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# Working Paper

Science, Innovation and Electronic Information Division

# **Biotechnology Use and Development** Survey: methodology, issues and responses

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Statistics

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This paper represents the views of the author and does not necessarily reflect the opinions of Statistics Canada.





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#### The science and innovation information program

The purpose of this program is to develop **useful indicators of science and technology activity** in Canada based on a framework that ties them together into a coherent picture. To achieve the purpose, statistical indicators are being developed in five key entities:

- Actors: are persons and institutions engaged in S&T activities. Measures include distinguishing R&D performers, identifying universities that license their technologies, and determining the field of study of graduates.
- **Activities**: include the creation, transmission or use of S&T knowledge including research and development, innovation, and use of technologies.
- Linkages: are the means by which S&T knowledge is transferred among actors. Measures include the flow of graduates to industries, the licensing of a university's technology to a company, co-authorship of scientific papers, the source of ideas for innovation in industry.
- Outcomes: are the medium-term consequences of activities. An outcome of an innovation in a firm may be more highly skilled jobs. An outcome of a firm adopting a new technology may be a greater market share for that firm.
- Impacts: are the longer-term consequences of activities, linkages and outcomes. Wireless telephony is the result of many activities, linkages and outcomes. It has wide-ranging economic and social impacts such as increased connectedness.

The development of these indicators and their further elaboration is being done at Statistics Canada, in collaboration with other government departments and agencies, and a network of contractors.

Prior to the start of this work, the ongoing measurements of S&T activities were limited to the investment of money and human resources in research and development (R&D). For governments, there were also measures of related scientific activity (RSA) such as surveys and routine testing. These measures presented a limited picture of science and technology in Canada. More measures were needed to improve the picture.

Innovation makes firms competitive and we are continuing with our efforts to understand the characteristics of innovative and non-innovative firms, especially in the service sector that dominates the Canadian Economy. The capacity to innovate resides in people and measures are being developed of the characteristics of people in those industries that lead science and technology activity. In these same industries, measures are being made of the creation and the loss of jobs as part of understanding the impact of technological change.

The federal government is a principal player in science and technology in which it invests over five billion dollars each year. In the past, it has been possible to say only *how much* the federal government spends and *where* it spends it. Our report **Federal Scientific Activities**, **1998 (Cat. No. 88-204)** first published socio-economic objectives indicators to show *what* the S&T money is spent on. As well as offering a basis for a public debate on the priorities of government spending, all of this information has been used to provide a context for performance reports of individual departments and agencies.

As of April 1999, the Program has been established as a part of Statistics Canada's Science, Innovation and Electronic Information Division.

The final version of the framework that guides the future elaboration of indicators was published in December, 1998 (Science and Technology Activities and Impacts: A Framework for a Statistical Information System, Cat. No. 88-522). The framework has given rise to A Five-Year Strategic Plan for the Development of an Information System for Science and Technology (Cat. No. 88-523).

It is now possible to report on the Canadian system on science and technology and show the role of the federal government in that system.

Our working papers and research papers are available at no cost on the Statistics Canada Internet site at http://www.statcan.ca/cgi-bin/downpub/research.cgi?subject=193.

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# Biotechnology Use and Development Survey: methodology issues and responses

#### I Background

Scientific breakthroughs in genomics, proteomics and bioinformatics have given an "élan" to the development of biotechnology activities. This is apparent in 1) the increasing number of innovative firms in biotechnology and 2) the rise in the leading indicators of biotechnology activities, namely, the number of biotechnology employees, biotech revenues, the number of products at both developmental and production stages, and R&D expenditures.

Collecting sound data on biotechnology is a real challenge (Rose, 2000). Biotechnology is a pervasive technology used in several industrial sectors. Second, unlike most industrial activities where "physical" components are assembled to obtain the end product, biotechnology is a set of techniques that are used in current operation or to develop new products and processes.

In 1996, the first ever survey of biotechnology by a national statistical agency was undertaken by Statistics Canada. Entitled "Survey of Biotechnology Use in Canadian Industries – 1996", it aimed at assessing the use of biotechnology by selected Canadian industries: Aquaculture and Forestry, Agro-industry, Wood, Pulp and Paper, Coal/Oil/Gas, and Chemicals. Strong indications of widespread biotechnology utilization by the Canadian industry gave rise to the Biotechnology Firm Survey - 1997. This was followed by the 1999 Biotechnology Use and Development (BUDS) and the 2001 BUDS. These last three surveys went beyond measuring the use of biotechnologies to emphasize the development of new products and processes using biotechnology. The objective of the extension in the scope of biotech surveys was two-fold: 1) to satisfy the information needs of both policy makers and the industry as they related to biotechnology adoption, development and diffusion; and, 2) to document and measure the characteristics of Canadian firms that are using biotechnology to develop new products and processes.

This paper explains the methodologies used in these surveys, the problems encountered, the responses to these problems, and the lessons learned. The organization of the paper is as follows: definitional issues will be presented in the second section; the target population and the sampling issues, as they relate to the management of response burden, in section 3; data quality in section 4; a brief description of the 1997 Biotechnology Firm Survey, the 1999 and the 2001 BUDS in section 5; issues related to the change in the survey methodology, as well as their impacts on the estimates, in section 6; the lessons learned in section 7; and the conclusions in section 8.

# II Defining biotechnology and an Innovative Biotechnology Firm: Issues and Responses

#### 2.1 What is Biotechnology?

Because of its pervasive nature, biotechnology is best thought of as a cross-sectoral or cross-industrial activity, rather than a sector or an industry. In fact, as a set of scientific techniques, it has multiple usage across sectors and industrial fields and does not fall under any single code of the North American Industry Classification System (NAICS)<sup>1</sup> (Statistics Canada, 1997). Consequently, defining biotechnology statistically is a challenge. Most of the earlier definitions are found in Rose (2000).

Statistics Canada uses a double-definition strategy: a "normative" or single definition and an "operational" or list-based definition. According to the single definition biotechnology is the "application of science and engineering in the direct or indirect use of living organisms or parts of organisms in their natural or modified forms in an innovative manner in the production of goods and services or to improve existing processes"<sup>2</sup>.

The problem associated with the use of such a general definition is that the interpretation of the definition can vary broadly and is left to the respondents. To help with the interpretation of this definition, a list-based approach was developed. The use of a list was preferred by respondents during questionnaire testing as being more precise.

In 1997, the list-based definition included 17 specific biotechnologies grouped under 3 main blocks (Appendix1): DNA-based biotechnologies; Biochemistry/Immunology; and Bioprocessing-based biotechnologies. The same grouping was used in the 1999 BUDS, except that the last block was divided into Bioprocessing based biotechnologies and Environment (Appendix 2). In the 2001 BUDS, the list-based definition comprised 19 specific biotechnologies in 5 main blocks (Appendix 4): DNA – the coding, Proteins and Molecules – the Functional Blocks, Cell and Tissue Culture, and Engineering, Process Biotechnologies, Sub-cellular Organisms. The provisional definition of biotechnology adopted by the OECD used a similar structure (OECD, 2003).

#### 2.2 What is an innovative biotechnology Firm?

Biotechnologies are techniques or set of techniques. Firms may use these techniques within their production processes for different reasons. For example, biotechnologies may be used for environmental purposes. Other firms may use them to develop products and processes. Many firms in this latter group, the main focus of the Biotechnology Use and Development Surveys, have no product on the market. For example, in 1997, 67% of the

<sup>&</sup>lt;sup>1</sup> This is why, at Statistics Canada, the term "biotechnology activities" instead of "biotechnology sector" or "biotechnology industry" is used to describe the activities related to the use of biotechnology techniques to develop new products/processes.

<sup>&</sup>lt;sup>2</sup> Under the leadership of Canada, the provisional single definition from the OECD to be used in the 2003 BUDS is as follows "the application of S&T to living organisms as well as parts, products and models thereof, to alter living or non-living materials for the production of knowledge, goods and services".

firms had no product on the market. This figure was 47% in 1999 and 52% in 2001. Thus, defining an innovative biotechnology firm <u>solely</u> in terms of the number of products on the market would have excluded a great number of firms. This problem was overcome by defining an innovative biotech firm as "a firm developing new products and processes that require the use of biotechnologies".

In practice, an innovative biotech firms is identified by doing two things. First, respondent firms have to be using at least one biotechnology. Second, this use has to be geared toward developing new products or processes. This latter condition is met when a firm reports having at least one product or process at any developmental stage, in production, approved, or on the market that require the use of biotechnologies. More often than not, such firms have employees dedicated to biotechnology activities and are conducting biotechnology related R&D.

An innovative biotechnology firm is thus defined in terms of its ability to use biotechnologies in its production process (process innovation), rather than having introduced new products on the market (product innovation). In this respect, the definition used is more in line with the OECD/EUROSTAT's definition of a process innovation, i.e. "the use of a technologically new or significantly technologically improved processes within a production process" (OECD/EUROSTAT, 1997).

#### III Target population, Sampling Unit

#### 3.1 Target Population

The target population of the biotechnology surveys are all firms that use biotechnology in their production process or to develop products or processes. These firms are not found under any single NAICS code. Thus reaching them requires selecting specific NAICS codes. The choice of these NAICS codes is based on the probability of biotechnology use, past experiences, and expert opinion. It involves Statistics Canada, industry experts, and federal departments such as Industry Canada, and Agriculture and Agri-food Canada. Firms that provide only biotechnology related services are excluded.

The 1997 Biotechnology Firm survey was based on 475 firms the name and addresses of which were obtained from Industry Canada, the 1998 Canadian Biotechnology Directory maintained by Contact Canada (Contact Canada, 1997), and the Statistics Canada Industrial R&D survey (Traoré, 2001; Laroche, 2001). The 1999 and the 2001 BUDS included both a "must-take-all" list of firms put in place using the 1997 methodology and a sample of firms from Statistics Canada's Business Register (BR). The specific NAICS codes from which the sample of firms were drawn are presented in Table 1.

Table 1: Selected NAICS from which the Survey Sample was drawn, 1999, 2001

NAICS	Associated Industry
1125	Animal Aquaculture
1132	Forest Nurseries and Gathering of Forest Products
2111	Oil and Gas Extraction
2122	Metal Ore Mining
2123	Non-Metal Mineral Mining and Quarrying
311	Food Manufacturing
3121	Beverage Manufacturing
31311 <sup>(a)</sup>	Fibre, Yarn and Thread Mills
3221	Pulp, Paper and Paperboard Mills
3222 <sup>(a)</sup>	Converted Paper Product Manufacturing
32411	Petroleum Refineries
325	Chemical Manufacturing
3254	Pharmaceutical and Medicine Manufacturing
4145	Pharmaceuticals, Toiletries, Cosmetics and Sundries Wholesalers-Distributors
4183	Agricultural Supplies Wholesalers-Distributors
5417 <sup>(a)</sup>	Scientific Research and Development Services
6215 <sup>(a)</sup>	Medical and Diagnostic Laboratories

Sources: Statistics Canada 1997. North American Industry Classification Systems, Cat.No. 12-501-XPE,

Note: (a) refers to NAICS codes that were surveyed only in 2001.

#### 3.2 Sampling Unit

Two types of firms are found in the Biotechnology surveys: single location firms and multiple location firms. The former group is made of firms for which the enterprise and the establishment as defined by Statistics Canada are the same. The latter group concerns all the enterprises with multiple establishments. In some cases, not all these establishments are in the same NAICS code. In other words, some multiple location firms have more than one NAICS code.

The need for any Statistics Canada's survey to produce data at the provincial and territorial levels was an overriding reason for the choice of the sampling unit. Thus, for the multi-establishment firms, all the establishments in a given province or territory were grouped to form a single unit. The NAICS code assigned to this new statistical unit is that of the establishment with the largest share of gross business income (GBI). In cases where two or more establishments have the same GBI, the assigned NAICS code is randomly chosen among these units (Morin et al., 2001; Lavigne, 2003). Consequently, the sampling unit is the "provincial" enterprise<sup>3</sup>.

<sup>&</sup>lt;sup>3</sup> The "provincial" enterprise as a sampling unit was also used in the 1999 Innovation Survey. Because of increased demand by data users for data at the Census Agglomerations (CA) and the Census Metropolitan Areas (CMA) levels, the survey on Innovation in Services and the 2003 Biotechnology Use and Development surveys will be using the "establishment" as the sampling unit. This refers to an operation, in contrast to the "provincial" enterprise which is the "sum" of all the operations of a given firm in a province.

#### 3.3 Data strata

Producing estimates at the provincial and territorial levels is an important requirement for all the surveys at Statistics Canada. Consequently, province/territory was an important dimension in stratifying the sample. These dimension has 12 levels: 10 provinces and 2 territories<sup>4</sup>. Two other dimensions, namely, the NAICS code (industrial sector) and firm size were deemed appropriate levels at which to produce estimates. The former is made of the 13 NAICS codes shown in table 2 for the 1999 BUDS; and 17 NAICS codes in the 2001 BUDS (Table 1). Size has 3 levels, small, medium, and large. As shown in Table 2, in 1999, this resulted in 468 (= 12\*13\*3) different strata and in 2001, 612 strata (=12\*17\*3).

Given that 1) Statistics Canada has no single standardized definition of "small businesses" and 2) most innovative biotechnology firms are years away from any revenues, it was not deemed appropriate to define size in terms of revenues. Instead, it was decided to use the number of employees in any given firm. As a result, small biotech firms are all the firms with 50 employees or fewer, medium-sized firms are firms that have between 51 and 150 employees, and large firms have over 150 employees. The choice of these size thresholds came about in the 1997 Biotechnology Firm Survey. Statistics Canada wanted to retain some measure of comparability with other existing sources of biotechnology data in Canada. Subsequently, to enable comparison of the estimates from one survey to the other, these thresholds were retained and continue to be used<sup>5</sup>. We use Statistics Canada's standard classes in any subsequent surveys.

Table 2: Number of Strata Used in the 1999 and 2001 BUDS

	1999	2001
Number of provinces and territories <sup>(a)</sup>	12	12
Number of industries (NAICS)	13	17
Number of firm size categories	3	3
Total	468	612

Source: Statistics Canada, BUDS 1999 & 2001

Note:

#### 3.4 Respondent Selection and Respondent Burden

#### 3.4.1 Respondent Selection

Respondent selection was undertaken based on two main considerations: to reach the target population; and, to minimize respondent burden. To this end, 1) GBI, R&D expenditures and number of employees were used as selection criteria, 2) industries were

<sup>(</sup>a) Nunavut was not surveyed

<sup>&</sup>lt;sup>4</sup> Nunavut was not surveyed.

<sup>&</sup>lt;sup>5</sup> Size classes in the BR are less than 50 employees, 50 to 149 employees and 150 employees and up. While the use of different size classes in the Biotechnology surveys results in an insignificant difference in the results, the BR classes will be used as soon as resources are available to reanalyze the earlier surveys.

surveyed at levels other than 4-digits, and 3) small firms with 50 employees or fewer were not rotated out of the survey even after being included in two consecutive surveys.

Based on results from the 1997 Biotechnology Firm Survey, 65% of firms conducting 35% of all R&D in biotechnology would have been excluded had the GBI been the only selection criterion. This problem was overcome by using the GBI along with the number of employees and the amount of R&D expenditures to select respondents from both the BR and the must-take-all list.

Firms with less than 5 employees and spending less than \$100,000 in R&D expenditures. These firms contribute less than 1% to both expenditures in R&D and to the number of new products/processes. Thus, the exclusion does not result in any noticeable bias in the estimates. In addition, it reduces respondent burden.

Furthermore, as shown in table 1, even though most industries were surveyed at the 4-digit level, some industries such as Food Manufacturing (NAICS 311) and Chemical Manufacturing (NAICS 325) were surveyed at the 3-digit level. Petroleum Refineries was surveyed at the 5-digit level (NAICS 32411).

The universe of firms involved in using biotechnology to develop new products and processes was made of 282 firms in 1997, 358 in 1999, and 375 in 2001. Over 70% of these are small firms with 50 employees or fewer. Thus, rotating these firms out of the sample after two years in the survey would invalidate the estimates.

#### 3.4.2 Respondent Burden

Based on respondent reports, 90 minutes were necessary to answer to the full 1997 Biotechnology Firm Survey. In the 1999 BUDS, a multi-stage screening process was implemented. It helped in reducing the response burden by progressively screening out firms based on the extent of their biotechnology activities. Thus, the time required to fill in the survey was 15 minutes for non-users of biotechnology, 30 minutes for users, and 90 minutes for innovative biotech firms. In 2001, the pre-contact questionnaire could be filled in within 5 minutes. Applying the same screening technique as in 1999, in the 2001 BUDS, users were able to complete the full questionnaire of the second phase within 30 minutes and innovative firms, within 90 minutes.

#### 3.5 Other exclusions from the survey

In addition to small firms with less than 5 employees and spending less than \$100,000 in R&D, additional exclusions from the surveys include not-for-profit organizations, universities, government laboratories, and hospitals. Also excluded from the survey are contract research organizations (CROs) that provide only services to biotechnology firms. These entities, even though related to biotechnologies through the provision of R&D services and the creation of spin-off firms, do not meet the main criterion of biotechnology surveys, i.e. to provide information on firms that use biotechnology to develop new products and processes.

#### IV Data Quality, Data Collection, Edit and Imputation

In all the biotechnology surveys, the data quality was insured by taking into account and applying throughout the survey process all 6 dimensions of data quality control at Statistics Canada, namely, relevance of data collected, their accuracy, their timeliness, their accessibility, their interpretability, and their coherence (Statistics Canada, 2003; 2002; 1998).

#### 4.1 Data Quality

#### 4.1.1 Relevance of the data

To insure the relevance of the data to policy makers, industry and analysts, all the questionnaires were designed by the Life Sciences Unit of the Science, Innovation and Electronic Division at Statistics Canada in collaboration with industry experts and federal stakeholders such as Industry Canada and Agriculture and Agri-Food Canada. Also, as shown below, the relevance of the data was insured by making the survey results and data available to users in a timely manner.

