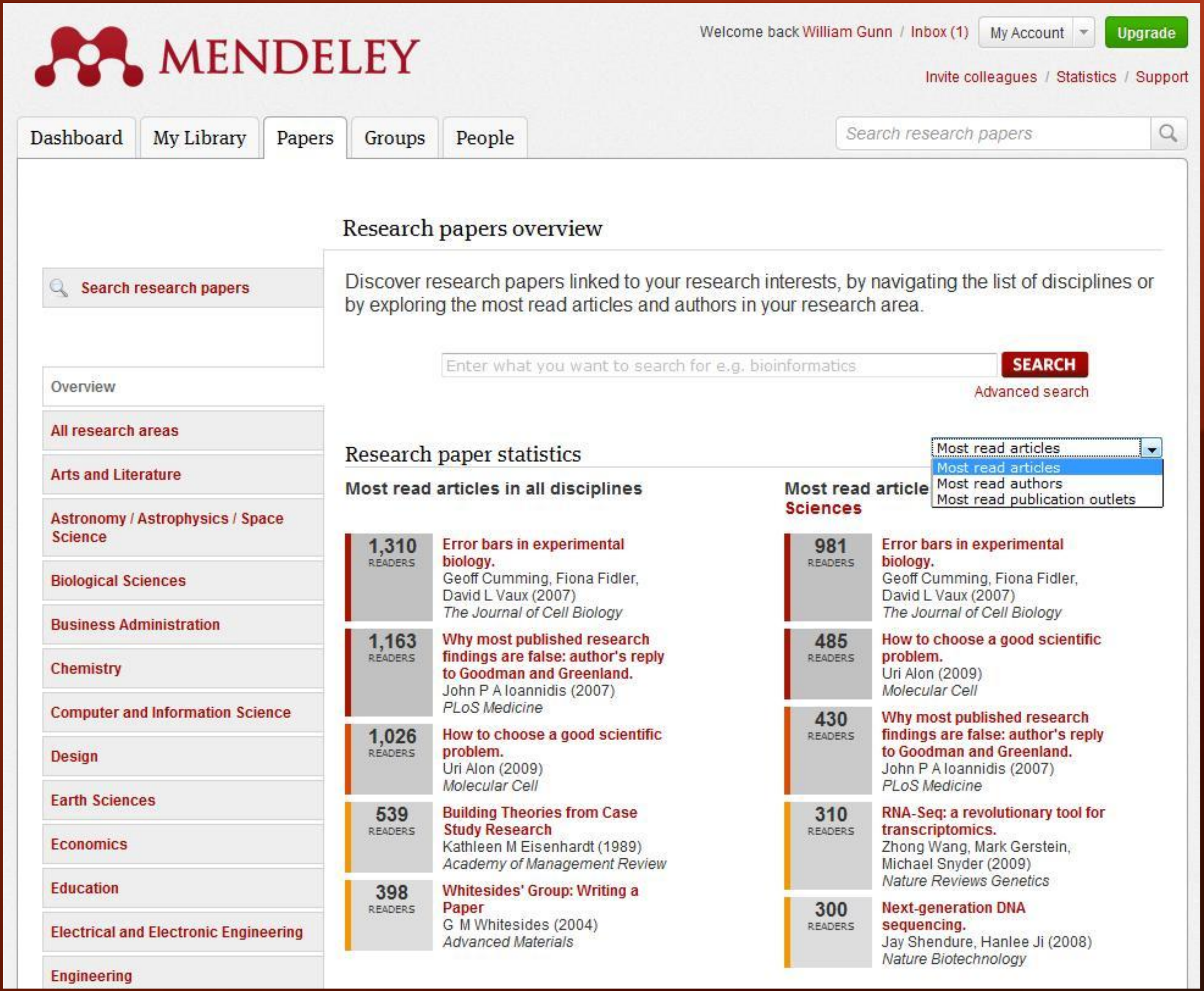
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