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Quick Start Guide to Springer Protocols

Springer Protocols is an online database of reproducible laboratory protocols in the biomedical and life sciences.

PLEASE NOTE that the Online Library does not subscribe to this database, so you can only access free protocols.

All free protocols have a yellow F next to the title or can be found by clicking 'Free Protocols' on the right side of the page:



Figure 1. How to access freely available protocols.



Browsing and Searching

Springer Protocols can be accessed via this URL: <u>http://www.springerprotocols.com/</u> This website offers basic and advanced searching, as well as browsing by subject.

Browsing

On the homepage there is 'Browse by Subject' area. When clicking on a subject area, it will take you to a list of all protocols relating to that subject.

Please note that this will contain material that is inaccessible. Keep an eye out for the **I** symbol for free material.



Figure 2. Browsing options.

On clicking on a subject you will then be taken to a results page, as shown in Figure 3. From here you can narrow your search using the options on the left hand side of the screen as shown in Figure 3.

You can either search within these results, or there is the option to browse further subject categorisations within the subject area you have already chosen. The numbers in brackets are the numbers of items relevant to that topic.



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	Welcome. Sign in <u>here</u> . New user? Register <u>here</u> .				
Search Within These Results	Results 1 - 10 of 128	1 2 3 4 5 6 7 8 9 10 Next>>			
	Search results for: Text "cytochemical techniqu	ues" - all of the words/ (Protocol search)			
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	Sort results by: Relevance 10	per page Collapse View			
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Cancer Research (8)	Cytochemical techniques and energy-	filtering transmission electron			
Cell Biology (39)	microscopy applied to the study of parasitic protozoa				
Genetics/Genomics (12)	Author(s): Marcos A. Vannier-Santos, Ulysses Lins				
Imaging/Radiology (4)	Summary: Cytochemical techniques and energy-filtering transmission electron				
Immunology (7)	microscopy applied to the study of parasitic protozoa The study of parasitic protozoa				
Infectious Diseases (1)	Abstract Full Text PDF (2326K) Open Access				
Microbiology (2)					
Malandan Madisira (10)	Electron Microscopic Enzyme Cytochemistry				
Molecular Medicine (10)	Author(s): Nobukazu Araki, Tanenori Hatae				
Neuroscience (21)	Pub. Date: Mar-09-1999; DOI:10.1385/1-59259-201-5:159				
Pharmacology/Toxicology (2)	cytochemistry is well established and one of the most common techniques. The original				
Plant Sciences (4)	method for the cytochemical demonstration				
Protein Science (3)					
Durance by Mann	Signal Amplification for DNA and mRNA				
Browse by Year	Author(s): Ernst J. Speel, Anton H. Hopman,	, Paul Komminoth			
2014-2016 (7)	Pub. Date: Sept-01-1999; DOI:10.1385/1-5	59259-677-0:195			
2011-2013 (22)	Summary: fluorescence of enzyme cytochemical visualization. HRP, horseradish peroxidase; AP, alkaline phosphatase. On the other hand, more and more literature is				
2008-2010 (14)	becoming available that describes approaches a	to amplify			
2005-2007 (14)					
2002-2004 (19)	Human Myoblasts from Skeletal Muscle B	iopsies: In Vitro Culture			
1999-2001 (13)	Preparations for Morphological and Cytoc Electron Microscopy	chemical Analyses at Light and			
1996-1998 (19)	Author(s): Manuela Malatesta, Marzia Giagnacovo, Rosanna Cardani, Giovanni Meda, Carlo Pellicriari				
1993-1995 (7)	Pub. Date: Mar-13-2013; DOI:10.1007/978-1-62703-317-6_6				
1990-1992 (5)	Summary: index (i.e., the percentage of myoblasts in the cell population) ranges				
1987-1989 (6)	petween 50 and 80, in the cell cultures at the first passage. Among the manifold cytochemical and immunocytochemical techniques				
1984-1986 (2)	Abstract Full Text PDF (577K)				
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Figure 3. Searching and ordering your browsing search results.