#### 4.1.2 Accuracy of the data

Accuracy of data collected was insured by conducting cognitive interviews with small samples of respondents in both official languages with help from the Questionnaire Design Resource Centre (QDRC). Feedbacks and comments from these interviews were integrated into the questionnaire to improve the design and most importantly, to insure that the questions were well understood by respondents.

A knowledgeable person in the respondent firms, such as an R&D manager or a production manager, was asked to answer the questionnaire. Such individuals were able to provide with accuracy the type of information requested in the questionnaires.

As shown below, further checks into the accuracy of the data were conducted by analyzing non-response rate. Furthermore, the sampling process was specifically designed to reach the target population.

#### 4.1.3 Timeliness of the data

To date, all the biotech surveys, from the inception period to the publication of the results, have been completed on time, i.e. within a year time. For example, the consultation process for the 2001 BUDS began in the Fall of 2001 and the survey results were published in the Fall of 2002.

#### 4.1.4 Accessibility to the data

Data from the biotechnology surveys are made available to users through a number of venues, including the sharing of data and statistics with provincial statistical institutions. Other venues include the Facilitated access program whereby researchers from academia are given access to the survey data to conduct research on their own topics.

Analytical and working papers by staff of the Life Sciences section of the Science, Innovation and Electronic Information Division are published in scientific journal or made available on the web-site of Statistics Canada. Some of these studies are also presented in international conferences. These are in addition to descriptive tables provided to different stakeholders such as Industry Canada and Agriculture and Agri-Food Canada

#### 4.1.5 Interpretability

All the Biotechnology surveys are registered in the Integrated Metadatabase (IMDB). The information available concern the definitions of the underlying concepts, the classifications used, the methodology of data collection and processing, and indications of the accuracy of the data and other related statistical information. These records are available to data users on Statistics Canada web-site.

#### 4.1.6 Coherence

Coherence in the data is insured by using the same definitions from across surveys, as well as the same classifications. The target population has remained the same so as the data collection process.

There was a methodology change between the 1997 Biotechnology Firm survey and the 1999 Biotechnology Use and Development survey. However, as shown below, this change had minor effects on the estimates.

### 4.1.7 Response Rates<sup>6</sup>

Of the 475 firms that were sent the 1997 Biotechnology Firm Survey, 392 filled out and returned the questionnaires, resulting into an overall response rate of 83%. The 1999 BUDS was mailed to 3,377 firms. The response rate was 66% or 2,229 returned questionnaires. In the 2001 BUDS, the first phase or "pre-contact" questionnaire was mailed in the Winter of 2002 to a sample of 11,262 firms. The response rate was 70%. The full or second stage questionnaire was mailed in the Spring of 2002 to 900 firms and had a response rate of 84%.

In each of the surveys, analyses of non-respondents were conducted. No evidence of bias was found as these analyses showed that non-respondents had the same characteristics as respondents to the surveys. Consequently, estimates from the surveys provide a clear picture of the biotechnology universe for each period and are a valid representation of the Canadian biotechnology firm population in each year.

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#### 4.2 Data Collection, Edit and Imputation

<sup>&</sup>lt;sup>6</sup> The detailed response rates are found in tables in section 5 below

#### 4 2.1 Data Collection and Edit

The data collection was done by staff of the Science, Innovation and Electronic Information Division. The surveys were direct and mandatory, meaning that the data were requested directly from respondents who were required to provide the requested information. The surveys were mail-out/mail back surveys. Reminder cards and telephone calls were used for subsequent follow-ups.

Post collection edits were undertaken in collaboration with the Business Survey Methodology Division (BSMD) to insure consistency in the data. These edits were applied to complete questionnaires. In 1997, a questionnaire was considered completed when questions 2, 3, 5 and 7 (Appendix 1) were answered. In 1999, it referred to one with responses to questions 1, 5, 7, and 13 (Appendix 2); and in 2001, it applied to a questionnaire with answers to questions 1, 4, 9, 10, and 16 (Appendix 4)<sup>7</sup>.

#### 4.2.2 Imputation techniques and Imputation rates

Two imputation techniques were used, namely, deterministic and hot-deck. Deterministic or deductive imputation was used in cases where logical relations existed among variables. In such cases, the "revealed" behavior of the respondent unit in a preceding question influenced the response to a subsequent question(s). Consequently, the revealed answer was used to impute the logical value of the missing response. Examples of such related questions were skip pattern questions. For instance, in the 2001 BUDS, in question 5a, respondents were asked if they had any unfilled biotechnology position. Those that answered "NO" to that question were asked to skip to question 5b. Otherwise, they were to fill-in a table. In cases where the table was filled and question 5a was not either answered or the given answer was "NO", the response was changed to "YES".

In cases where deterministic imputation did not apply, hot-deck imputation was used. It consisted of choosing a random donor from the same homogenous response group as the non-respondent, starting with the smallest, i.e. Province/NAICS/Size. For example, to find a donor for a small firm active in Agriculture in Quebec, the search would start in the Quebec/Agriculture/Small response group. If no donor is found in this group, the response group is extended to Province/NAICS. This process continues until a donor is found.

Overall, the imputation rates, obtained by dividing the number of imputed respondents to a question by the total number of respondents, were low for all three biotechnology surveys: on average 3.7% in 1997, 12.9% in 1999 and about 7% in 2001. For the 1997 Biotechnology Firm Survey, the rates of imputation range from 0 for a series of questions to a high of 5.3% for Q8. In the 1999 BUDS, except for Q11, i.e. where the imputation rate was 35%, all other questions had a rate of imputation between 0, for Q2, and 19.5%

<sup>&</sup>lt;sup>7</sup> In all the cases, these questions referred to 1) the use of biotechnologies in either current production or for environmental purposes, 2) the presence in the firm of employees dedicated to biotechnology, 3) the development of products/processes using biotechnologies, 4) having biotechnology products/processes at either developmental or approved/production stages, and 5) conducting biotechnology R&D.

for Q13. As for the 2001 BUDS, the rates range from 1.9% to a high of 18.3%. The imputation rates for "quantitative" questions were quite low, 2.9% for Q4, i.e. Human resources, .8% for Q10, biotechnology products, 7% for Intellectual Properties, 3.8% for Capital raised, 4.8% for Exports and Imports question (Table 3).

Table 3: Imputation rates, Biotechnology Firm Survey – 1997, BUDS 1999, BUDS 2001

Questions Imputation rates (%)			
	Biotechnology Firm Survey - 1997		BUDS - 2001
Q1	1	7.21	6.70
Q2	5.19	0	7.74
Q3	0	15.13	8.96
Q4	2.75	15.76	2.95
Q5	9	13.62	7.98
Q6	8	16.83	5.99
Q7	0	12.89	5.34
Q8	5.20	1.46	9.29
Q9	0	16.38	8.99
Q10	10	16.7	.79
Q11	0	34.96	17.99
Q12		16.59	8.80
Q13		19.45	3.16
Q14			6.98
Q15			10.18
Q16			7.35
Q17			8.96
Q18			2.57
Q19			9.47
Q20			1.98
Q21			1.98
Q22			3.49
Q23			3.81
Q24			12.38
Q25			18.26
Q26			4.77
Q27			1.93
Q28			9.52
Q29			9.25

Source: Statistics Canada

#### 4.3 Estimates

Estimates from surveys are used to produce population estimates. This requires that the sample weights of the responding units be adjusted to take into account non-respondents.

This procedure was applied to all the biotechnology surveys. All the estimates were produced using the Generalized Estimation System (GES) in the Statistical Analysis System (SAS) computer package (Statistics Canada, 2002). Estimates were produced at the firm size, industry and province/territory levels. However for this latter level, to meet confidentiality requirements, no estimates could be produced separately for the Atlantic provinces. It was then decided to produce a single estimate for this region, under Atlantic. Similarly, no estimates could be produced for the two territories that were surveyed, namely, Yukon and the Northwest Territories.

Publication of the estimates follows Statistics Canada's guidelines in this matter (Table 4). In cases where the estimates were either unreliable because of high coefficient of variations (CV), i.e. CV above 50%, or did not meet the confidentiality requirements of the Statistics Act, they were suppressed and replaced by the appropriate symbol (Table 4). Whenever the estimates had a low reliability, i.e. a CV between 30% and 50%, they were marked with an E to warn data users that they should be used with some caution (Table 4).

**Table 4: Symbols Used When Publishing the Estimates** 

Symbol	Meaning
	Not available for any reference period
	Not available for a specific reference period
	Not applicable
p	Preliminary
r	Revised
X	Suppressed to meet the requirements of the Statistics Act
E	Use with caution, (CV between 30% and 50%)
F	Too unreliable to be published, (CV more than 50%).

Source: Statistics Canada

### V Brief Description of the Surveys

#### 5.1 1997 Biotechnology Firm Survey - 1997

The 1997 Biotechnology Firm Survey was based on a list of 475 firms<sup>8</sup>. Of these, 138 were out of scope, 44 were out of business, 51 could not be reached or refused to answer the questionnaire, 32 did not return any questionnaire and 210 provided the requested data (Table 5). These resulted in 392 returned questionnaires of 83% survey response rate. After accounting for non-responses and applying post-stratification techniques, it was estimated that 282 firms constituted the core of Canadian biotechnology activities in 1997.

Statistics Canada

<sup>&</sup>lt;sup>8</sup> More information on the 1997 Biotechnology Firm Survey may be found in Traoré (2001) and Laroche (2001).

Table 5: Response Categories and Corresponding Number of Respondents, Biotechnology Firm Survey – 1997

Response categories	Number of respondents
Out of scope	138
Out of business	44
Could not be reached or refused to answer	51
Provided requested information	210
Non-response	32
Total	475

Source: Statistics Canada, Biotechnology Firm Survey - 1997

#### 5.2 Biotechnology Use and Development Survey – 1999

Drawing upon the 1997 Biotechnology Firm Survey, the 1999 BUDS contained additional questions. They related to 1) the barriers to using biotechnologies, 2) the sources of information on biotechnology, 3) the benefits from using biotechnology, and 4) Intellectual property (Appendix 2).

The sample was made of 3,377 firms of which 2,999 came from the BR and 378 from a must-take-all list of firms provided by Statistics Canada and industry experts. Sixty two (62) of these firms were out of business, 40 were out of scope, 13 had either merged with or were acquired by other firms, 123 could not be reached, 20 refused to answer the questionnaire, 1,982 filled out and returned the questionnaire, and no response was received from 1,148 firms (Table 6). Of the 1,982 returned questionnaires, 1,568 were not involved in biotechnology, 192 were biotechnology users, and 223 were classified as innovative firms. As shown in Table 7, 35 of the 223 innovative firms came from the BR and 188 from the Must-take-all list. Likewise,165 of the users came from the BR and 27 from the Must-take-all list.

After accounting for non-respondents, it was estimated that 358 firms made up the population of innovative biotechnology population in Canada in 1999.

Table 6: Response Categories and Corresponding Number of Respondents, Biotechnology Use and Development Survey – 1999

Response categories	Number of respondents
Out of scope	40
Out of business	62
Could not be reached	123
Refused to answer	20
Merger/Acquisition	13
Provided requested data	1,982
Non-response	1,148
Total	3,377

Source: Statistics Canada, Biotechnology Use and Development Survey – 1999

Table 7: Distribution of Innovative, Users among Respondents in the BR Sample and the Must-take-all List, 1999

Origin	Status		Total
	Innovative	Users	
Sample from the BR	35	165	200
Must-take-all	188	27	215
Total	223	192	415

Source: Statistics Canada, 1999 BUDS

#### 5.3 Biotechnology Use and Development Survey – 2001

The 2001 BUDS was done in two phases. The pre-contact or first phase questionnaire was a one page questionnaire (Appendix 3) and was mailed out to 11,262 firms from the BR. It was intended to identify firms that used biotechnology as part of their activities. Seven thousand eight hundred and eighty three (7,883) firms returned the questionnaire. Of these, 512 firms or 6.5% reported using or developing biotechnology products/processes in their daily activities.

This list of 512 firms was supplemented by a "must-take-all" list of 388 firms. The list was constructed using the following sources: Statistics Canada, Agriculture and Agri-Food Canada. The resulting 900 firms where sent the full questionnaire in the Spring of 2002 (Appendix 4). Twenty three (23) were out of scope, 10 were out of business, 45 had either merged with or been acquired by another firm, 24 could not be reached, 9 refused to answer the questionnaire, no questionnaire was received from 143 (Table 8). Of the 646 completed questionnaires, 334 were not involved in biotechnology, 59 were biotechnology users, and 253 were involved in developing biotechnology products/processes. As shown in Table 9, 65 of the innovative came from the BR and 188 came from the Must-take-all list. Forty one (41) of the users came from the BR and 18 from the Must-take-all list.

Taking into account non-respondents, estimates from this second stage indicated that the core of Canadian biotechnology was made of 375 innovative firms in 2001.

Table 8: Response Categories and Corresponding Number of Respondents, Biotechnology Use and Development Survey – 2001

Response categories	Number of respondents
Out of scope	23
Out of business	10
Could not be reached	24
Refused to answer	9
Merger/Acquisition	45
Provided requested data	646
Non-response	143
Total	900

Source: Statistics Canada, Biotechnology Use and Development Survey - 2001

Table 9: Distribution of Innovative, Users among Respondents in the BR Sample and the Must-take-all List, 2001

Origin	Status		Total
	Innovative	Users	
Sample from the BR	65	41	106
Must-take-all	188	18	206
Total	253	59	312

Source: Statistics Canada, 2001 BUDS

# VI Changes in the Survey Methodology and their Impacts on the Estimates

#### 6.1 1997 Biotechnology Firm Survey Vs 1999 BUDS

As stated earlier, the 1997 Biotechnology Firm survey was based on a list of 475 firms (Traoré, 2001; Laroche, 2001). The 1999 BUDS was a combination of a sample of 2,999 firms from the BR and a must-take-all list of 378 firms (McNiven 2001 a&b). Altogether, they returned 2,229 questionnaires of which 1,971 came from the BR and 258 from the must-take-all list. As shown by figures in Table 4, 188 of these 258 firms from the must-take-all list were classified as innovative biotech firms. This amounts to 84% of the 223 firms on which the 1999 biotechnology population estimates were based. Comparatively, 35 firms or 16% of the innovative firms came from the BR sample. In other words, 16% of the 223 innovative firms would not have been accounted for had the must-take-all list been the sole source of firms surveyed in 1999. When the population weights are taken into account, this translates into one third (1/3) of the 358 innovative firms in 1999 coming from the BR and two thirds (2/3) from the must-take-all list. Applying these percentages to the 76 additional firms accounted for in 1999, gives 25 firms from the BR and 51 from the must-take-all list. Thus, 25 additional firms have been identified by the sampling procedure. Put differently, had the 1997 survey methodology been applied in 1999, this would have resulted in an underestimation of the biotechnology firm population by 25 firms.

Since these 76 additional firms represent a 27% increase in biotech firm population over the 1997-1999 period, the actual impact of the change in the survey methodology on the change in any biotechnology activity indicator is 27%\*(1/3) = 9%. For biotechnology leading indicators, this implies that \$121 million of the \$1.35 billion increase in biotech revenues are the arte-fact of the change in the survey methodology; so are \$30 million of the \$333 million increase in biotech R&D, \$37 million of the \$407 million increase in biotech exports, 119 employees in the 1,324 decrease in biotech employment, \$151 million of the \$1.68 billion increase in the amount of capital raised, and 779 products in the 8,650 increase in the total number of biotech products (Table 10).

Comparison between the list of firms in 1997 and the must-take-all list in 1999 shows: 1) 81% of the 258 must-take-all firms that returned their questionnaires in 1999 were from the 1997 survey and 2) all the 388 firms on the must-take-all list in 1999 were from the same NAICS as firms in the 1997 survey. Thus, the 1997 list of firms and the 1999 must-take-all list are two samples from the same population. Consequently, they are representative samples of the same population.

Table 10: Share of the Change in the Leading Biotechnology Activity Indicators
Attributed to the Change in the Survey Methodology between 1997 and 1999

	The survey internounting, seeween 1997 and 1999
Indicators	Actual Impact
Number of innovative biotech firms	25
Biotech R&D	\$30 million
Biotech Revenues	\$121 million
Biotech employment	(119) employees <sup>(a)</sup>
Biotech Exports	\$37 million
Biotech products	779 products
Amount of capital raised	\$151 million

Source: Statistics Canada, 1997 Biotechnology Firm Survey and 1999 BUDS Note:

#### 6.2 1999 BUDS Vs 2001 BUDS

No methodology changes occurred between the last two surveys, i.e. 1999 and 2001 surveys, only the population surveyed in 2001 was larger and the "pre-contact" was done by mail. Like for the 1997 and 1999 surveys, the must-take-all list in 2001 came from the same NAICS as that of 1999. Thus, the two must-take-all lists are representative of the same population. Also all the NAICS codes of 1999 were surveyed in 2001 and represented 80% of the sample in the 2001 BUDS. The remaining 20% came from 4 new NAICS codes, namely, 31311, 3222, 5417, and 6215 (Table 1). The first NAICS code contributed nothing to the final count of innovative firms and the second contributed only one unit. In other words, the increase in the sample in 2001 came principally from the same population surveyed in 1999. Furthermore, this larger population size resulted in 17 additional firms, less than 5% of the estimated biotechnology firm population in 2001.

<sup>(</sup>a) Implies that the figure is negative

#### 6.3 Impacts of the Change in Survey Methodology on the Estimates

Given the aforementioned facts, changes in survey methodology resulted in 25 additional firms being captured in 1999. In other words, had the methodology remained unchanged between 1997 and 1999, the biotechnology firm population would have been underestimated by 25 firms or 7%. However, as shown in Table 10, the impact of this under-estimation on the key biotechnology activity indicators is small. Between 1999 and 2001, the larger sample size resulted in 17 additional firms, less than 5% of the 2001 biotech firm population in 2001.