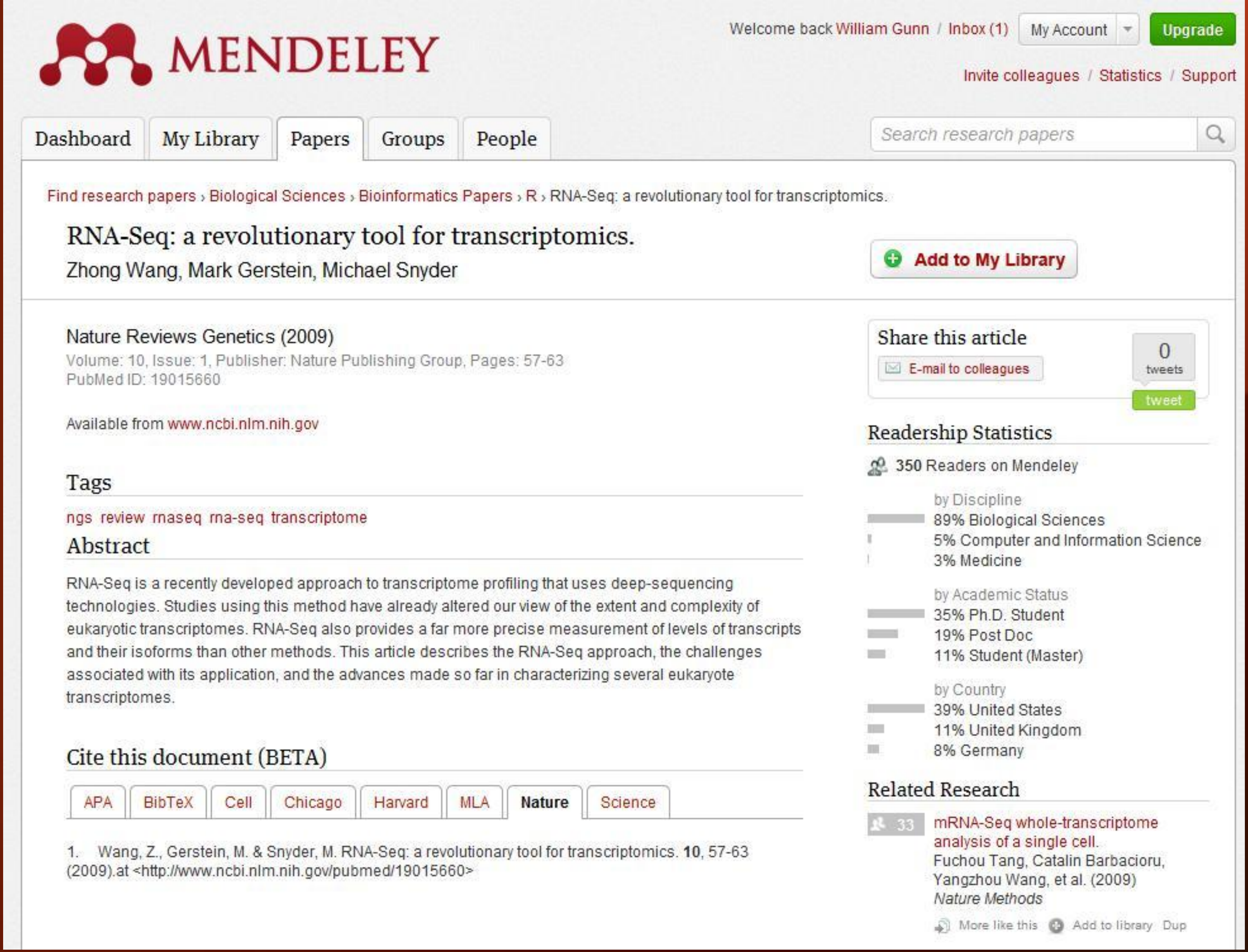
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Abstract