There is also the option to 'Browse by Year' on the bottom left hand side of the screen. You can also sort the results that appear by selecting the drop down box as shown at the top of Figure 3. You can order results by relevance, date (most recent), author name and title.

To see further information, click on the title of the item.



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Searching

The homepage provides a basic search box for you to conduct a search with.

Springer Pr	otocols	us a RSS 🛛 HELP			
	HOME MY A	CCOUNT MY PROTOCOLS			
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Mobile		Inside SpringerProtocols			
SpringerProtocols goes mobile! Learn about our new mobile site, available now.	Search Protocols Search	Source Title List			
😻 Upload a Protocol	Advanced Search	★ Free Protocols ♣ Popular Protocols			
Upload your own protocols for		79			

Figure 4. Search box on homepage.

When searching using this box, you will see a similar screen to the one when you have browsed. The results appear as in Figure 3.

There is also an advanced search option which allows you to fill in a variety of different fields to bring back a smaller, more specific number of results.

Advanced S	Search		
Select Option	Protocols Books		
Anywhere in Text:	🔹 all 🔍 any 🔍 exact	phrase	
Abstract:	💿 all 🔍 any 🔍 exact	phrase	
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Series:	Select Series 🔻		
Volume No:			
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Subject:	Select Subject 🔻		
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Figure 5. Advanced search.

Please note that when using the browse and search functions of this resource, the results will include anything relevant from its collection and not all of the content may be accessible as the Online Library does not subscribe to this resource. Remember to keep an eye out for the yellow F next to the title

I to find those that are freely available:



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Figure 6. Example of a freely available text.

Once you have clicked on the title of the item, you will see a page like the one in Figure 7:

Comparison of Enzymatic and Non-Enzymatic Means of Dissociating Adherent Monolayers of Mesenchymal Stem Cells

By: Boon C. Heng², Catherine M. Cowan², Shubhayu Basu¹ ⊠

Abstract



The dissociation of adherent mesenchymal stem cell (MSC) monolayers with trypsin and enzyme-free dissociation buffer was compared. A significantly lower proportion of viable cells were obtained with enzyme-free dissociation buffers compared to trypsin. Subsequently, the dissociated cells were re-seeded on new cell culture dishes and were subjected to the MTT assay 24 h later. The proportion of viable cells that reattached was significantly lower for cells obtained by dissociation with enzyme-free dissociation buffer compared to trypsin. Frozen-thawed MSC displayed a similar trend, yielding consistently higher cell viability and reattachment rates when dissociated with trypsin compared to enzyme-free dissociation buffer. It was also demonstrated that exposure of trypsin-dissociated MSC to enzyme-free dissociation buffer for 1 h had no significant detrimental effect on cell viability.

Images from this Protocol

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Affiliation(s): (1) New Business Ventures, Abbott Vascular, 3200 Lakeside Dr., Santa Clara, CA 95054, USA

(2) Abbott Vascular, 3200 Lakeside Dr., Santa Clara, CA 95054, USA Journal Title: Biological Procedures Online

Volume: 11 | Issue: 1 | Pub. Date: Dec-01-2009 | Page Range: 161-169 | DOI: 10.1007/s12575-009-9001-4

Subject: Biochemistry

Key Words: Dissociation - Enzyme - Mesenchymal - Stem cells - Trypsin

2

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Post comment/View all comments

By LAC Springer Training Apr-06-2015 02:53 AM This is a test about an example of comments Report Violation

Figure 7. Example of a freely available text record and how to access the full text.



You will be able to download the full text by selecting on the hyperlink 'Full Text' or 'Download PDF' options as shown in Figure 7.

There is a helpful video provided by Springer to guide users on how to use and conduct searches on their resource:

http://www.springerprotocols.com/tour/Protocols_Introduction/Protocols_Introduction.vm

Further Help

If you need help using this or any other information resources, please contact the **Online Library** by:

Telephone at: +44 (0)20 7862 8478 (between 09.00 and 17.00 GMT),

By email at: OnlineLibrary@shl.lon.ac.uk

By the Enquiries Form at: http://onlinelibrary.london.ac.uk/about/contact-us