#### VII Lessons Learned

#### 7.1 Defining biotechnology

Comments from respondents in the cognitive interviews during the questionnaire testing showed that the strategy consisting in using a double-definition of biotechnology is very helpful. Even though concise, the wording of the single definition was not clear to all the respondents. In fact, some respondents were not sure of the meaning or interpretation of key words such as science, living organisms or parts of living organisms, innovative manner. In contrast, the list based-definition which used specific biotechnologies grouped under different domains, was well understood. Second, by so doing, it allowed them to know whether or not their activities fell into the biotechnology domain. Otherwise, they could quickly exit the survey and save time. Thus, an additional advantage of the double-definition strategy was to minimize respondent burden.

#### 7.2 Defining an Innovative biotech firm

According to the Oslo Manual (OECD/EUROSTAT, 1997), a technological product/process (TPP) innovating firm is "one that has implemented technologically new or significantly technologically improved products or processes". A TPP innovation has been implemented if "it has been introduced on the market (product innovation) or used within a production process (process innovation)".

Given that most firms in the target population of the Biotechnology Use and Development surveys do not have any product on the market, and given Biotechnology is best thought as a set techniques which may be used by firms in their production processes, an innovative biotech firms has been defined along the lines of process innovation. This allows to capture the target population of the biotech surveys.

#### 7.3 Targeting the right population

Biotechnology is a cross-sectoral activity that cannot be classified under any single NAICS code. Two things were done to solve this problem. NAICS codes where the use of biotechnology was probable were selected. Then, within these NAICS, care was taken to insure that only innovative biotechnology firms, i.e. firms are actively involved in using biotechnologies to develop new biotechnology products and processes are captured.

To this end, a screening process was put in place. First and foremost, respondents have to be using at least one of a series of biotechnologies. Second, they have to be using biotechnologies to develop new products and processes. This latter condition implies that the firm has at least one product or process in development, in production, or on the market. Only respondents that fulfilled all these conditions were classified as innovative biotechnology firms.

Firms with biotech employees and using biotechnologies in their current production process for purposes other than developing new products and processes are classified as biotechnology users.

Using this screening scheme allows to target and survey the right population and to attain the main objective of the biotechnology survey, i.e. to collect information on firms that use biotechnology to develop products and processes.

#### 7.4 Measuring Biotechnology Employment

A large percentage of firms are involved in activities other than biotechnology. As a result, any given employee may be involved in multiple tasks, including those related to biotechnology. Consequently, more than the level of education, an employee's involvement in biotechnology activities better measures if he/she is classified as a full-time or part-time biotech employee or not. During the data collection, it was observed that the number of biotechnology employees in question 4b in the 2001 BUDS, was less than the sum of 4c and 4d (Appendix 4). And more often than not, this difference corresponded to the number of employees in production, finance, marketing and management. In other words, most respondents equate employment in biotechnology to research and clinical activities. This could potentially lead to an under-estimation of the number of employees dedicated to biotechnology. By using an activity-based accounting, we avoid this problem.

#### VIII Conclusion

From its beginning in 1996 with the "Survey of Biotechnology Use in Canadian Industries – 1996", to the 2001 Biotechnology Use and Development Survey, surveying biotechnology has evolved in many ways. First of all, the focus of the surveys has changed from investigating the use of biotechnologies to include the use of biotechnologies to develop new products and processes. Second, along this change in focus, have emerged new questions such as those related to intellectual property (IP) rights. Third, the definition of Biotechnology has been refined and international comparability, mainly between OECD countries, is now possible.

Measures of biotechnology activity have been developed. These allow for a better understanding of Biotechnology, not only as a set of techniques, but also as a means to develop new products and processes. NAICS codes where the use of Biotechnology is probable are now clearly identified.

The quality of the data is improving along common standards, allowing for comparability between the survey data and other data sources, be it national or international.

Whenever needed, appropriate adjustments to definitions and concepts were made to better capture the target population. For example, in selecting the respondents, the Gross Business Income (GBI) was used in conjunction with R&D expenditures and the number of employees. Likewise, innovative biotech firms were defined in terms of their use of biotechnologies to develop new products and processes (process innovation), rather than having new products on the market (product innovation).

Insights gained from the iteration of the biotech surveys have allowed a better understanding of both the extent and the intensity of biotechnology activities in Canada. The lessons learned will help improve upcoming surveys.

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# **Appendix 1: Biotechnology Firm Survey – 1997**



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### Biotechnology Firm Survey - 1997

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#### 3

#### Purpose

Statistics Canada is undertaking this survey under the sponsorship of Industry Canada and BIOTECanada to produce information of use to firms engaged in biotechnology activities by adressing the following question: What are the main characteristics of the firms which choose to develop biotechnologies as an important component of their business? The information from the survey can be used by businesses for market analysis, by trade associations to study performance and other characteristics of their industries, by government to develop national and regional economic policies, and by other users involved in research or policy making. Statistics Canada will create a database combining survey responses with existing Revenue Canada and Statistics Canada records.

#### Confidentiality

While participation in this survey is voluntary, your cooperation is important to ensure that the information collected in this survey is as accurate and as comprehensive as possible. Statistics Canada is prohibited by law from publishing or releasing, in any manner, any statistics which would divulge information obtained from this survey that relates to any identifiable business without the previous written consent of that business. The data reported on the survey questionnaire will be treated in strict confidence, used for statistical purposes and released in aggregated form only. The confidentiality provisions of the Statistics Act are not affected by either the Access to Information Act or any other Legislation.

#### **Biotechnology Definition**

Biotechnology is defined as the application of science and engineering in the direct or indirect use of living organisms or parts of organisms in their natural or modified forms in an innovative manner in the production of goods and services or to improve existing processes.

Please report only on Canadian biotechnology activities of your company. Complete a separate questionnaire for each company engaged in biotechnology activities in Canada.

#### Questions?

If you require assistance in the completion of this questionnaire or have any questions regarding this survey please contact Lloyd Lizotte (tel: 613-951-2188 (call collect), fax: 613-951-9920 or e-mail: lizollo@statcan.ca)

#### **Survey Contact**

Please indicate the name of the person completing this form so we know who to contact should we have questions about this report.

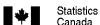
Name	Title
Telephone Number	Fax Number

Combined Report			
If your records do not permit separate report			

 If your records do not permit separate reporting, list the names of other companies included in this report and indicate whether they are engaged in biotechnology activities by writing YES or NO in the second column.

Name of company	Biotechnology activities YES or NO	Indicate type of affiliation with reporting company (i.e. parent, subsidiary or other)
	¹ Yes ² No	

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istics Statistique ada Canada



	0 "	IF "YES" ➤	Principal Use <i>(ch</i>	IF "NO"		
Biotechnologies	Currently Used in Operations?	Products/ Processes Development	Clinical/ Field Trials	Current Production	Do you PLAN TO USE within 3 years?	
DNA Based	¹ Yes	1	2	3		
Genetic Engineering	<sup>2</sup> No )	-		_	→ ¹○ Yes ²○ No	
	¹() Yes )	1 ( )	2()	3 ( )		
Gene Probes	<sup>2</sup> No )	-			→ ¹○ Yes ²○ No	
	1 Yes	1 ( )	2 ( )	3 ( )		
Bio-informatics / Genomics / Pharmacogenetics	2	Ü			→ ¹○ Yes ²○ No	
	1 Voc. 1	1 (	2 (	3 (		
DNA Sequencing / Synthesis / Amplification	<sup>2</sup> No )				→ ¹○ Yes ²○ No	
		1 (	2 (	3 (	<u>,                                     </u>	
Gene Therapy			-0		→ ¹○ Yes ²○ No	
Biochemistry / Immunochemistry Based					<u> </u>	
Vaccines / Immune Stimulants / Drug		1	2	3		
Design & Delivery / Combinatorial Chemistry	<sup>2</sup> No )	-			→ ¹ Yes ² No	
	¹ Yes >	1	2	3		
Diagnostic Tests / Antibodies	<sup>2</sup> No )				→ ¹○ Yes ²○ No	
	¹ Yes	1	2	3		
Peptide / Protein Sequencing or Synthesis	<sup>2</sup> No )	-			→ ¹○ Yes ²○ No	
Cell Receptors / Cell Signalling /	¹ Yes	1	2	3		
Pheromones / Three Dimensional Molecular Modelling / Structural Biology	<sup>2</sup> No )	-			→ ¹○ Yes ²○ No	
	¹() Yes	1 ( )	2()	3 ( )		
Biosensors	~			_	→ ¹○ Yes ²○ No	
	1 ( Yes )	1 ( )	2 ( )	3 ( )		
Biomaterials		_		0	→ ¹○ Yes ²○ No	
			2 (	3 (		
Microbiology / Virology / Microbial Ecology		_		$\overline{}$	→ ¹○ Yes ²○ No	
Bioprocessing Based		_	2 (		7 0 100 0 110	
•		_	2	<u> </u>	► 1 O Voc. 2 O No.	
Cell / Tissue / Embryo Culture & Manipulation			200		Tes O NO	
Fermentation / Bioprocessing / Biotransformation / Bioleaching / Biopulping /		_	2		> 1 ○ V 2 ○ N-	
Biobleaching / Biodesulphurization					Yes 2 NO	
	$\tilde{}$		2	3	20	
Extraction / Purification / Separation	<sup>2</sup> No )	-			→ ¹	
Bioremediation / Phytoremediation /	Currently					
Biofiltration / Bioindicators	Current   Do you PLAN TO USE   Processors   Clinical   Production   Do you PLAN TO USE within 3 years?					
	¹ Yes	1 )	2	3		
Natural Products Chemistry	<sup>2</sup> No )	-			→ ¹○ Yes ²○ No	
Others (Please specify)	¹ Yes	1	2	3		
	<sup>2</sup> No )	<b>-</b>			→ ¹○ Yes ²○ No	
	¹() Yes	1 ( )	2	3		
	$\tilde{}$	_			→ ¹○ Yes ²○ No	
1	1 Yes	1 ( )	2()	3 ( )		
	2	_		0	→ ¹○ Yes ²○ No	

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#### **BIO-INDUSTRY SECTOR**

3. a) Please indicate the top 3 bio-industry sectors, where applicable, in numbered order, that best describe your firm's biotechnology activities, (use 1 for the Most important and 3 for the least important). Please also indicate the number of products based on genetic engineering (rDNA) your company has in each of the relevant development stages:

#### Human Health - Bio

- Diagnostics (e.g. immunodiagnostics, gene probes, biosensors)
- Therapeutics (e.g. vaccines, immune stimulants, biopharmaceuticals, rational drug design, drug delivery, combinatorial chemistry)
- 3) **Gene Therapy** (e.g. gene identification, gene constructs, gene delivery)

#### **Bio-Informatics**

 Genomics and Molecular Modelling (e.g. DNA/RNA/protein sequencing & databases for humans,plants, animals and microorganisms)

#### Ag - Bio

- Plant Biotechnology (e.g. tissue culture, embryogenesis, genetic markers, genetic engineering)
- 6) **Animal Biotechnology** (e.g. diagnostics, therapeutics, embryo transplantation, genetic markers, genetic engineering)
- 7) Biofertilizers/Biopesticides/Biological Feed Additives/Microbial pest control (e.g. bacteria, fungi, yeasts)
- Non-Food Applications of Agricultural Products (e.g. fuels, lubricants,commodity and fine chemical feedstocks, cosmetics)

#### **Food Processing**

- 9) Bioprocessing
  - (e.g. using enzymes and bacteria culture)
- 10) Functional Foods/Nutriceuticals (e.g. probiotics, unsaturated fatty acids)

#### Aquaculture

- 11) Fish health (e.g. diagnostics, therapeutics)
- Broodstock genetics (e.g. tracking superior traits, genetic modification / engineering)
- 13) **Bioextraction** (e.g. karageenan from seaweed, antifreeze proteins from fish, flavours)

#### Mining/Energy/Petroleum/Chemicals

- 14) Microbiologically enhanced petroleum/mineral recovery
- (Cleaner) Industrial Bioprocessing(e.g. biodesulphurization, bio-cracking, bio-recovery)

#### **Forest Products**

- 16) **Silviculture** (e.g. ectomycorrhizae, tissue culture, somatic embryogenesis, genetic markers, genetic engineering)
- 17) (Cleaner) Industrial Bioprocessing (e.g. biopulping, biobleaching, biological prevention of sapstain)

#### **Environment**

- 18) **Biofiltration** (e.g. treatment of organic emissions to air/water)
- Bioremediation and Phytoremediation (e.g. cleanup of toxic waste sites using microorganisms)
- 20) **Diagnostics** (e.g. detection of toxic substances using bioindicators, biosensors, immunodiagnostics)

#### Other

- 21) Custom synthesis- chemical or biological (e.g. peptides, proteins, nucleotides, hormones, growth factors, biochemicals
- 22) Other (please specify)

	Number of	products by developme	ent stages
Top Bio-sectors (1 to 3 by importance) Please write number and sub-headings from above list	Approved/ On Market	Clinical/Field Trial stage	Under development
	(number)	(number)	(number)
1			
2			
3			
b) Did you implement a new or significantly improved PROCESS of biotechnologies?	<b>3</b> in the last 3 year	rs that required th	e development
1  Yes ➤ How many?  2 No			
c) Do you PLAN to implement in the NEXT 3 years a new or s development of biotechnologies?	significantly improv	ed <b>PROCESS</b> th	at requires the
Number  1  Yes ➤ How many?  2  No			

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Financial Inform	nation					
4. Total Business A	Activity (All	activit	ies including	Biotech)		
a) Is your firm a pub	olic company?	1 (	Yes			
		2 (	○ No			
b) Please report da Do not include sa	ta for 1997 or ales and opera	the late	st fiscal year av f your <b>subsidia</b>	railable. ries located outsic	le Canada.	
Fiscal year (if different from	1997)					
Balance Sheet	1997		Revenue	1997	Expenses	1997
Cash & securities	000	\$ Pro	oduct sales	000\$	R&D	000\$
Total Assets	000	\$ Co	ntract Research	000\$	All other expens	ses 000\$
Total liabilities	000	\$ Oth	ner revenue	000\$		
Shareholders Equity	000	\$				
TOTAL Exports (as % of product sales)			%	Total exports to the		%
Biotechnology Related (as % of product sales)			%	Total exports to Eu (as % of product s		%
				Total exports to As (as % of product s		%
				Total exports to So (as % of product s		%
				Total exports to ot (as % of product s		%
Total number of employ (average for the year):	yees					
What percentage of you are based on biotechno			C	% OR bio	product sales	000\$
What percentage of you are based on biotechno		es	C	% OR bio	) R&D	000\$
R&D						
5. a) Did your firm hav	/e biotech R&I	) expen	ditures			
in any of the yea			1 (	Yes No		

5. a) Did your firm have biotech R&D expenditures in any of the years 1995-1997?

1 Yes
2 No

b) In the past 5 years, did your company apply for the tax benefit under the R&D (SR&ED) tax programme?

1 Yes
2 No

What was the most recent year?

2 No

Why did your company not apply? (check most important only)

1 complexity of the application process
2 uncertainty of eligibility
3 did not meet eligibility requirements
4 other (specify)

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			For	eign Partner, (	Please identif	y country of partn	er)
trategic Alliance for:		Canadian Partner	USA	E-U	Asia	South/ Latin America	Othe
larketing/Distribution	¹	1	2	3	4	5	6
lanufacturing	¹	1	2	3	4	5	6
egulatory Affairs	¹	1	2	3	4	5	6
inance	¹	1	2	3	4	5	6
ther	¹	1	2	3	4	5	6
<b>&amp;D Partnership:</b> Biotech Company	¹	1 🔵	2	3	4	5	6
Other company	¹	1	2	3	4	5	6
University	¹○ Yes ➤ ²○ No	1	2	3	4	5	6
Hospital	¹○ Yes ➤ ²○ No	1	2	3	4	5	6
Research Institute	¹	1	2	3	4	5	6
Federal Lab (Canadian)	¹ Yes ² No						
Provincial Lab (Canadian)	¹ Yes ² No						
Network of Centres of Excellence (Canadian)	¹ Yes ² No						
<ul> <li>a) In your current biotechnology (include contract personnel a</li> </ul>	y activities, pleas	e give the nu	mber of e	mployees	on staff in ling more	the listed potential	ositior e).
			<u> </u>			Estimated N	

Position	Number Now Employed	Positions Unfilled Now	Estimated Number Employed in year 2001
R&D			
Clinical Affairs/Quality Assurance			
Regulatory/Legal/Government Affairs			
Manufacturing			
Marketing & Sales			
Business Development/Finance			
Administration/Human Resources			

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Hullian Kesourc	es engagea in Biotec	chnology Activ	ities (continued	ı
7. b) Does your firm h	ave a formal employee devel	opment program (con	tinuing education and t	raining program)?
¹ Yes ➤ If fo	yes, estimate your firm's tota ormal education and training \$000) or as a percentage of p	al expenditures on in 1997? oroduct sales	\$000	OR %
c) Does your firm e	mploy co-op program studen	ts from universities?		
¹ Yes				
<sup>2</sup> No				
d) Do you have a fu	ıll time person solely respons	sible for Human Resou	urces in your organizati	on?
<sup>1</sup> Yes				
<sup>2</sup> No				
e) What is your ap biotechnology ac	oproximate professional staf ctivity in 1997?	ff turnover rate (pers	ons leaving as a %	of total staff) for the
f) Are you experien	ncing problems in recruiting B	Susiness Operations st	taff?	
1 Ves ➤ If	YES. In which specialties?	<sup>1</sup> Finance		
<sup>2</sup> No	TEO. III WIIIOII Specialitico:	<sup>2</sup> Marketing		
<u> </u>		<sup>3</sup> Regulatory, Le	egal	
		<sup>4</sup> Clinical Affairs		
g) Are you experien	ncing problems in recruiting T	echnical / Production	/ Scientific / R&D staff?	)
¹○ Yes ➤ If	YES. In which specialties?	<sup>1</sup> Scientist		
<sup>2</sup> No		<sup>2</sup> Engineering		
-		<sup>3</sup> Technicians		
h) Have you tried to	hire personnel from outside	Canada in 1997?		
¹○ Yes		<sup>2</sup> No		
If Yes, From v	vhich areas?			
¹ US				
<sup>2</sup> EU				
<sup>3</sup> Asia				
	tin America			
<sup>5</sup> Other   ▼				
If Yes, were	you successful?			
¹ Yes				
<sup>2</sup> ○ No ➤	If No, Was the problem rel to the following issues?	lated		
	<sup>1</sup> Personal income taxe	es		
	<sup>2</sup> Immigration Rules			
	3 Advancement opportu	unities		
	<sup>4</sup> Other			