RNA-Seq is a recently developed approach to transcriptome profiling that uses deep-sequencing technologies. Studies using this method have already altered our view of the extent and complexity of eukaryotic transcriptomes. RNA-Seq also provides a far more precise measurement of levels of transcripts and their isoforms than other methods. This article describes the RNA-Seq approach, the challenges associated with its application, and the advances made so far in characterizing several eukaryote transcriptomes.

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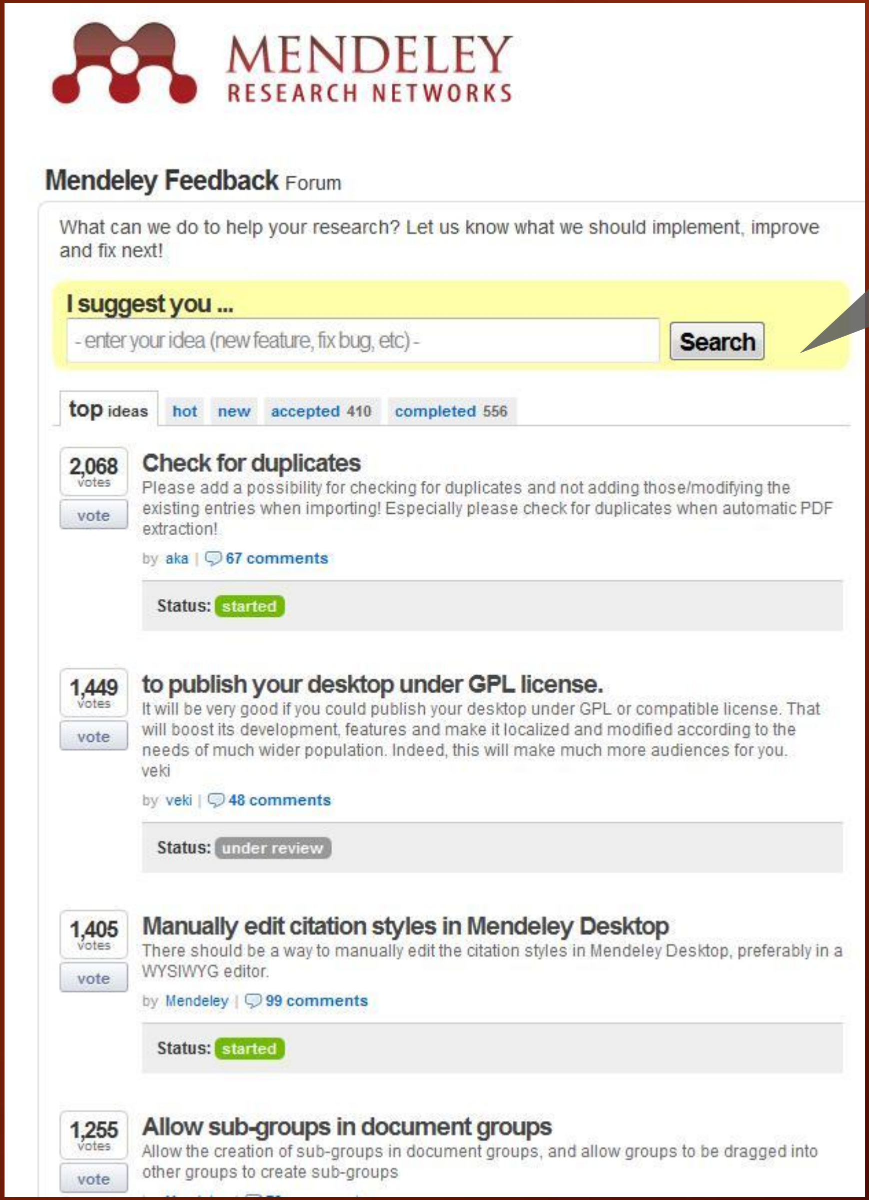