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8. a) Have you ever had to abandon an important biotech project because further work was blocked by IP rights held by another organization?	of pate	his an issue of so ent in Canada as ared to other cou	s ·	
b) Has your firm been involved in litigation related to patent infringement in the past year?	¹ Yes ➤ How m ² No	nany different ca	ises?	
c) During the last three years has your firm assigned the right to use		In Canada	Outside Canada	
intellectual property <b>TO</b> :  (check where applicable)	Another firm	1	2	
c) During the last three years has your firm assigned the right to use intellectual property TO: (check where applicable)  d) During the last three years has your firm acquired the right to use intellectual property FROM: (check where applicable)  roblems for Biotechnology Com  Select the three issues that you consider biotechnology products/processes:  101  Access to capital  20  Access to smart capital	Government lab	3	4	
	University	5	6	
	Hospital	7	8	
		In Canada	Outside Canada	
intellectual property <b>FROM</b> :	Another firm	1	2	
(check where applicable)	Government lab	3	4	
	University	5	6	
	Hospital	7	8	
<ul><li>9. Select the three issues that you consider a biotechnology products/processes:</li></ul>			ccessfully commercia	lize your
O1 Access to capital				
Access to smart capital (money plus management expertise	∍)			
Access to technology				
Skilled human resources				
Consumer acceptance				
Lack of information about markets				
Regulations: 07 Labeling				
Limited internation	onal harmonization			
<sup>09</sup> IP protection				
Cost for gaining i	regulatory approval			
Time required to	gain regulatory approve	al		

Intellectual property (IP) instruments

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Other (please specify)

10		000¢	10		000¢
¹ Yes ➤ How Much? L		000\$	¹○ Yes ➤ How Much? [		000\$
<sup>2</sup> No			<sup>2</sup> No		
For what purposes? (check most important)			For what purposes? (check most important)		
¹○ R&D			¹○ R&D		
<sup>2</sup> Regulatory approvals			<sup>2</sup> Regulatory approvals		
<sup>3</sup> Process Scaleup			<sup>3</sup> Process Scaleup		
<sup>4</sup> Other			<sup>4</sup> Other		
Sources? (check most important)			Sources? (check most important)		
<sup>1</sup> Friends / "Angel Investor	s"		<sup>1</sup> Friends / "Angel Investo	ors"	
<sup>2</sup> Venture Capital / Labour	Sponsored	Funds	<sup>2</sup> Venture Capital / Labou	ur Sponsored Fu	ınds
<sup>3</sup> Private Placement			<sup>3</sup> Private Placement		
<sup>4</sup> IPO (Initial Public Offerin	g)		<sup>4</sup> O IPO (Initial Public Offer	ing)	
<sup>5</sup> Secondary Public Offerin	g		<sup>5</sup> Secondary Public Offer	ring	
	r		<sup>6</sup> Strategic Alliance Partr	ner	
<sup>6</sup> Strategic Alliance Partne					
Other  rategic Decisions e following question should be		-	O of your company.  Outher	or this year (1998	8)?
Other  rategic Decisions e following question should be	important d	ecisions y	O of your company.		•
Other  rategic Decisions  e following question should be  As the CEO, what were the most (check up to three for each year)	important d	-	O of your company.	or this year (1998 	•
Other  rategic Decisions  e following question should be  As the CEO, what were the most	important d	ecisions y	O of your company.		_1
Other  rategic Decisions  e following question should be  As the CEO, what were the most (check up to three for each year)  Refocused current product	<u>1997</u>	ecisions y	O of your company. ou made over last year (1997)? Fo	<u>1997</u>	
Other  rategic Decisions  e following question should be  As the CEO, what were the most (check up to three for each year)  Refocused current product development	1997 01 0	ecisions y	O of your company.  ou made over last year (1997)? Fo	1997	8)? 
Other  rategic Decisions  e following question should be  As the CEO, what were the most (check up to three for each year)  Refocused current product development  Downsized the organization  Increased the size of the organization	1997 01 0	ecisions ye	O of your company.  ou made over last year (1997)? For the company of the company.  Licenced out technology  Alliances with academia  Alliances with government	1997 19 <u>21</u>	20 22 24
Other  rategic Decisions  e following question should be  As the CEO, what were the most (check up to three for each year)  Refocused current product development  Downsized the organization	1997 01 0 03 0 05 0	ecisions ye  1998 02 04 06 08	O of your company.  ou made over last year (1997)? For Licenced out technology  Alliances with academia	1997 19 0 21 0 23 0	20 22 24 26
Other  rategic Decisions  e following question should be  As the CEO, what were the most (check up to three for each year)  Refocused current product development  Downsized the organization  Increased the size of the organization	1997 01 0 03 0	ecisions ye  1998 02 04 06 06	O of your company.  ou made over last year (1997)? For the company of the company.  Licenced out technology  Alliances with academia  Alliances with government	1997 19 0 21 0	20 22 24 26
Other  Prategic Decisions  e following question should be  As the CEO, what were the most (check up to three for each year)  Refocused current product development  Downsized the organization  Increased the size of the organization  Entered product trials	1997 01 0 03 0 05 0	ecisions ye  1998 02 04 06 08	O of your company. ou made over last year (1997)? Fo  Licenced out technology  Alliances with academia  Alliances with government  Alliances with companies	1997 19 0 21 0 23 0	20
rategic Decisions e following question should be As the CEO, what were the most (check up to three for each year) Refocused current product development  Downsized the organization  Increased the size of the organization  Entered product trials  Launched new product	1997 01 0 03 0 05 0 09 0	ecisions ye  1998 02 04 06 08 10	O of your company. ou made over last year (1997)? For Licenced out technology  Alliances with academia  Alliances with government  Alliances with companies  Raised private capital	1997 19 21 23 25 27 27	20 22 24 26 28
rategic Decisions e following question should be As the CEO, what were the most (check up to three for each year) Refocused current product development  Downsized the organization  Increased the size of the organization  Entered product trials  Launched new product  Acquired a company	1997 01 0 03 0 05 0 09 0	ecisions ye  1998 02 04 06 08 10 12	O of your company.  ou made over last year (1997)? For Licenced out technology  Alliances with academia  Alliances with government  Alliances with companies  Raised private capital  Raised public capital	1997 19 21 23 25 27 29 0	1 20 22 24 26 28 30
rategic Decisions e following question should be As the CEO, what were the most (check up to three for each year) Refocused current product development  Downsized the organization  Increased the size of the organization  Entered product trials  Launched new product  Acquired a company  Acquired by a company	1997 01 0 03 0 05 0 09 0 11 0	ecisions ye  1998 02 04 06 08 10 12 14	O of your company.  ou made over last year (1997)? For Licenced out technology  Alliances with academia  Alliances with government  Alliances with companies  Raised private capital  Raised \$ from alliance	1997 19 21 23 25 27 29 31 0	1 20 22 24 26 28 30 32

Comments			
			<del></del>

Thank you for your cooperation!

### **Appendix 2: Biotechnology Use and Development Survey – 1999**



## Biotechnology Use and Development Survey - 1999

#### Confidential when completed

Collected under the authority of the Statistics Act, Revised Statutes of Canada, 1985, c. S-19.

Completion of the questionnaire is a legal requirement under the Statistics

Si vous préférez ce questionnaire	
en français, veuillez cocher	



#### **Survey Purpose**

Statistics Canada is undertaking this survey in support of the Canadian Biotechnology Strategy. The purpose is to produce information about firms engaged in biotechnology activities by addressing the following question. What are the characteristics and activities of firms that use or develop biotechnology as an important part of their firm's activity?

Biotechnology is a dynamic emerging sector of the Canadian economy and its impact has the potential to be felt through all parts of Canadian society. An accurate understanding of biotechnology requires comprehensive data. Information from this survey may be used by businesses for economic or market analysis, by trade associations to study industry performance, government departments and agencies to assist policy formation, and the academic community for research purposes. Statistics Canada will create a database combining survey responses with existing Statistics Canada data records. An executive summary of the results will be sent to all respondents.

Please report on Canadian biotechnology activities of your firm. Complete a separate questionnaire for each firm engaged in biotechnology activity in Canada.

#### **Authority**

Collected under the authority of the Statistics Act, Revised Statutes of Canada, Chapter S19. Completion of this questionnaire is a legal requirement under the Statistics Act.

#### Confidentiality

Statistics Canada is prohibited from publishing or releasing any statistics that would divulge information obtained from this survey that relates to any identifiable firm without the previous written consent of that firm. The data reported in this questionnaire will be treated in strict confidence, used for statistical purposes and released in aggregate form only. The confidentiality provisions of the Statistics Act are not affected by either the Access to Information Act or any other Legislation.

If you require assistance in the completion of the questionnaire or have any questions regarding the survey, please contact:

Claire Racine-Lebel
Science, Innovation and Electronic Information Division
Statistics Canada
Tunney's Pasture
Ottawa, Ontario
K1A 0T6

Phone: (613) 951-6309 (please call collect) - Fax: (613) 951-9920 e-mail: Claire.Racine-Lebel@statcan.ca

Please indicate the name of the person completing this form so we kn	ow who to contact should we have questions about this report.
Name	Title
Telephone Number	Email
Fax Number	

5-4900-500.1: 2000-01-13 STC/SAT-430-75177





		If currently us	ing, do you	use them for	Number	If N	lo
Biotechnologies	Currently Used in Operations	Development	Current Production	Environmental Purposes	of Years in Use	Do you use withir	
DNA Based	0	1	2	3	4	5	
Gene Probes/DNA Markers		<b>-</b>	0	0		Yes	
Bio-Informatics	Yes -	<b>→</b> ○	$\bigcirc$	0		Yes	
Genomics/Pharmacogenetics		<b>-</b>	$\bigcirc$	$\bigcirc$		Yes	
Genetic Engineering/DNA Sequencing/Synthesis/Amplification	Yes -	<b>→</b> ○	$\bigcirc$	0		Yes	
Biochemistry/Immunochemistry							
Vaccines/Immune Stimulants		<b>-</b>	0	$\circ$		Yes	$\subset$
Drug Design & Delivery	Yes ■ No ■	<b>-</b>	0	0	<b>&gt;</b>	Yes	
Diagnostic Tests/Antibodies	Yes ■ No ■	<b>-</b>	<u> </u>	0	<b>_</b>	Yes	
Peptide/Protein Sequencing/ Synthesis	Yes -	<b>-</b>		<u> </u>	-	Yes	
Cell Receptors/Signalling/ Pheromones/Structural Biology	Yes -			<u> </u>	-	Yes	
Combinatorial Chemistry/ 3D Molecular Modelling	Yes -			<u> </u>	<b>—</b>	Yes	
Biomaterials	Yes -	<b>→</b> ○	<u> </u>	<u> </u>	-	Yes	
Microbiology/Virology/Microbial Ecology  Bioprocessing Based	Yes -	<b>→</b> ○	<u> </u>	<u> </u>	<u> </u>	Yes	<u></u>
Cell/Tissue/Embryo Culture Manipulation		<b>-</b>		0		Yes	
Extraction/Purification/Separation		<b></b> 0	$\bigcirc$	$\circ$		Yes	
Fermentation/Bioprocessing/ Biotransformation/Natural Products Chemistry		<b>-</b>	0	0		Yes	
Environment							
Bioleaching/Biopulping/Biobleaching/ Biodesulphurization	Yes ■ No ■	<b>-</b>	0	0	-	Yes	$\subset$
Bioremediation/Biofiltration/ Phytoremediation	Yes ■ No ■	<b>-</b>	0	0	<b>—</b>	Yes	
Other (please specify)							
	Yes -	<b>-</b>	$\bigcirc$	$\bigcirc$		Yes	
	Yes -	<b>-</b>	$\bigcirc$	$\circ$		Yes	
						- U 168	

				nportanc	your firr	n. 	Not
		Low 1	2	3	4	High 5	Applicable 0
	Lack of Financial Justification						
2100	Small market size	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
2110	High cost of equipment	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
2120	High cost to implement/integrate biotechnology	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
2130	Cost of capital	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
	Human Resources						
2140	Shortage of skilled or trained staff	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
2150	Worker resistance	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
2160	Increased labour costs	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
	External						
2170	Government regulations	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
2180	Public acceptance/perception of biotechnology	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
	Technology						
2190	Biotechnology not sufficiently developed	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
2200	Lack of external technical expertise/support	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
	Other (please specify)						
2210		$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	

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		In	nportanc	е		Not
Sources of Information on Biotechnology	Low 1	2	3	4	High 5 <del>→</del>	Applicab 0
Internal resources/staff or parent/subsidiary firm	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Academic journals/trade publications	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Universities/colleges/private training institutes	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Federal government department/agency		$\bigcirc$		$\bigcirc$	$\bigcirc$	
Personal contact with others (tacit knowledge)	$\bigcap$	$\bigcirc$		$\bigcirc$	$\bigcirc$	
Other companies	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Provincial government department/agency	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Professional/industry associations	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Library/literature search	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Database retrieval services	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Conferences/workshops/trade shows		$\overline{}$	$\overline{}$	$\overline{}$	$\bigcirc$	
Other (please specify)						
V 1 7/						
<ul> <li>No → Go to Question 5.</li> <li>Yes</li> <li>Rate the benefits from using biotechnologies in your firm</li> <li>1 is low importance and 5 is high importance. Indicate if</li> </ul>	•	or proces	• .	rations. I	Jse the follo	wing scale wh
○ Yes ↓	m's production o	or proces to your fi	rm.		Jse the follo	
Yes  Rate the benefits from using biotechnologies in your firm	m's production o	or proces to your fi	• .		Jse the follo High 5	Not
Yes  Rate the benefits from using biotechnologies in your firr 1 is low importance and 5 is high importance. Indicate if	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Yes  Rate the benefits from using biotechnologies in your firr 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Yes  Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Yes  Rate the benefits from using biotechnologies in your firr 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Yes  Rate the benefits from using biotechnologies in your firr 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applicat
Yes  Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement  Develop new products or processes	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applicat
Productivity Improvement  Lower labour costs  Lower energy costs  Product Improvement  Develop new product range	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Productivity Improvement Lower labour costs Lower energy costs  Product Improvement Develop new product quality  Yes  Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement Lower labour costs  Lower capital costs  Extend product sor processes	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Productivity Improvement Lower labour costs Lower energy costs Product Improvement Develop new products or processes Extend product range Improvement in product quality Plant Organization	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement  Develop new products or processes  Extend product range  Improvement in product quality  Plant Organization  Increase production flexibility	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement  Develop new products or processes  Extend product range  Improvement in product quality  Plant Organization  Increase production flexibility  Lower maintenance expenses	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement  Develop new products or processes  Extend product range  Improvement in product quality  Plant Organization  Increase production flexibility  Lower maintenance expenses  Cleaner production/pollution reduction	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement  Develop new products or processes  Extend product range  Improvement in product quality  Plant Organization  Increase production flexibility  Lower maintenance expenses  Cleaner production/pollution reduction  Market Performance	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applical
Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement  Develop new products or processes  Extend product range  Improvement in product quality  Plant Organization  Increase production flexibility  Lower maintenance expenses  Cleaner production/pollution reduction  Market Performance  Improve market position	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applicat
Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if 1 is low importance and 5 is high importance. Indicate if 1 is low importance and 5 is high importance. Indicate if 1 is low importance and 5 is high importance. Indicate if 1 is low importance and 5 is high importance. Indicate if 1 is low importance in productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement  Develop new products or processes  Extend product range  Improvement in product quality  Plant Organization  Increase production flexibility  Lower maintenance expenses  Cleaner production/pollution reduction  Market Performance  Improve market position  Increase sales	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applicat
Rate the benefits from using biotechnologies in your firm 1 is low importance and 5 is high importance. Indicate if  Benefit of Using Biotechnology  Productivity Improvement  Lower labour costs  Lower capital costs  Lower energy costs  Product Improvement  Develop new products or processes  Extend product range  Improvement in product quality  Plant Organization  Increase production flexibility  Lower maintenance expenses  Cleaner production/pollution reduction  Market Performance  Improve market position	m's production of f not applicable Low	or proces to your fi	nportanc	e	High	Not Applicat

	e purposes of this survey Employees are defined as atement of Remuneration Paid Form for the 1999 tax					
a)	How many employees does your firm currently employ?	? 5100				
b)	How many employees have biotechnology-related resp	oonsibilities?	5110			
	In the table below provide the number of biotechnology. For example, a person working 60% of their time on scientific/research direction.					
	Position	_	Numbe	er Currently Em	ployed	
			Working full- time on biotechnology (more than 50% of time)	Working part time on biotechnology (less than 50% of time)	Estimated number to be employed in biotechnology in 2002	
	Biotechnology R&D Activities					
5120	Scientific/Research Direction					
5130	Technicians/Engineering					
5140	Regulatory/Clinical Affairs					
	Biotechnology Administration & Production					
5150	Production					
5160	Finance/Marketing					
5170	Management/Licensing/Administration					
d) 518	Does your firm currently have unfilled full time biotechn  80	ology-related positions?				
			If Yes, was the reason due to			
		Г		0000000	Т	
	Position	Number of Unfilled Full-Time Positions	Lack of qualified candidates	Compensation required by qualified candidated too high	Other 4	
	Position  Biotechnology R&D Activities	Unfilled Full-Time Positions	qualified candidates	tion required by qualified candidated too high		
519	Biotechnology R&D Activities	Unfilled Full-Time Positions	qualified candidates	tion required by qualified candidated too high		
51 <u>9</u>	Biotechnology R&D Activities  Scientific/Research Direction	Unfilled Full-Time Positions	qualified candidates	tion required by qualified candidated too high 3	4	
	Biotechnology R&D Activities  Scientific/Research Direction  Technicians/Engineering	Unfilled Full-Time Positions	qualified candidates	tion required by qualified candidated too high 3	4	
520	Biotechnology R&D Activities  Scientific/Research Direction  Technicians/Engineering	Unfilled Full-Time Positions	qualified candidates	tion required by qualified candidated too high 3	4	
520	Biotechnology R&D Activities  Scientific/Research Direction  Technicians/Engineering  Regulatory/Clinical Affairs  Biotechnology Administration & Production	Unfilled Full-Time Positions	qualified candidates	tion required by qualified candidated too high 3	4	
520 521	Biotechnology R&D Activities  Scientific/Research Direction  Technicians/Engineering Regulatory/Clinical Affairs  Biotechnology Administration & Production  Production	Unfilled Full-Time Positions	qualified candidates	tion required by qualified candidated too high 3	4	