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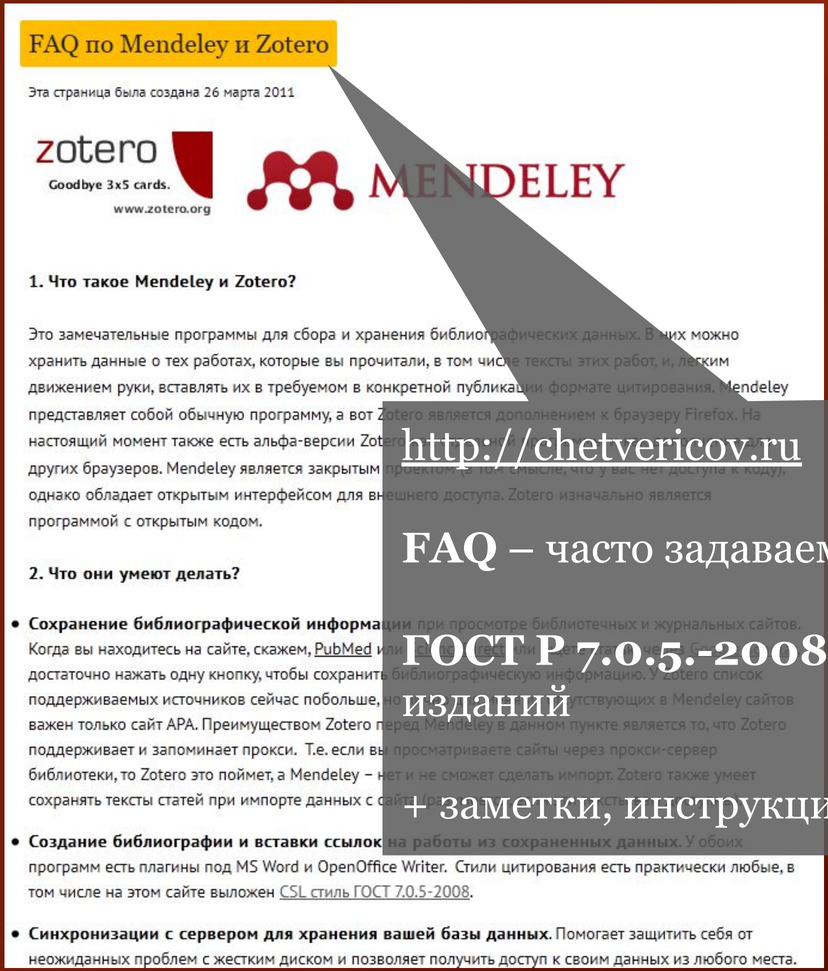

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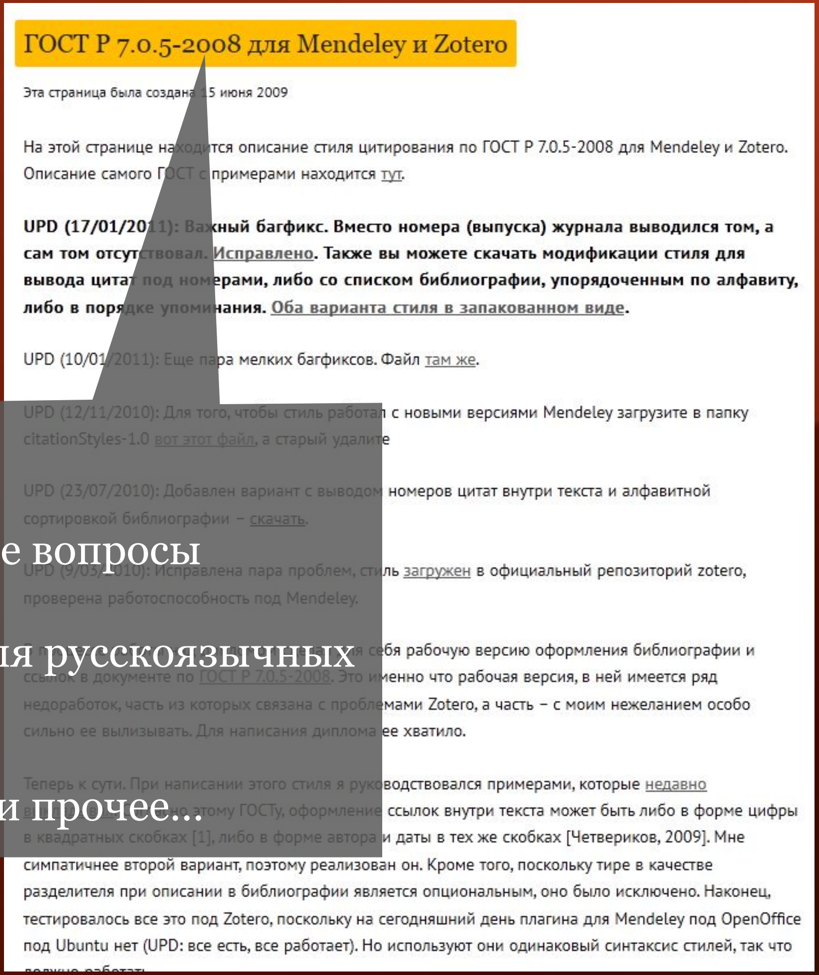
Эта страница была создана 26 марта 2011

1. Что такое Mendeley и Zotero?

Это замечательные программы для сбора и хранения библиографических данных. В них можно хранить данные о тех работах, которые вы прочитали, в том числе тексты этих работ, и, легким движением руки, вставлять их в требуемом в конкретной публикации формате цитирования. Mendeley представляет собой обычную программу, а вот Zotero является дополнением к браузеру Firefox. На настоящий момент также есть альфа-версии Zotero [illegible] других браузеров. Mendeley является закрытым проектом (в том смысле, что у вас нет доступа к коду), однако обладает открытым интерфейсом для внешнего доступа. Zotero изначально является программой с открытым кодом.

2. Что они умеют делать?

- **Сохранение библиографической информации** при просмотре библиотечных и журнальных сайтов. Когда вы находитесь на сайте, скажем, PubMed или [illegible] достаточно нажать одну кнопку, чтобы сохранить библиографическую информацию. У Zotero список поддерживаемых источников сейчас побольше, но [illegible]утствующих в Mendeley сайтов важен только сайт APA. Преимуществом Zotero перед Mendeley в данном пункте является то, что Zotero поддерживает и запоминает прокси. Т.е. если вы просматриваете сайты через прокси-сервер библиотеки, то Zotero это поймет, а Mendeley – нет и не сможет сделать импорт. Zotero также умеет сохранять тексты статей при импорте данных с сайта [illegible]
- **Создание библиографии и вставки ссылок на работы из сохраненных данных.** У обоих программ есть плагины под MS Word и OpenOffice Writer. Стили цитирования есть практически любые, в том числе на этом сайте выложен CSL стиль ГОСТ 7.0.5-2008.
- **Синхронизации с сервером для хранения вашей базы данных.** Помогает защитить себя от неожиданных проблем с жестким диском и позволяет получить доступ к своим данных из любого места.



ГОСТ Р 7.0.5-2008 для Mendeley и Zotero

Эта страница была создана 15 июня 2009

На этой странице находится описание стиля цитирования по ГОСТ Р 7.0.5-2008 для Mendeley и Zotero. Описание самого ГОСТ с примерами находится тут.

UPD (17/01/2011): Важный багфикс. Вместо номера (выпуска) журнала выводился том, а сам том отсутствовал. Исправлено. Также вы можете скачать модификации стиля для вывода цитат под номерами, либо со списком библиографии, упорядоченным по алфавиту, либо в порядке упоминания. Оба варианта стиля в запакованном виде.

UPD (10/01/2011): Еще пара мелких багфиксов. Файл там же.

UPD (12/11/2010): Для того, чтобы стиль работал с новыми версиями Mendeley загрузите в папку citationStyles-1.0 вот этот файл, а старый удалите

UPD (23/07/2010): Добавлен вариант с выводом номеров цитат внутри текста и алфавитной сортировкой библиографии – скачать.

UPD (9/03/2010): Исправлена пара проблем, стиль загружен в официальный репозиторий zotero, проверена работоспособность под Mendeley.

[illegible] себя рабочую версию оформления библиографии и ссылок в документе по ГОСТ Р 7.0.5-2008. Это именно что рабочая версия, в ней имеется ряд недоработок, часть из которых связана с проблемами Zotero, а часть – с моим нежеланием особо сильно ее вылизывать. Для написания диплома ее хватило.

Теперь к сути. При написании этого стиля я руководствовался примерами, которые недавно [illegible] этому ГОСТу, оформление ссылок внутри текста может быть либо в форме цифры в квадратных скобках [1], либо в форме автора и даты в тех же скобках [Четвериков, 2009]. Мне симпатичнее второй вариант, поэтому реализован он. Кроме того, поскольку тире в качестве разделителя при описании в библиографии является опциональным, оно было исключено. Наконец, тестировалось все это под Zotero, поскольку на сегодняшний день плагина для Mendeley под OpenOffice под Ubuntu нет (UPD: все есть, все работает). Но используют они одинаковый синтаксис стилей, так что

http://chetvericov.ru

FAQ – часто задаваемые вопросы

ГОСТ Р 7.0.5.-2008 для русскоязычных изданий

+ заметки, инструкции и прочее…

MENDELEY

www.mendeley.com