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5250	No → Go to Question 5 f)					
		Unde	nical/Trad ergraduate uate level	e/College level		
) Doe	es your firm contract out any of the following biotechnology-	related a	activities?			
_	Biotechnology Activity		No 0	Yes	(in \$000) of cor If more than the tota	is the value ntracts in 1999 n one what is al value?
5260 I	Research & Development		Ö	$\bigcirc$ $\rightarrow$	\$	,000
5270 I	Regulatory/Clinical Affairs			$\bigcirc$ $\rightarrow$	\$	,000
5280	Marketing/Distribution			$\bigcirc$ $\rightarrow$	\$	,000
5290	Management/Licensing/Administration			$\bigcirc$	\$	,000
_						
	4:u.u. Duostissa					
	ting Practices  any of the following methods used to fill biotechnology-relat	ed nocit	ione			
6000	1 Internet resources 7	_		alified staff		
	2 University recruitment 8		e over-qu tworking	aiiileu Staii		
	<sup>3</sup> Use under-qualified staff <sup>9</sup>		_	ournal ads		
	4 Temporary/contract staff			associations		
	5 Employment agencies 11					
	6 In-house training	<i>)</i> Oii	nei ( <i>piea</i> s	e specify)		
	notice training					
a) Did	I you attempt to hire biotechnology staff from outside Canad	a in 199	9?			
6100	No → Go to Question 7 c)					
	Yes → From where? → 1 USA	4 (	Latir	n America		
	<sup>2</sup> Europe	5(	Othe	er		
	<sup>3</sup> Asia					
o) We	ere you successful in hiring biotechnology staff from outside	Canada	?			
6120	○ No					
	Yes → How many biotechnology staff did you hire	from ou	tside Can	ada in 1999?	1	ı
c) Did	I biotechnology personnel leave your firm in 1999?					
6130	○ No					
	Yes → How many?					
roduc	ct/Process Development					
	ct/Process Development your firm currently developing product that requires the use	e of biote	echnologie	es?		
		e of biote	echnologie	es?		
a) Isy	your firm currently <b>developing</b> product that <b>requires</b> the use	e of biote	echnologie	es?		

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8.	b) Is your firm currently <b>developing</b> processes that requi	res the use of	biotechnologies?							
	7110 Yes									
	○ No									
-	c) Does your firm consider biotechnology central to its ac	ctivities?								
	7120 Yes									
	○ No									
-	Did you answer "Yes" to any part of Question 8?									
	7130									
	○ No → Please stop here. Return the ques	tionnaire in the	prepaid return envelo	pe. Thank you for your	cooperation.					
8 E	iotechnology Products									
9.	Please provide the <b>number</b> of biotechnology products or	processes you	r firm has at each stag	ge of development.						
		Numb	per of biotechnolog	gy products/process	ses by					
	Biotechnology Sector		develop	ment stage						
		Research & Development	Pre-clinical trials/ Confined field trials	Regulatory phase/ Unconfined release assessment	Approved/ On market/In production					
	Human Health	0	1	2	3					
8110										
0400	gene probes)									
8120	<b>Therapeutics</b> (e.g. vaccines, immune stimulants, biopharmaceuticals, rational drug design, drug delivery, combinatorial chemistry)									
	Agriculture Biotechnology									
8130	Plant Biotechnology (e.g. tissue culture, embryogenesis, genetic markers, genetic engineering)									
8140	<b>Animal Biotechnology</b> (e.g. diagnostics, therapeutics, embryo transplantation, genetic markers, genetic engineering)									
8150	Non-food Agriculture (e.g. fuels, lubricants, commodity and fine chemical feedstocks, cosmetics)									
	Natural Resources									
8160	<b>Energy</b> (e.g. microbiologically enhanced petroleum recovery, industrial bioprocessing, biodesulphurization)									
8170	<b>Mining</b> (e.g. microbiologically enhanced mineral recovery, industrial bioprocessing, biodesulphurization)									
8180	Forest Products (e.g. biopulping, biobleaching, biopesticides, tree biotechnology, industrial bioprocessing)									
	Environment									
8190	Air (e.g. bioremediation, diagnostics, phytoremediation, biofiltration)									
8200	phytoremediation)									
8210	<b>Soil</b> (e.g. biofiltration, diagnostics, bioremediation, phytoremediation)									

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-			Number of biotechnology products/processes by development stage						
	Biotechn	ology Sector	Research & Development	Pre-clinical trials/ Confined field trials	Regulatory phase/ Unconfined release assessment	Approved/ On market/In production			
	Aquaculture			<u>.</u>					
8220	Fish health, broodstoc	k genetics, bioextraction							
	BioInformatics								
		ar modelling (e.g. DNA/RNA/ databases for humans, plants, ganisms)							
8240 ·	Gene therapy (e.g. go constructs, gene delive	ene identification, gene ery)							
	Food Processing								
	<b>Bioprocessing</b> (e.g. ι culture)	using enzymes and bacteria							
	Functional Foods/Nu unsaturated fatty acids	traceuticals (e.g. probiotics,							
	Other (please specify	<i>(</i> )							
8270									
0270									
8280									
8280 -									
8280 	-	rative Arrangements							
8280 Co	Was your firm involved n 1999?  Cooperative and colla or organizations in order	rative Arrangements in biotechnology-related coopera aborative arrangements involve to develop and/or continue work contracting-out is not regarded as	the active partic k on new or sign	ipation in projects by	/ your company and othe	er companies			
8280	Was your firm involved n 1999? Cooperative and collabor organizations in orderand/or services. Pure of	in biotechnology-related cooperative arrangements involve for to develop and/or continue work	the active partic k on new or sign	ipation in projects by	/ your company and othe	er companies			
8280	Was your firm involved n 1999? Cooperative and collabor organizations in order and/or services. Pure €	in biotechnology-related <b>coopera aborative arrangements</b> involve the state of t	the active partic k on new or sign	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Was your firm involved n 1999?  Cooperative and collabor organizations in order and/or services. Pure €  P100 No → €  Yes → I	in biotechnology-related <b>coopera aborative arrangements</b> involve to develop and/or continue work contracting-out is not regarded as  Go to question 13	the active partic k on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Was your firm involved n 1999?  Cooperative and collabor organizations in order and/or services. Pure €  P100 No → €  Yes → I	in biotechnology-related <b>coopera aborative arrangements</b> involve to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   1  ch purposes. Check any that are a	the active partic k on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Was your firm involved in 1999? Cooperative and collabor organizations in order and/or services. Pure of the property of the p	in biotechnology-related <b>coopera aborative arrangements</b> involve to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   1  ch purposes. Check any that are a	the active partic k on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Was your firm involved in 1999? Cooperative and collabor organizations in order and/or services. Pure of the properties	in biotechnology-related <b>coopera</b> aborative arrangements involve to to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   the purposes. Check any that are a search & development (R&D)/ Acceptable in the purpose in	the active partic k on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Arrangement  9110  Arrangement  Regulatory aff	in biotechnology-related <b>coopera</b> aborative arrangements involve to to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   the purposes. Check any that are a series are a court of the purpose of the p	the active partic k on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Arrangement  9110  Arrangement  9120  Regulatory aff  9130  Nas your firm involved in 1999?  Cooperative and collabor organizations in order and/or services. Pure of the properties of the prop	in biotechnology-related <b>coopera</b> aborative arrangements involve to to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   the purposes. Check any that are a series are a court of the purpose of the p	the active partic k on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Was your firm involved in 1999? Cooperative and collabor organizations in order and/or services. Pure of the properties	in biotechnology-related <b>coopera</b> aborative arrangements involve to to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   the purposes. Check any that are a countracting to the co	the active particle on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Arrangement  Please indicate for whice  Arrangement  Prototype dev  9150  Access marke	in biotechnology-related <b>coopera</b> aborative arrangements involve to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   the purposes. Check any that are a countracting of the purposes of the purposes of the purpose	the active particle on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Was your firm involved n 1999? Cooperative and collabor organizations in order and/or services. Pure of the process of the pro	in biotechnology-related <b>coopera</b> aborative arrangements involve to to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   the purposes. Check any that are a series are a development (R&D)/ Accounts  The purpose search & development (R&D)/ Accounts	the active particle on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			
8280 10. \ ii 0	Was your firm involved n 1999? Cooperative and collabor organizations in order and/or services. Pure of the process of the pro	in biotechnology-related <b>coopera</b> aborative arrangements involve to to develop and/or continue work contracting-out is not regarded as  Go to question 13  How many?   the purposes. Check any that are a series are a development (R&D)/ Accounts  The purpose search & development (R&D)/ Accounts	the active particle on new or sign collaboration.	ipation in projects by	/ your company and othe	er companies			

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A firm of smaller or equal size  On A larger firm  Government department/agency  University/Hospital/Research network  Was your firm as a spin-off?  A Spin-off is defined as a new firm created to transfer and commercialize inventions and technology developed in universities, firms or laboratories.  Devial No — Go to Question 14  Yes — Was your firm a spin-off from; — 1 University/hospital  2 Another company  3 Government agency/lab  4 Office (please specity)  Other (please specity)  Other (please specity)   Other (please specity)  A firm of smaller or equal size  A firm of small or equal size  Constraints  Imputs  Constraints  Domestic market too small  Lack of distribution & marketing channels  Constraints  Transportation regulations on biotechnology  Department protection for plants  Constraints  Constraints  Constraints  Control plants protection for plants  Lack of patient protection for markets  Control plants protection for plants  Lack of patient protection for plants  Lack of patient protection for markets  Cother (please specity)	2.	Check collaboration/co-operation arrangements by each type and	their geog	graphic I	ocation.				
A firm of smaller or equal size  A larger firm  Covernment department/agency  Cither (please specify)  Cither (please sp		Partner Category						America	Asia 4
A larger from	90	A firm of smaller or equal size			$\bigcirc$		$\bigcirc$	$\bigcirc$	
Converting the Special Programment department department department department department department department department department or substitutions and technology developed in universities, firms or laboratories.    Several   No   Second or   No   Second or		A larger firm			$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
University/riospital ricesearch network  Would you describe your firm as a 'spin-off?  A Spin-off is defined as a new firm created to transfer and commercialize inventions and technology developed in universities, firms or laboratories.  249 No Go to Question 14  Yes — Was your firm a spin-off from; — ! University/hospital  2 Another company  3 Government agency/lab  4 Other (please specify)   Obstacles to Biotechnology Commercialization  4. Rate the following obstacles to advancement of biotechnology commercialization activities in your firm. Use the following scale where 1 is low importance and 5 is high importance: indicate if not applicable to your firm.  Inputs  Access to capital  Access to capital  Access to technology/information  Access to technology/information  Company  Access to technology/information  Access to international markets  Domestic market too small  Constraints  Transportation regulations on biotechnology  Domestic market too small  Constraints  Transportation regulations on biotechnology  Patent rights held by others  Patent rights held by others  Lack of patent protection for nammals  Lack of patent protection for human components (e.g., organs, itssues)	10	Government department/agency			$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Would you describe your firm as a 'spin-off'?   A Spin-off is defined as a new firm created to transfer and commercialize inventions and technology developed in universities, firms or laboratories.   1	20	University/Hospital/Research network			$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
A Spin-off is defined as a new firm created to transfer and commercialize inventions and technology developed in universities, firms or laboratories.  9240 No — Go to Question 14  Yes — Was your firm a spin-off from; — 1 University/hospital  2 Another company  3 Government agency/lab  4 Other (please specify)  Other (please specify)  Low to following obstacles to advancement of biotechnology commercialization activities in your firm.  Use the following scale where 1 is low importance and 5 is high importance. Indicate if not applicable to your firm.  Inputs  Inputs  Access to capital  Access to capital  Access to technology/information  Access to technology/information  Access to thuman resources  Markets  Tansportation regulations on biotechnology  Constraints  To Public perception/acceptance  Regulatory requirements  Time/cost  Time/cost  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for human components (e.g., organs, tissues)		Other ( <i>please specify</i> )			0	$\bigcirc$	$\bigcirc$	0	$\bigcirc$
Use the following scale where 1 is low importance and 5 is high importance. Indicate if not applicable to your firm.    Low   1   2   3   4   5   5	0	A Spin-off is defined as a new firm created to transfer and comme firms or laboratories.  9240 No → Go to Question 14  Yes → Was your firm a spin-off from; → 1  2  3  4  bstacles to Biotechnology Commercialization	University Another (	sity/hosper compa nment aq ( <i>please</i> :	oital any gency/lab <i>specify</i> )			d in universiti	es,
Low   1	4.	Rate the following obstacles to advancement of biotechnology	commer						
Access to capital			portance.				e to your fii		Not
Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for human components (e.g., organs, tissues)  Access to technology/information  O OOO  Domestic market too small  O OOO  OOO  Domestic market too small  O OOO  OOO  OOO  Domestic market too small  O OOO  OOO  OOO  OOO  Domestic market too small  O OOO  OOO  OOO  OOO  Domestic market too small  O OOO  OOO  OOO  OOO  OOO  OOO  OOO			Low	lm	nportance	<b>.</b>	High	ı	licable
Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for naimals  Lack of patent protection for human components (e.g., organs, tissues)		Use the following scale where 1 is low importance and 5 is high im	Low	lm	nportance	<b>.</b>	High	ı	licable
Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements  Time/cost  Date of patent rights held by others  Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100	Use the following scale where 1 is low importance and 5 is high im	Low	lm	nportance	<b>.</b>	High	ı	licable
Domestic market too small	1100	Use the following scale where 1 is low importance and 5 is high im  Inputs  Access to capital	Low	lm	nportance	<b>.</b>	High	ı	licable
Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Constraints  Public perception/acceptance  Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	1100	Use the following scale where 1 is low importance and 5 is high im  Inputs  Access to capital  Access to technology/information	Low	lm	nportance	<b>.</b>	High	ı	licable
Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Public perception/acceptance  Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for human components (e.g., organs, tissues)	100 1110 120	Inputs  Access to capital  Access to technology/information  Access to human resources	Low	lm	nportance	<b>.</b>	High	ı	licable
Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets	Low	lm	nportance	<b>.</b>	High	ı	licable
Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small	Low	lm	nportance	<b>.</b>	High	ı	licable
Public perception/acceptance  Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130 140	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets	Low	lm	nportance	<b>.</b>	High	ı	licable
Public perception/acceptance  Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130 140	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology	Low	lm	nportance	<b>.</b>	High	ı	licable
Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130 140 150	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels	Low	lm	nportance	<b>.</b>	High	ı	licable
Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130 140 150 160	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints	Low	lm	nportance	<b>.</b>	High	ı	licable
Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130 140 150 160	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance	Low	lm	nportance	<b>.</b>	High	ı	licable
Lack of patent protection for plants  Lack of patent protection for animals  Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130 140 150 160	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements	Low	lm	nportance	<b>.</b>	High	ı	licable
Lack of patent protection for animals  Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130 140 150 160 170 180	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements  Time/cost	Low	lm	nportance	<b>.</b>	High	ı	licable
Lack of patent protection for human components (e.g., organs, tissues)	100 110 120 130 140 150 160 170 180 190	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements  Time/cost  Patent rights held by others	Low	lm	nportance	<b>.</b>	High	ı	licable
	100 110 120 130 140 150 160 170 190 200 210	Inputs  Access to capital Access to technology/information Access to human resources  Markets  Domestic market too small Lack of access to international markets  Transportation regulations on biotechnology Lack of distribution & marketing channels  Constraints  Public perception/acceptance Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants	Low	lm	nportance	<b>.</b>	High	ı	licable
	100 110 120 130 140 150 160 170 190 200 210	Inputs  Access to capital  Access to technology/information  Access to human resources  Markets  Domestic market too small  Lack of access to international markets  Transportation regulations on biotechnology  Lack of distribution & marketing channels  Constraints  Public perception/acceptance  Regulatory requirements  Time/cost  Patent rights held by others  Lack of patent protection for plants  Lack of patent protection for human components	Low	lm	nportance	<b>.</b>	High	ı	licable

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							Geogr	raphic Lo	cation	
					None 5	Canada 0	USA 1	Europe	Latin Americ	a .
11100	Existing patents				<u> </u>		'	2	3	
11110	Pending patents									
	se indicate the number of cate '0' if none)	f patent appli	cations you	r company s	ubmitte	ed to the fo	llowing	Patent Off	ices.	
11120	Patent Office/Year						<b>1998</b>		<b>1999</b>	
11130	Canadian Intellectual P	roperty Office	(CIPO)						'	
11140	United States Patent &	Trademark Of	ffice (USPTC	D)						
11150	European Patent Office	e (EPO)								
11160	Other (please specify)									
11180 11190 11120	Plant Variety Protection  Community Plant Varie  Other (please specify)	n Office, USDA								
During 1	ual Property  the last two years, 1998  the right to use intelle  No → Go to Que	ctual propert	y from anot	her firm?					ther firm or did	your
	Yes → Please inc	dicate the type	and direction	on of such in	tellectu	al property	/ transfe	er.		
	tual Property	Canadi	Rights to an Firms	Granted Foreigr	Rights Firms	to Acc	quired R Canadia 2		Acquired R Foreign	ights Firm
Intellec		Yes	No	Yes	No	) \	'es	No	Yes	N
Intellec			$\bigcirc$		C	) (		0	0	(
Trade S	Secrets/Licensing nents					,				
	nents	0	0	0	C	) (	<u> </u>	$\bigcirc$	0	(

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Revenues, Expenditures & Trade  17. Please provide financial details in the following table. P	lease report for fiscal y	years and	in thou	sands of do	ollars (	\$,000's).		
Indicate "0" if none					ase provide details in \$,000's for the years			
		<b>1998</b>		<b>1999</b>		for 20 200 2		
Total Firm Sales/Revenue	\$		000	\$	,000	\$	,000	
13110 % of Total Sales/Revenue From Biotechnology			%		%		%	
Total R&D Spending	\$	,(	000	\$	,000	\$	,000	
% of R&D Spending on Biotechnology R&D			%		%		%	
Total Exports (including licensing agreements)	\$	,(	000	\$	,000	\$	,000	
% of Exports from Biotechnology			%		%		%	
Total Imports	\$	,(	,00	\$	,000	\$	,000	
% of Imports from Biotechnology			%		%		%	
18. If your firm <b>exported</b> biotechnologies, what percentage 1999? Include licensing agreements. What is your forec		2002?	(	Geographi	c Loca	ition		
		Canada 0	<b>USA</b>	Europe 2	Latir	n America	Asia 4	
13180 1999								
<sup>13190</sup> Forecast for 2002								
Year		Canada	USA	Geographic		ntion n America	Asia	
		O	1 1	2	Latii	3	4	
1999								
Forcast for 2002								
20. a) Did your firm attempt to raise capital for biotechnolog  13220 No → Go to Question 20 c)  Yes	y in fiscal year 1999?							
b) Were you successful in raising capital?								
$_{13230}$ $\bigcirc$ No $\longrightarrow$ Go to Question 20 c)								
	\$ ,000	0						
Indicate the sources of capital and the percentage (%	b) of total capital that s	ource pro	vided ir	า 1999.				
Source		%	of To	tal Capital	•			
13240 Angel investors/family/friends					•			
13250 Government loans/grants/incentives								
13260 Venture Capital funds								
13270 Conventional sources (i.e. banks)								
13280 Initial Public Offering (IPO)								
13290 Collaborative alliance								
13300 Other (please specify)								
Sales (please openly)		TOTAL	10	0%				

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20. c)	) Does	your	firm plar	n to raise capital in 2002?				
	13310		No →	Go to Question 21				
			Yes→	How much do you plan to ra	ise in 2002? -	→ 1	Less than \$500,000	
						2	\$500,000 to \$5,000,000	
						3	More than \$5,000,000	
21. 1	n the n	ast 5 v	veare did	your firm apply for the tay he	anefit for hiotec	hnology rel	elated activities under the R&D (SRED) tax pro-	aram?
,	ii tiie p	asi o y	ears uic	a your mini apply for the tax be	erielit for blotec	illiology lei	lated activities drider the NXD (SIXED) tax pro-	grain:
	13320		No →	Why? $\longrightarrow$ <sup>1</sup> $\bigcirc$ Complex	ity of application	n process		
				<u> </u>	nty of eligibility			
				9	neet eligibility r	•		
			Yes	<sup>4</sup> Other (p	lease specify)			
22. [	Does yo	our firn	n use th	e Internet?				
				0 1 0 1 00				
	13330	$\bigcirc$	No →	Go to Question 23				
		$\bigcirc$	Yes→	Indicate for what purposes y (Check any that are applicable)		ne Internet.		
			1		•	6		
			2	Silalling research & dev	elopment	6	Truman resource search	
			3	- Warksting/seiling	orviooo	8	Public relations  Conoral communication	
			4	Purchasing goods and s  Accessing databases/in		9	General communication Other (please specify)	
			5	E-commerce	ioimation sour	ces	Other (please specify)	
				L commerce				
			_	strategies did your firm use i	n 1999?			
	13400	1 (	Ref	ocused product development	8	Licensed in	in technology	
		2 (	O Dov	vnsized	9	Licensed of	out technology	
		3 (	Incr	reased size	10	Merged wi	ith other company	
		4 (	Ent	ered product trials	11 🔾	Formed a	joint venture	
		5 (	O Lau	nched new product	12 🔾	Expanded	l into foreign markets	
		6 (	O Acq	uired a company	13 🔾	No change	e	
		7 (	Out	-source production	14 🔾	Other (plea	ease specify)	
Com	ments							
14100								
ļ	f you ha	ave ar	ny comm	nents regarding this survey, pl	ease provide th	nem in the	space below.	
_								
_								
_								
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-								
=								
=								
-								
=								

Thank you for your co-operation

Please return the questionnaire in the return prepaid envelope.

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# Appendix 3: Biotechnology Use and Development Survey – 2001; First phase questionnaire



## Biotechnology Use and **Development Survey -**

Confidential once completed

Collected under the authority of the Statistics Act, Revised Statutes of Canada, 1985, Chapter S19. Completion of this questionnaire is a legal requirement under the Statistics Act.

Version française au verso



### Information for the Respondent

#### **Purpose of Survey**

Statistics Canada is conducting this survey in order to develop information on biotechnology and related technologies such as functional foods, nutraceutical and bioproducts by identifying industry sectors where these activities take place. Please report on Canadian activities of your firm in biotechnology, functional foods, nutraceutical or bioproducts. Your firm may have responded to biotechnology questions in previous surveys, but there is also an increasing demand for information on other technologies and their impact on the Canadian economy.

#### **Authority**

Collected under the authority of the Statistics Act, Revised Statutes of Canada, 1985, Chapter S19. Completion of this questionnaire is a legal requirement under the Statistics Act.

#### Confidentiality

Statistics Canada is prohibited from publishing any statistics that would divulge information obtained from this survey that relates to any identifiable business, institution or individual. Data is treated in strict confidence, used for statistical purposes and released in aggregate form only. The confidentiality provisions of the Statistics Act are not affected by either the Access to Information Act or any other Legislation.

#### **Federal-Provincial Agreement**

In order to avoid duplication of enquiry, reduce the cost of collection, and provide consistent statistics, Statistics Canada has entered into an agreement with the Institut de la Statistique du Québec, under Section 11 of the Statistics Act. Data collected from Québec firms in this survey will be transmitted to the Institut de la Statistique du Québec. The Statistics Act of provisions for Quebec includes the same confidentiality and penalties for disclosure of information as the Federal Statistics Act.

#### Instruction

A knowledgeable senior person in your firm, such as an R&D manager or production manager, can quickly complete this questionnaire. Please fill in the contact information below, answer all 3 questions and return the completed questionnaire in the accompanying self addressed prepaid envelope to Statistics Canada by March 7, 2002.

#### Assistance

If you have questions or require assistance please contact:

> Claire Racine-Lebel 7th floor, RHCoats Building Statistics Canada

Telephone: 613-951-6309 Fax: 613-951-9920 E-mail: Sieidinfo@statcan.ca

## Please provide the following information: Name of person completing this form Telephone number Title Fax number E-mail Web address

5-4900-505: 2001-01-24 STC/SAT-465-75330



Statistics Statistique Canada Canada



1.	Does your firm currently use or develop biotechnology in its activities?
	O Yes
	O No
	○ INO
	Examples of biotechnologies:
	DNA genomics, pharmaco-genetics gene probes, DNA sequencing/synthesis/amplification, genetic engineering. Protein/peptide sequencing/synthesis, lipid/protein engineering, proteomics, hormones and growth factors, cell receptors/signalling/pheromones, cell/tissue culture, tissue engineering, hybridisation, cellular fusion, vaccine/immune stimulants, embryo manipulation, bioreactors, fermentation, bioprocessing, bioleaching, bio-pulping, bio-bleaching, biodesulphurization, bioremediation, biofiltration, gene therapy, viral vectors, bioinformatics, other.
2.	Does your firm currently make or develop functional foods or nutraceutical products?
	O Yes
	○ No
	Functional food
	is a conventional food, beverage, or ingredient enriched with functional components beneficial in
	disease prevention or disease-risk management, beyond basic nutritional functions. A food, beverage or ingredient may be made functional through a variety of means, such as the addition of components, extraction, fractionation, processing, plant or livestock breeding, livestock feeding techniques, genetic modification, other.
	Nutraceutical
	is a product isolated or purified from foods (includes herbs and botanicals) that is generally sold in medicinal forms not usually associated with food. A nutraceutical is demonstrated to have a physiological benefit or provide protection against chronic disease.
3.	Does your firm currently make or develop a bioproduct?
	O Yes
	O No
	Bioproduct
	a commercial or industrial product (other than food and feed) made with biological or renewable domestic agricultural (plant, animal), marine or forestry materials, such as, bio-energy (heating and electricity), bio-fuels (ethanol and bio-diesel), biochemicals, fiberboard, textiles and bio-plastics, other.
	Thank you for your cooperation

Please return the completed questionnaire in the accompanying self addressed prepaid envelope

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### Appendix 4: Biotechnology Use and Development Survey – 2001; Second phase questionnaire



### **Biotechnology Use and Development Survey - 2001**

#### Confidential when completed

Collected under the authority of the Statistics Act, Revised Statutes of Canada, 1985, c. S-19. Completion of the questionnaire is a legal requirement under the Statistics

Si vous préférez ce questionnaire	
en français, veuillez cocher	



#### Information for the Respondent

#### **Survey Purpose**

Statistics Canada is undertaking this survey to produce a profile of firms engaged in biotechnology activities in Canada. The survey focuses on the characteristics and activities of firms that use or develop biotechnology as part of their company's activity.

Biotechnology is an emerging sector of the Canadian economy and its impact has the potential to be felt through all parts of Canada's society. An accurate understanding of biotechnology requires comprehensive data. Information from this survey may be used by businesses for economic or market analysis, by trade associations to study industry performance, government departments and agencies to assist policy formation, and by the academic community for research purposes. Statistics Canada may create a database by combining survey data with existing Statistics Canada data records.

Please report 2001 on Canadian biotechnology activities of your firm unless a specific question indicates otherwise. Complete a separate questionnaire for each company engaged in biotechnology activities in Canada.

#### Confidentiality

Statistics Canada is prohibited from publishing any statistics that would divulge information obtained from this survey that relates to any identifiable business, institution or individual. Data is treated in strict confidence, used for statistical purposes and released in aggregate form only. The confidentiality provisions of the Statistics Act are not affected by either the Access to Information Act or any other Legislation.

#### **Federal-Provincial Agreement**

In order to avoid duplication of enquiry, reduce the cost of collection and provide consistent statistics, Statistics Canada has entered into an agreement with the Institute de la Statistique du Québec. Under Section 11 of the Statistics Act data collected from Quebec firms in this survey will be transmitted to the Institut de la Statistique du Québec. The Statistics Act of Quebec includes the same provisions for confidentiality and penalties for disclosure of information as the Federal Statistics Act.

#### Who Should Complete This Questionnaire?

A senior manager, scientist/researcher or production manager should complete this questionnaire.

#### Assistance

If you have questions or require assistance please contact:

> Science, Innovation and Electronic Information Division Statistics Canada **Tunneys Pasture** Ottawa K1A 0T6

Claire Racine-Lebel

Telephone: 613-951-6309 (Call collect)

Fax: 613-951-9920 E-mail: Sieidinfo@statcan.ca

Name of person completing this form	Telephone number  Area code
Title	Fax number
Web address	E-mail

5-5300-500.1: 2002-02-06 STC/SAT-430-75177



Statistics Canada

Statistique Canada



	Currently	If currently us	sing, do you	use them for	Number	If <b>No</b> ▼
Biotechnologies	Used in Operation	Product/ Process Development	Current Production	Environmental Purposes	of Years in Use	Do you plan t use within 3 years?
DNA - the coding	0	1	2	3	4	5
Genomics/Pharmaco-genetics	○ Yes -	<b></b> ()	$\bigcirc$	0		→ ○ Yes ○
Gene probes	○ Yes -	<b></b>	$\bigcirc$	$\circ$		→ ○ Yes ○
DNA sequencing synthesis amplification, Genetic Engineering	○ Yes -	<b></b>	$\bigcirc$	$\bigcirc$		→ ○ Yes ○
Proteins and Molecules - the	functional b	locks				
Protein/peptide sequencing/ synthesis	○ Yes -	<b></b>	0	$\bigcirc$		→ ○ Yes ○
Lipid/protein engineering	Yes -	<b>\rightarrow</b> ()	<u> </u>			→ ○ Yes ○
Proteomics	○ Yes -	<b></b>	$\bigcirc$	<u> </u>		→ ○ Yes ○
Hormones, growth factors, pheromones	○ Yes -		$\bigcirc$	<u> </u>		→ ○ Yes ○
Cell receptors signalling	$\sim$	<b></b>	$\bigcirc$	<u> </u>		→ ○ Yes ○
Cell and Tissue Culture, and	Engineering					
Cell/ tissue culture, Embryo manipulation	$\sim$		<u> </u>			→ ○ Yes ○
Tissue engineering	<u> </u>	<b></b>	$\bigcirc$	<u> </u>		→ ○ Yes ○
Hybridization	○ Yes -	<b>→</b> ()	$\bigcirc$	0		→ ○ Yes ○
Cellular fusion		<b>\rightarrow</b> ()	<u> </u>	0		→ ○ Yes ○
Vaccine/immune stimulants	$\circ$	<b></b>	0			→ ○ Yes ○
Process Biotechnologies						
0 Bioreactors	$\tilde{}$	$\longrightarrow$ $\bigcirc$				► O Vac O
	○ No -					→ ○ Yes ○
Fermentation, Bioprocessing	Yes -		$\bigcup$	$\bigcup$		→ ( ) Yes ( )

echnologies	Used in Operation	Product/ Process Development	Current Production	Environmental	of Years	Do you plan t	
	1	20.0.0pmont		Purposes	in Use	Do you plan use within 3 years?	
	0	1	2	3	4	5	
Bio-pulping, Biodesulphurization	Yes -	<b>-</b>		$\bigcirc$		→ Yes	
n, Biofiltration	○ Yes -	<b></b> ()		<u> </u>		→	
r Organisms							
у		<b></b> ()		0		→ ○ Yes ○	
	○ Yes -	<b></b>		$\bigcirc$		→	
3	○ Yes -	<b>-</b>	<u> </u>	0		→ ○ Yes ○	
nologies	○ Yes -	<b></b>	<u> </u>	0		→ ○ Yes ○	
Specify:	Yes -	<b>-</b>	0	0		→ ( ) Yes ( )	
	-						
	pn, Biofiltration  ar Organisms  y  s  nologies  e Specify:	Yes — No —  Ar Organisms  Yes — No —  No —  Yes — No —  Yes — No —  Yes — No —  Yes — No —  Yes — No —  Se Specify:  Yes — Yes — No —	Yes No  Ar Organisms  Yes No  No  Yes No  Yes No  No  Yes No	Ann, Biofiltration  Yes  No  Ar Organisms  Yes  No  Yes  No  Yes  No  No  Yes  No  Se Specify:  Yes  Yes  No  Yes  No  Yes  No  Yes  No  Yes  No  Yes  No  No  No  Yes  No  No  Yes  No  No  Yes  No  No  No  No  No  No  No  No  No  N	on, Biofiltration  Yes  No  Ar Organisms  Yes  No  No  Yes  No  Yes  No  Yes  No  No  Se Specify:  Yes  No  Yes  No  Yes  No  Yes  No  No  No  Yes  No  No  No  No  No  No  No  No  No  N	on, Biofiltration  Yes  No  Ar Organisms  Yes  No  Yes  No  Yes  No  No  Se Specify:  Yes  Yes  Yes  No  Se Specify:  Yes  No  No  No  No  No  No  No  No  No  N	

If you do not use any of the biotechnologies listed in Question 1

Please return the questionnaire in the accompanying prepaid return envelope.

Thank you for your assistance.

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Using the table below, please rate the level of influence	Low	lm	portano	e	High
of each factor on increasing your use of biotechnology.	1	2	3	4	5
Inputs					<b>→</b>
Access to capital	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Access to technology/information		$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Access to human resources		$\bigcirc$	$\bigcirc$	$\bigcirc$	
Markets					
Size of Domestic Market		$\bigcirc$	$\bigcirc$	$\bigcirc$	
Access to international markets	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Information about markets		$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Distribution & marketing channels		$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Constraints					
Public perception/acceptance		$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Cost of regulatory approval	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Time required for regulatory approval		$\bigcirc$	$\bigcirc$	$\bigcirc$	
Limited international harmonization					$\overline{\bigcirc}$
Patent rights held by others		$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Lack of protection for intellectual property					
Lack of protection for intellectual property  Other, Please specify:	0				
Lack of protection for intellectual property	0	0	0	0	0
Other, Please specify:	0		0	0	0
Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use	Low	Im	portanc	ce	High
Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.	Low 1	 	portance		High 5
Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity					_
Contact of protection for intellectual property  Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs				:e 4	_
Capital costs  Other, Please specify:  Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs				©	_
Cack of protection for intellectual property  Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs				©	_
Cack of protection for intellectual property  Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products				©	_
Cack of protection for intellectual property  Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced				©	_
Copital costs  Improved Products  New products or processes introduced  Product range increased  Other, Please specify:  Other, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Improved Products  New products or processes introduced				©	_
Cother, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced  Product range increased  Product quality increased				©	_
Cother, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced  Product range increased  Product quality increased  Knowledge Based				©	_
Cother, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced  Product range increased  Product quality increased  Knowledge Based  Developing new areas for R&D				©	_
Cother, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced  Product range increased  Product quality increased  Knowledge Based  Developing new areas for R&D  Increase efficiency for R&D				©	_
Cother, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced  Product range increased  Product quality increased  Knowledge Based  Developing new areas for R&D  Improved Market Performance				©	_
Cother, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced  Product range increased  Product quality increased  Knowledge Based  Developing new areas for R&D  Improved Market Performance  Market position improved				©	_
Cother, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced  Product range increased  Product quality increased  Knowledge Based  Developing new areas for R&D  Improved Market Performance  Market position improved  New Market Niche Developed				©	_
Cother, Please specify:  For each of the performance factors listed below, please rate the level of impacts of biotechnology use on your firm's performance.  Increased Productivity  Labour costs  Capital costs  Energy costs  Improved Products  New products or processes introduced  Product range increased  Product quality increased  Knowledge Based  Developing new areas for R&D  Improved Market Performance  Market position improved				<ul><li>Calcal Control</li><li>Calcal Control</li></ul>	_

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#### Section 3 - Human Resources in Biotechnology

Concerns have been expressed about the availability of skilled biotechnology employees. Your cooperation in careful completion of this section is essential in developing an accurate understanding of human resources in biotechnology. For the purpose of this survey Employees are defined as those workers for whom you completed a Canada Customs and Revenue Agency T-4 statement for the 2001 tax year. Include working owners. Do not include students. Only count employees working in Canada. If '0' (zero) indicate '0'.

#### **Number of Biotechnology Employees**

4.	a)	How many employees does your firm employ in Canada? Please Report Typical Employment Level for 2001.	4000
	b)	How many employees have biotechnology-related responsibilities?	4010

c) Full-time Biotechnology Employees

Please Report Typical Employment Level for 2001.

For each group listed below indicate how many are full-time biotechnology employees (50% or more of their time spent on biotech related activities)? If an employee fulfils more than 1 duty, report their primary responsibility. Count each person only once. Please Report Typical Employment Level for 2001.

Position	Number of full-time
Scientific Research & Direction	4100
Technicians	4110
Regulatory/Clinical Affairs	4120
Production	4130
Finance/Marketing	4140
Management	4150
Other, Please Specify:	4160
Total Full-time employees	4170

#### Part-time Biotechnology Employees

d) For each group listed below indicate how many are Part-time biotechnology employees (less than 50% of their time spent on biotech related activities)? If an employee fulfils more than 1 duty, report their primary responsibility. Count each person only once. Please Report Typical Employment Level for 2001.

Position	Number of part-time
Scientific Research & Direction	4200
Technicians	4210
Regulatory/Clinical Affairs	4220
Production	4230
Finance/Marketing	4240
Management	4250
Other, Please Specify:	4260
Total Part-time employees	4270

e) Total Number of biotechnology employees.

Total full-time and part-time employees with biotechnology-related responsibility (Box 4170 + Box 4270)

4011	This
	equ

This number must equal 4010 above.

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<b>a)</b> D		Co to augotion Fh					
50	000 ( No	Go to question 5b					
	Yes Yes	In the table below indicate the number o category.	of unfilled positions	by			
		Position	Number o Unfilled Positions				
	l	Scientific Research & Direction	5100				
		Technicians	5110				
		Regulatory/Clinical Affairs	5120				
		Production	5130				
			5140				
		Finance/Marketing	5150				
	,	Management	5160				
		Other, Please Specify:	3100				
		Total unfilled positions	5170				
	Yes      Yes	<ul> <li>Were you successful?</li> <li>™ No</li></ul>	5310				
54 54	Vhat sources w  400 Univers  410 Tempor  420 Employe  430 Profess  440 Own Sta	00 No Go to question 6	5450 5460 5470 5480 5490	Other Pharm Newsp Stude Interna	naceutica paper/Jou nt Interns al Trainin	urnal ship g of Staff	
54 54	Vhat sources w  400 Univers  410 Tempor  420 Employe  430 Profess  440 Own Sta	OOO No Go to question 6 Yes How many did you hire?  Were successfully used in recruiting biot ity Recruitment rary/Contract Staff ment agencies/Headhunters ional Associations  aff/Incentive program	technology staff?  5450  5460  5470  5480  5490   rts in filling bioted	Other Pharm News Stude Interna	naceutica paper/Jou nt Interns al Trainin	al Firms urnal ship ag of Staff	cies.
54 54	Vhat sources w  400 Univers  410 Tempor  420 Employe  430 Profess  440 Own Sta	OOO No Go to question 6 Yes How many did you hire?  Were successfully used in recruiting biot ity Recruitment rary/Contract Staff ment agencies/Headhunters ional Associations  aff/Incentive program	5450 5460 5470 5480 5490	Other Pharm News Stude Interna	naceutica paper/Joi nt Interns al Trainin gy-relate	al Firms urnal ship ag of Staff	
54 54	Vhat sources w  400 Univers  410 Tempor  420 Employ  430 Profess  440 Own Sta	No So to question 6 Yes How many did you hire?  Vere successfully used in recruiting biot hity Recruitment frary/Contract Staff  ment agencies/Headhunters  ional Associations  aff/Incentive program  act of the following factors on your effort	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm Newsp Stude International	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.
54 54	Vhat sources w  400 Univers  410 Tempor  420 Employ  430 Profess  440 Own Sta	No So to question 6 Yes How many did you hire?  Vere successfully used in recruiting biot hity Recruitment frary/Contract Staff  ment agencies/Headhunters  ional Associations  aff/Incentive program  act of the following factors on your effort	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm Newsp Stude International	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.
54 54 54 Pleas	Vhat sources w  400 Univers  410 Tempor  420 Employ  430 Profess  440 Own Sta  re rate the impa  Factors  Candidate I  Compensation	OO No So Go to question 6 Yes How many did you hire?  Vere successfully used in recruiting biotority Recruitment  Farary/Contract Staff  Ment agencies/Headhunters  John Associations  J	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm Newsp Stude International	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.
54 54 54 Pleas	Vhat sources w  400 Univers  410 Tempor  420 Employ  430 Profess  440 Own Sta  Factors  Candidate I  Compensation  Candidates ur	No So to question 6 Yes How many did you hire?  Vere successfully used in recruiting biot ity Recruitment rary/Contract Staff ment agencies/Headhunters ional Associations aff/Incentive program  Factors  In requirements by candidates too high mwilling to relocate	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm News Stude Interna	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.
54 54 54 54 Pleas 6000 6010 6020	Vhat sources w  400 Univers  410 Tempor  420 Employ  430 Profess  440 Own Sta  Factors  Candidate I  Compensation  Candidates ur  Lack of experi	No So to question 6 Yes Men How many did you hire?  Vere successfully used in recruiting biotority Recruitment  Farary/Contract Staff  Ment agencies/Headhunters  Minional Associations  Margeff/Incentive program  Meact of the following factors on your effort  Factors  Marger requirements by candidates too high  Menwilling to relocate	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm News Stude Interna	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.
54 54 54 54 Pleas 6000 6010	Vhat sources wath with the sources wath wath with the sources wath wath with the sources wath wath wath with the sources wath wath wath with the sources wath wath wath wath wath wath wath wath	No So to question 6 Yes How many did you hire?  Vere successfully used in recruiting biot ity Recruitment rary/Contract Staff ment agencies/Headhunters ional Associations aff/Incentive program  Act of the following factors on your effort  requirements by candidates too high  mwilling to relocate  fence  res  reces insufficient to attract candidates	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm News Stude Interna	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.
54 54 54 54 Pleas 6000 6010 6020	Vhat sources wath with the sources wath wath with the sources wath wath with the sources wath wath wath with the source wath wath wath with the source wath wath wath wath wath wath wath wath	No So to question 6 Yes Mow many did you hire?  Vere successfully used in recruiting biot ity Recruitment rary/Contract Staff ment agencies/Headhunters ional Associations aff/Incentive program  Factors In requirements by candidates too high Inwilling to relocate Interest of the following factors on your efforts  The requirements by candidates too high Invilling to relocate Interest of the following factors on your efforts  The requirements by candidates too high Invilling to relocate Interest of the following factors on your efforts  The requirements by candidates too high Interest of the following factors on your efforts  The requirements by candidates too high  The requirements by candidates too high  The requirements by candidates too high  The requirements of the following factors on your efforts  The requirements by candidates too high  The requirements by the requirements by candidates too high  The requirements by the	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm News Stude Interna	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.
54 54 54 54 54 Pleas 6000 6010 6020 6100	/hat sources w 400 Univers 410 Tempor 420 Employ 430 Profess 440 Own Sta  e rate the impa  Factors  Candidate I Compensation Candidates ur Lack of experi Firm Factor Capital/resour External Fa Lack of qualific	No So to question 6 Yes Mow many did you hire?  Vere successfully used in recruiting biot ity Recruitment rary/Contract Staff ment agencies/Headhunters ional Associations aff/Incentive program  Factors In requirements by candidates too high Inwilling to relocate Items Incentive program  Factors In requirements by candidates too high Inwilling to relocate Items Ite	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm News Stude Interna	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.
54 54 54 54 Pleas 6000 6010 6020	/hat sources wath with the sources wath wath with the sources wath wath with the sources wath wath wath with the source wath wath wath with the source wath wath wath wath wath wath wath wath	No So to question 6 Yes Mow many did you hire?  Vere successfully used in recruiting biotity Recruitment rary/Contract Staff ment agencies/Headhunters ional Associations aff/Incentive program  act of the following factors on your effort  Factors In requirements by candidates too high mwilling to relocate fence fence fence fres freces insufficient to attract candidates for qualified candidates for qualified candidates	technology staff?  5450  5460  5470  5480  5490  Trts in filling bioted	Other Pharm News Stude Interna	naceutica paper/Jou nt Interns al Trainin gy-relate nportane	al Firms  urnal  ship  g of Staff  ed vacan	cies.

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7.	Dic	you attempt to hire biotechnology staff from outside of Canada in 2001?
	700	No No Go to question 8
		Yes Was your firm successful in hiring from outside of Canada?
		7010 No So to question 8.
		Yes How many staff from outside Canada did you hire?
8.	Dic	d any biotechnology personnel leave your firm in 2001?
	800	
		8010
		Yes How many?
9	octic	on 4 - Biotechnology Products
Ih	IS S	ection measures the development of new biotechnology products and processes by your firm.
9.	a)	Do you have biotechnology products/processes on the market?
		9000 No Go to question 9b)
		Yes What year was the most significant product first introduced?
	b)	Is your firm currently developing products that require the use of biotechnology?  9100 No Go to question 9c)
		Yes What year will the most significant of these products reach market?
		Vitat year will the most significant of these products reach market:
	c)	Is your firm currently developing processes that require the use of biotechnology?
	٠,	9200 No Go to question 9d)
		Yes What year will the most significant of these processes be completed?
	d)	Do you consider biotechnology central to your firm's activities or strategies?
		9300 No
Ī		( ) Yes
1		
	e)	If you answered "Yes" to any  Go to Q10
		Part of Question 9
		Otherwise Please return the questionnaire in the
		accompanying prepaid return envelope.
		Thank you for your assistance.

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10. In the table below, for each sector listed please indicate the number of biotechnology products or processes your firm currently has for each stage of development. Number of biotechnology products/processes by development stage Research Regulatory phase/ Approved/ **Biotechnology Sector** Pre-clinical trials/ Unconfined release On market/In Confined field trials assessment production Development 0 1 2 3 **Human Health** Diagnostics (e.g. biosensors, immunodiagnostics, gene probes) Therapeutics (e.g. vaccines, immune stimulants, biopharmaceuticals) 10020 Drug Delivery **Agriculture Biotechnology** Plant Biotechnology (e.g. tissue culture, embryogenesis, genetic markers, genetic engineering) <sup>10110</sup> **Animal Biotechnology** (e.g. diagnostics, therapeutics, embryo transplantation, genetic markers, genetic engineering) Non-food Agriculture (e.g. fuels, lubricants, commodity and fine chemical feedstocks, cosmetics) **Natural Resources** Energy (e.g. microbiologically enhanced petroleum recovery, industrial bioprocessing, biodesulphurization) Mining (e.g. microbiologically enhanced mineral recovery, industrial bioprocessing, biodesulphurization) <sup>10220</sup> Forest Products (e.g. biopulping, biobleaching, biopesticides, tree biotechnology, industrial bioprocessing) **Environment**  $^{10300}\,\text{Air}$  (e.g. bioremediation, diagnostics, phytoremediation, biofiltration) Water (e.g. biofiltration, diagnostics, bioremediation, phytoremediation) Soil (e.g. biofiltration, diagnostics, bioremediation, phytoremediation) **Aquaculture**  $^{10400}\,\mathrm{Fish}$  health, broodstock genetics, bioextraction **BioInformatics** 10500 Genomics & molecular modelling (e.g. DNA/RNA/protein synthesising & databases for humans, plants, animals, and micro-organisms) Gene therapy (e.g. gene identification, gene constructs, gene delivery) **Food Processing** Bioprocessing (e.g. using enzymes and bacteria culture) Functional Foods/Nutraceuticals (e.g. probiotics, unsaturated fatty acids 10620 Other, Please Specify

	11000	Years	11001	Months	<b>S</b>								
<b>b)</b> V	Vhat is the	e total co	st to bring	vour p	rincipal biote	chnol	oav product	or pro	cess from	the ini	tial develo	opment	
					market? If st							эрттотт	
	\$		,000										
Section	15 - Bu	usiness	Praction	ces									
Contrac	_												
•					ogy related a	ctivit	ies in 2001?						
'			o to questi		P 4 11 1					,			
	Y	es Fo	or each pai	tner type	e listed below,	pleas					in 2001 f		ip listed
							10	otai va		000)	III 2001 I	Of	
	P	artner T	ype		Number of Contracts				Purpose	of Cont	ract	T	
							R&D		ulatory/ inical		gement/ duction	0	ther
12100 Privat	te Entities	(C.R.O's /	other Firm	s, etc)	0	<b>e</b>	,000	<b>e</b>	,000	\$	,000	<b>e</b>	,00
12110 Public		Universitie	es / Govern	ment		\$		\$				\$	-
Labs.	.)					\$	,000	\$	,000	\$	,000	\$	,00
		12310 Unit 12320 Gov 12330 Oth	ate resea versity/Ho vernment er biotech er, Please	spital ab nology					% % % %				
<b>c)</b> R	Rate the le	evel of im	portance	of each	of the follow	ing re	easons on yo	our dec		ontract	out.		
								Low	lm	portan	ce	Lliab	
R	Reasons 1	or Conti	acting O	ut				1	2	3	4	High 5	
12400 <sub>k</sub>	Knowledge	not avails	ıhle interna	llv				$\overline{}$	$\bigcap$	$\bigcap$	$\stackrel{\longrightarrow}{\cap}$	<b>&gt;</b>	
	Access out							$\overline{\bigcirc}$		$\bigcirc$			
_	Cost Redu												
12420	R&D Ac							$\bigcap$		$\bigcirc$			
12430		ory/Clinica	al Affairs					$\bigcirc$			$\overline{}$		
12440	Producti							$\bigcirc$	$\bigcirc$	$\bigcirc$	$\overline{}$	$\overline{\bigcirc}$	
<sup>12450</sup> F	Precursor t		agreemen	t				$\bigcirc$	$\bigcirc$	$\bigcirc$	$\overline{}$	$\overline{\bigcirc}$	
40400	Reduce risl							$\bigcirc$	$\bigcirc$	$\bigcirc$	$\overline{}$	$\overline{\bigcirc}$	
_	Other, Plea												

**11. a)** What is the total time required to bring your principal biotechnology product or process from the initial development phase/proof of concept stage to the market? If still in pre-market stages provide an estimate.

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12610 S	Contract Services  Routine Lab services  Specialized Lab services  Production/manufacturing services  Other, Please Specify:	er	nber of contracts ntered in 2001	Revenue receive source in 1	
12610 S 12620 P 12630 C	Specialized Lab services  Production/manufacturing services  Other, Please Specify:	5	0		
12620 P 12630 C	Production/manufacturing services Other, Please Specify:	5		Ψ	,000
12630 C	Other, Please Specify:	5		\$	,000
=				\$	,000
= 12640 <b>T</b> -	otal			\$	,000
				\$	,000
between you continue wor Pure contra	e and collaborative arranger company and other compark on new or significantly in cting-out work is not regard	panies or orga mproved biotec arded as collab	nizations in ord hnology proces poration.	der to develop a sses and/or prod	nd/or ducts.
13000	<ul><li>Go to question 14</li><li>Provide the number of arranger</li></ul>	ments by purpose a	and partner type		
		None	-b	manta ku Bartua	. T
Arrange	ement Purpose	Biotech Firm	Non-biotech Firm	Academic Institution/ Hospital	Government lab or agency
Arrange  3100  To conduct research &		Biotech	Non-biotech	Academic Institution/	Government
		Biotech Firm	Non-biotech Firm	Academic Institution/	Government lab or agency
<sup>3100</sup> To conduct research &	development (R&D)	Biotech Firm	Non-biotech Firm	Academic Institution/	Government
To conduct research & 3110 Regulatory affairs 3120 Access others' patents	development (R&D)	Biotech Firm	Non-biotech Firm	Academic Institution/	Government
To conduct research & Regulatory affairs	development (R&D)	Biotech Firm	Non-biotech Firm	Academic Institution/	Government
To conduct research & 3110 Regulatory affairs 3120 Access others' patents 3130 Production/manufactur 3140 Access markets/distrib	development (R&D)	Biotech Firm	Non-biotech Firm	Academic Institution/	Government lab or agence
To conduct research & 3110 Regulatory affairs 3120 Access others' patents 3130 Production/manufactur 3140 Access markets/distrib 3150 Access capital	development (R&D)  string pution channels	Biotech Firm	Non-biotech Firm	Academic Institution/	Government lab or agence
To conduct research & 3110 Regulatory affairs 3120 Access others' patents 3130 Production/manufactur	development (R&D)  string pution channels	Biotech Firm	Non-biotech Firm	Academic Institution/	Government lab or agence

d) Does your firm provide contract services to other firms or organizations?

12500 No Go to question 13

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Intellectual Property				
14. a) Did your firm grant biotechnology related intelle	ectual property (IF	rights to anothe	r firm?	
14000 No So to question 14b)				
Yes For each type of intellectual progranted by country and the total	•	•		of IP rights
Intellectual Property Instrument	Number with Canadian firms	Number with USA firms	Number with other country firms	Revenue from IP licensing in 2001
14100 Licensing Agreement				\$ ,000
Patents				\$ ,000
14120 Other, Please Specify				Φ 000
				\$ ,000
b) Did your firm obtain biotechnology related intell	ectual property ri	ghts from another	firm?	
<sup>14200</sup> No  Go to question 15				
Yes Complete the following table				
Intellectual Property Instrument	Number with Canadian firms	Number with USA firms	Number with other country firms	Cost to your firm of obtaining IP in 2001
14300 Licensing Agreement				\$ ,000
14310 Patents				\$ ,000
14320 Other, Please Specify				
				\$ ,000
15. a) Does your firm have biotechnology related pate  15000 No ▶ Go to question 16  Yes ▶ How many?  Indicate the distribution of biotechnology related pate			patents your firm ha	as by Patent Office
	Canadian Intellectual Property Office (CIPO)	U.S. Patent & Trademark Office (USPTO)	European Patent Office	Other 3
15100 Existing Patents			<del>-</del>	-
15110 Pending Patents				
h) Describe the course of original retard configuration				
<b>b)</b> Provide the number of unique patent applicatio	ns your company Number	submitted in		
15200 2000				
15210 2001				
Section 6 - Firm Characteristics and Finar	ncial Profile			

### Revenues and Research and Development (R&D) Expenditures

16. Please complete the following table. If information is not available please provide a carefully considered estimate. Report for fiscal years and in thousands of dollars (\$,000's). If '0' (ZERO) please indicate, do not leave blanks.

	2	2000	2	001	2	004 Forecast
		0		1		2
16000 Total Firm Sales/Revenues (all sources)	\$	,000	\$	,000	\$	,000
% of revenues from Biotechnology		%		%		%
Total R&D spending	\$	,000	\$	,000	\$	,000
Total spending on Biotechnology R&D	\$	,000	\$	,000	\$	,000
% of Biotechnology R&D spending contracted out		%		%		%

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17.	Does your firm have sales of biotechnology products?  17000 No Go to question 18	
	Yes What percentage of your sales of biotechnology	products came from.
		%
	Direct sales to consumers or distributors	17100
	Products sold to other firms to be used as inputs	17110
Firm F	History	
18.	Is your firm a public firm?	
	18000 No So to question 19	
	Yes What year was the Initial Public Offering (IPO)?	18100
19.	What year was your firm or spin-off established? 19000	
20.	Has your firm merged with another firm? (Include acquisition	of another firm or by another firm)
	20000 No Go to question 21	00400
	Yes What year did the merge take place?	20100
21.	Is your firm a subsidiary of a Multi-National Enterprise (MNE)	?
	21000 No So to question 22	
	Yes	
22. a)	Is your firm a spin-off? A spin-off is defined as a new firm createchnology developed in universities, firms or laboratories.	ated to transfer and commercialize inventions and
	22000 No So to question 23	
	Yes Was your firm a spin-off from University/h	ospital 22100
	Another Bio	tech company 22110
	Non-biotech	
	Governmen	t Agency/lab 22130
	Other, Pleas	se Specify 22140
Raisin	ng Capital	
	A great deal of attention has focused on the ability and the challenges of raising capital. Questions ir information in order to address this critical issue facing	n this section are intended to collect
23. a)	Did your firm attempt to raise capital for biotechnology related	d purposes in 2001?
	23000 No So to question 23h)	
	Yes Were you successful in raising capital?	
	23100 No Go to question 23c)	
		,000
b)	Did you reach your target?	
,		
	Go to question 230)	

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Check all that apply.  Biotechnology product/process not sufficiently developed  Biotechnology product line or portfolio limited in scope  Insufficient specific management skills/expertise  Capital not available due to market conditions  Further product development or proof of concept required  Lender does not fund development projects  Other, Please Specify	al?
Biotechnology product/process not sufficiently developed  Biotechnology product line or portfolio limited in scope  Insufficient specific management skills/expertise  Capital not available due to market conditions  Further product development or proof of concept required  Lender does not fund development projects  23300  23310  23320  23330  Further product development or proof of concept required  23340  Lender does not fund development projects	
Biotechnology product line or portfolio limited in scope  Insufficient specific management skills/expertise  Capital not available due to market conditions  Further product development or proof of concept required  Lender does not fund development projects  23310  23320  23330  Europe 23330  23340  23340  23350	
Insufficient specific management skills/expertise  Capital not available due to market conditions  Further product development or proof of concept required  Lender does not fund development projects  23320  23330  23340  23340  23350	
Capital not available due to market conditions  Further product development or proof of concept required  Lender does not fund development projects  23340  23350	
Further product development or proof of concept required 23340  Lender does not fund development projects 23350	
Lender does not fund development projects  23350	
Other, Please Specify 23360	
d) What sources provided funding?	
% of total raised each source	
Canadian based Venture Capital	%
American based Venture Capital	%
Conventional sources (i.e. banks)	%
Angel Investors/Family	%
Government sources 23440	%
Other, Please Specify 23450	
	%
e) For your most important biotechnology product or process, please indicate the current stage of development.  Stage of Development  R&D  23500  Pre-Clinical  23520	
please indicate the <b>current</b> stage of development.  Stage of Development  R&D  Pre-Clinical  Clinical Trials  23520  23530	
please indicate the <b>current</b> stage of development.  Stage of Development  R&D  23500  Pre-Clinical  Clinical Trials  23520	
please indicate the <b>current</b> stage of development.  Stage of Development  R&D  Pre-Clinical  Clinical Trials  23520  23530	
please indicate the <b>current</b> stage of development.  Stage of Development  R&D 23500  Pre-Clinical 23510  Clinical Trials 23520  Market Entry 23530  For your most important biotechnology product or process,	
please indicate the current stage of development.  Stage of Development  R&D 23500  Pre-Clinical 23510  Clinical Trials 23520  Market Entry 23530  For your most important biotechnology product or process, please indicate total spending since the beginning of development.  Stage of Development  Total spending up to and	

23620 Clinical Trials

<sup>23630</sup> Market Entry

For your most important biotechnology product or process, please estimate the total amount of capital required to complete each stage, as well as the total capital available.

	Stage of Development	Total additional capital required to complete stage	Total capital available to complete stage (include all committed funds)
23700	R&D	\$ ,000	\$ ,000
23710	Pre-Clinical	\$ ,000	\$ ,000
23720	Clinical Trials	\$ ,000	\$ ,000
23730	Market Entry	\$ ,000	\$ ,000

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23. f)	How long do you	anticipate this	capital (committed a	and on hand) last	ting?		
	23800 Years	23810 Mc	onths				
g)	Why did you raise	e or attempt to ı	raise capital? Indica	te each category	y that applies	s to your firm	
	23900 R&D pu	rposes/Expand R	R&D capacity				
	23910 Repay o	current investors					
	23920 Comme	rcialize current R	&D projects				
	23930 Clinical/	regulatory expen	ses				
		p production/man	ufacturing capability				
	23950 Other, F	Please Specify:					
h)	Do you plan on ra	aising capital in	2002?				
	24000 No	Go to question	24				
	○ Yes ▶	How much do y	ou plan to raise?	<\$1,000,000		4010	
				\$1,000,000-\$5,000	0,000	4020 4030	
				>\$5,000,000		1000	
Tax In	centives						
Tax In 24. a)		e biotechnology	/ R&D expenditures	in any of the pre	evious 5 yea	rs?	
			-	in any of the pre	evious 5 yea	rs?	
	Did your firm hav	Go to question and the past 5 ye related activities	-	ofor benefits for bi	otechnology	rs?	
	Did your firm hav	Go to question and the past 5 ye related activities	26 ars did your firm apply s under the Scientific I	for benefits for bi Research and Exp	otechnology erimental		
	Did your firm hav	Go to question in the past 5 yearelated activities Development (\$	ars did your firm apply sunder the Scientific ISR&ED) tax program?  How much did you apply for in 2001?	r for benefits for bi Research and Exp	otechnology	Go to question 24b	
	Did your firm hav	Go to question in the past 5 yearelated activities Development (\$	ars did your firm apply sunder the Scientific ISR&ED) tax program?	for benefits for bi Research and Exp	otechnology erimental		
	Did your firm hav	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{200}\$ Yes	ars did your firm apply sunder the Scientific ISR&ED) tax program?  How much did you apply for in 2001?	o for benefits for bi Research and Exp 24210 \$	otechnology erimental		
	Did your firm hav	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{200}\$ Yes	ars did your firm apply ars did your firm apply under the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?	of for benefits for bickesearch and Exp  24210 \$ cation process	otechnology erimental ,000		
	Did your firm hav	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{200}\$ Yes	ars did your firm apply ars did your firm apply sunder the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of applic	r for benefits for bir Research and Exp 24210 \$ sation process	otechnology erimental ,000		
	Did your firm hav	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{200}\$ Yes	ars did your firm apply ars did your firm apply ander the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of applications are the second applications of the second applications are the second applications.	r for benefits for bir Research and Exp 24210 \$ sation process fility	otechnology erimental ,000 ,000		
	Did your firm hav	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{200}\$ Yes	ars did your firm applys under the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of applicution of eligibout not meet eligibilis	r for benefits for bir Research and Exp 24210 \$ sation process fility	otechnology erimental ,000 ,000 24300 24310 24320		
	Did your firm hav	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{200}\$ Yes	ars did your firm applys under the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of applicution of eligibout not meet eligibilis	r for benefits for bir Research and Exp 24210 \$ ration process fility	otechnology erimental ,000 ,000 24300 24310 24320		
24. a)	Did your firm hav	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{}\) Yes \rightarrow  \text{No} \text{No} \rightarrow	ars did your firm apply ars did your firm apply ander the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of application of the program of the prog	r for benefits for bir Research and Exp 24210 \$ ration process fility	otechnology erimental ,000 ,000 24300 24310 24320		
24. a)	Did your firm have 24100 No Yes >	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{}\) Yes \rightarrow  \text{No} \text{No} \rightarrow	ars did your firm apply ars did your firm apply ander the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of application of the program of the prog	r for benefits for bir Research and Exp 24210 \$ ration process fility	otechnology erimental ,000 ,000 24300 24310 24320		
24. a)	Did your firm have 24100 No Yes Yes	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{}\) Yes \rightarrow  \text{No} \text{No} \rightarrow	ars did your firm apply ars did your firm apply ander the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of application of the program of the prog	r for benefits for bir Research and Exp 24210 \$ ration process fility	otechnology erimental ,000 ,000 24300 24310 24320		
24. a)	Did your firm have 24100 No Yes Yes A	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{}\) Yes \rightarrow  \text{No} \text{No} \rightarrow	ars did your firm apply ars did your firm apply ander the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of application of the program of the prog	r for benefits for bir Research and Exp 24210 \$ ration process fility	otechnology erimental ,000 ,000 24300 24310 24320		
24. a)	Did your firm have 24100 No Yes Yes A	Go to question: In the past 5 ye related activities Development (\$\frac{4200}{}\) Yes \rightarrow  \text{No} \text{No} \rightarrow	ars did your firm apply ars did your firm apply ander the Scientific ISR&ED) tax program?  How much did you apply for in 2001?  Why?  Complexity of application of the program of the prog	r for benefits for bir Research and Exp 24210 \$ ration process fility	otechnology erimental ,000 ,000 24300 24310 24320		

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25.	Did your firm appl	y for any provincial R&D tax b	enefit or incentive?	>		
	25000 Yes					
	○ No ▶	Why did you not apply?				
		Complexity of application proces	25100			
		Uncertainty of eligibility	25110			
		Did not meet eligibility requirement	ents25120			
		Other, Please Specify	25130			
-	s & Exports					
26.	Did your firm expo	ort biotechnology products?				
	<sup>26000</sup> No	Go to question 27				
	○ Yes ▶	Please complete the following ta If '0' (ZERO) please indicate, do		years and in thousand	ds of dollars (\$,000's)	•
			2000	2001	Forecast for 2004	]
			0	1	2	
26100	Total Exports Reve	enues (all sources)	\$ ,000	\$ ,000	\$ ,000	
26110	% export revenues	from Biotechnology	%	%	%	
	Regional Distribut	tion				•
26200	% export revenues	to US	%	%	%	
26210	% export revenues	to Europe	%	%	%	
26220	% export revenues	to Asia	%	%	%	
26230	% export revenues	to other regions	%	%	%	
						1
27.	Did your firm impe	ort bioto obnology producto?				
21.	27000	ort biotechnology products?				
	Yes	Go to question 28  Please complete the following ta	ble Benert for fiscal	years and in thousans	do of dollars (\$ 000's)	
	O res	If '0' (ZERO) please indicate, do		years and in thousand	us of dollars (\$,000 s)	
			2000	2001	Forecast for 2004	
07400			0	1	2	
		ditures (all sources)	\$ ,000	\$ ,000	\$ ,000	
27110	% import expenditu	ires from Biotechnology	%	%	%	
	Regional Distribut	tion	7		1	T
27200	% import expenditu	ires to US	%	%	%	
27210	% import expenditu	ires to Europe	%	%	%	
27220	% import expenditu	ires to Asia	%	%	%	
27230	% import expenditu	ires to other regions	%	%	%	

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follo	ne table below rate the significance of each of the		Ir	nportan	ce	112
	owing strategies on your firm's performance in 2001.	Low 1	2	3	4	High 5
	owledge development strategies					<b>→</b>
	otured and used knowledge obtained from other industry sources h as industry associations, competitors, clients and suppliers	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	
	otured and used knowledge obtained from public research institutions uding universities and government laboratories	$\bigcirc$		$\bigcirc$	$\bigcirc$	$\bigcirc$
<sup>8020</sup> Use	ed and updated databases of scientific information	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
	reloped firm policies and practices for knowledge/intellectual property tection		$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Dev	/eloped/encouraged staff education/upgrading	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$	
and	nducted an Intellectual Property Audit to ensure protection of products I processes at all stages of development siness strategies	0	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
	reased firm size through acquisition, merger or joint venture				$\bigcirc$	
	wnsized operations of the firm	$\bigcirc$				
8120 Ente	ered product trials/adapted products or processes for increased		$\overline{}$			$\overline{}$
	rket penetration					
0130 -	gan new research & development project	$\bigcup$	$\bigcirc$	$\cup$		
Beg		$\overline{}$	$\overline{}$		, ,	
Beg 8140 Expa	panded into foreign markets	$\bigcirc$	$\bigcirc$	$\bigcirc$		
Exp. 3140 Exp. 3150 Other		sses a novel comb	biologica	of genetical entity of	c materia apable of	
Beg 8140 Expand 8150 Other	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living organism	sses a novel comb	biologica roids.	al entity c	apable of	f
Exp. 3140 Exp. 3150 Other	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living organism transferring or replicating genetic material, including sterile organisms	sses a novel comb anism means any ms, viruses and vi Source: Cartag	biologica roids. gena Pro	al entity c	apable of	f
Beg 8140 Expand 8150 Other	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living organism transferring or replicating genetic material, including sterile organism	sses a novel comb anism means any ms, viruses and vi Source: Cartag	biologica roids. gena Pro	al entity c	apable of	f
8140 Expansion Expansion Other	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living organism transferring or replicating genetic material, including sterile organism of the living organism that posses obtained through the use of modern biotechnology. A living organism transferring or replicating genetic material, including sterile organism of the living organism that posses obtained through the use of modern biotechnology. A living organism that posses obtained through the use of modern biotechnology. A living organism that posses obtained through the use of modern biotechnology. A living organism that posses obtained through the use of modern biotechnology. A living organism that posses of transferring or replicating genetic material, including sterile organism that posses obtained through the use of modern biotechnology. A living organism that posses obtained through the use of modern biotechnology. A living organism that posses of transferring or replicating genetic material, including sterile organism that posses obtained through the use of modern biotechnology. A living organism that posses obtained through the use of modern biotechnology. A living organism that posses obtained through the use of modern biotechnology.	sses a novel comb anism means any ms, viruses and vi Source: Cartag	biologica roids. gena Pro	al entity c	apable of	f
Exp. 3140 Exp. 3150 Other	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living orgatransferring or replicating genetic material, including sterile organism of the products based on living following stages?  Research & Development Stage Clinical/Regulatory stage	sses a novel comb anism means any ms, viruses and vi Source: Cartag	biologica roids. gena Pro	al entity c	apable of	f
Esp. 3140 Exp. 3150 Other 29 a)	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living organism transferring or replicating genetic material, including sterile organism of the living following stages?  Pessearch & Development Stage Clinical/Regulatory stage Market stage Total	sses a novel combanism means any sims, viruses and vir	biologica roids. gena Pro	al entity c	apable of	f
Eeg Expand Expan	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living orgatransferring or replicating genetic material, including sterile organism following stages?  Pessearch & Development Stage Clinical/Regulatory stage Market stage Total  Did your firm export living modified organisms in 2001?	sses a novel combanism means any sims, viruses and vir	biologica roids. gena Pro	al entity c	apable of	f
Eeg Expansion	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living orgation transferring or replicating genetic material, including sterile organism of the serile organism orga	sses a novel combanism means any sims, viruses and vir	biologica roids. gena Pro ms does	at entity contocol on	apable of Biosafety	f
8140 Exp. 8150 Other 29 a)	Does your firm develop, produce or sell Living Modified Orga  Living modified organism means any living organism that posses obtained through the use of modern biotechnology. A living orgativansferring or replicating genetic material, including sterile organism following stages?  Pessearch & Development Stage Clinical/Regulatory stage Market stage Total  Did your firm export living modified organisms in 2001?	sses a novel combanism means any sims, viruses and vir	biologica roids. gena Pro ms does	otocol on  your firm	apable of Biosafety	each of t

Thank you for your assistance.

Return the questionnaire in the accompanying self addressed prepaid envelope.

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