# Annual Report on Findings of Infectious Agents in Japan 2004

## CONTENTS

Foreword	7
Surveillance System	8
The Topics of the Month of IASR	21
Tables	
I. Reports on Isolation of Bacteria (including Fungi, Spirochetes and Protozoa)	47
II. Reports on Isolation/Detection of Viruses (including Rickettsiae and Chlamydiae)	103
Appendix	137

# This report was prepared by Editorial Committee of Infectious Agents Surveillance Report, 2005:

Chairman:	Nobuhiko OKABE, M.D.
Viaa ahairman:	Director, Infectious Disease Surveillance Center, National Institute of Infectious Diseases
vice-chairman.	Deputy Director-General National Institute of Infectious Diseases
	Director, Department of Bacteriology, National Institute of Infectious Diseases
	Yoshichika ARAKAWA, M.D.
	Director, Department of Bacterial Pathogenesis and Infection Control, National Institute
	of Infectious Diseases
	Takuro ENDO, Ph.D.
	Director, Department of Parasitology, National Institute of Infectious Diseases
	Chief Laboratory of Reservoir Control of Zoonoses Department of Veterinary Science
	National Institute of Infectious Diseases
	Norihisa ISHII, M.D.
	Director, Department of Bioregulation, Leprosy Research Center, National Institute of
	Infectious Diseases
	Kenichiro IIO, Ph.D.
	Center, Division of Dacteriological Laboratory Training, infectious Disease Survemance Center, National Institute of Infectious Diseases
	Mikio KIMURA, M.D.
	Chief, Surveillance and Information Division, Infectious Disease Surveillance Center,
	National Institute of Infectious Diseases
	Toshio KISHIMOTO, M.D.
	Chief, Laboratory of Rickettsia & Chlamydia, Department of Virology I, National
	Mutsuo KOBAVA SHL Ph D
	Director, Department of Medical Entomology, National Institute of Infectious Diseases
	Ichiro KURANE, M.D.
	Director, Department of Virology I, National Institute of Infectious Diseases
	Tatsuo MIYAMURA, M.D.
	Director, Department of Virology II, National Institute of Infectious Diseases
	Director Division of International Cooperation National Insultitute of Infectious
	Diseases
	Osamu NISHIO, D.V.M
	Chief, Division of Virological Laboratory Training, Infectious Disease Surveillance
	Center, National Institute of Infectious Diseases
	Itesutaro SAIA, M.D.
	Miharu SHINTANI, Ph D
	Department of Bacterial Pathogenesis and Infection Control. National Institute of
	Infectious Diseases
	Kiyosu TANIGUCHI, M.D.
	Chief, Intelligence and Policy Planning Division, Infectious Disease Surveillance
	Center, National Institute of Infectious Diseases
	Masalo IASHIKO, M.D. Director Department of Virology III National Institute of Infectious Diseases
	Keiko TAYA M D
	Chief, Immunization Program Division, Infectious Disease Surveillance Center,
	National Institute of Infectious Diseases
	Jun TERAJIMA, D.V.M.
	Chief, Laboratory of Enteric Infection I, Department of Bacteriology, National Institute
	of infectious Diseases
	Infectious Disease Surveillance Center, National Institute of Infectious Diseases

Advisory Board:		
•	Sakae INOUYE, M.D.	
	Emeritus Researcher, N	Vational Institute of Infectious Diseases
	Takahiro INOUE, M.D.	
	Technical Official, Safe of Health, Labour and	ety Division, Pharmaceutical and Food Safety Bureau, Ministry Welfare
	Yumiko KANARI, M.D.	
	Technical Official, Tub Health, Labour and We	erculosis and Infectious Diseases Control Division, Ministry of Ifare
	Takeshi KURATA, M.D.	
	Director-General, Natio	onal Institute of Infectious Diseases
	Satoshi MIYAKE, M.D.	
	Director, Planning and	Coodination, National Institute of Infectious Diseases
	Yuji NAKANIWA, D.V.N	И.
	Technical Official, Dep	bartment of Food Safety, Ministry of Health, Labour and Welfare
	Koji TAKIMOTO, M.D.	
	Technical Official, Tub	erculosis and Infectious Diseases Control Division, Ministry of
	Health, Labour and We	NTARE NYM
	Snigeki YAMAMOTO, D	J. V. MI. f Diamadical Food Descent National Institute of Health
	Sciences	i Biomedical Food Research, National Institute of Health
	Hiroshi YOSHIKURA M	( D
	Emeritus Researcher N	Jational Institute of Infectious Diseases
Statistics, data pro	ocessing and graphs:	
	Nobuko KATO	Motoko NOJI
	Yuki TADA	Mariko TOKUNAGA

Kazuyo YAMASHITA Masae YOSHIKAWA Surveillance and Information Division, Infectious Disease Surveillance Center, National Institute of Infectious Diseases

# The Reporting System of Findings of Infectious Agents in Japan is operating under Advisory Committee, 2005 organized by the members of the Association of Public Health Laboratories for Microbiological Technology:

Chairman: Nobuhiko OKABE, M.D., Director, Infectious Disease Surveillance Center, National Institute of Infectious Diseases Kazuo AKIYAMA, Miyagi Prefectural Institute of Public Health and Environment Yoshichika ARAKAWA, National Institute of Infectious Diseases Ritsushi FUJII, Okayama Prefectural Institute of Environmental Science and Public Health Kenichi HAYASHI, Shiga Prefectural Institute of Public Health and Environmental Sciences Kenichiro ITO, National Institute of Infectious Diseases Mikio KIMURA, National Institute of Infectious Diseases Ichiro KURANE, National Institute of Infectious Diseases Kazuo MATSUMOTO, Fukui Prefectural Institute of Public Health and Environmental Science Tatsuo MIYAMURA, National Institute of Infectious Diseases Masao OGAWA, Oita Prefectural Institute of Health and Environment Masaaki SUGIEDA, Shizuoka Institute of Environment and Hygine Masato TASHIRO, National Institute of Infectious Diseases Jun TERAJIMA, National Institute of Infectious Diseases Haruo WATANABE, National Institute of Infectious Diseases

### **Cooperative Surveillance Projects and Associations:**

National Epidemiological Surveillance of Infectious Diseases National Epidemiological Surveillance of Vaccine-Preventable Diseases Research Project on a Reference System for Microbiological Examination Research Group for Enteric Infection in Japan Association of Public Health Laboratories for Microbiological Technology Surveillance System of Typhoid and Paratyphoid Fevers

# List of prefectural and municipal public health institute participating in the reporting system, 2004

Code	Prefecture	Institute
number	/city	
011	Hokkaido P.	Hokkaido Institute of Public Health
012	Sapporo C.	Sapporo City Institute of Public Health
013	Hakodate C.	Hakodate City Institute of Public Health
021	Aomori P.	Aomori Prefectural Institute of Public Health and Environment
031	Iwate P.	Research Institute for Environmental Sciences and Public Health of Iwate
		Prefecture
041	Miyagi P.	Miyagi Prefectural Institute of Public Health and Environment
042	Sendai C.	Sendai City Institute of Public Health
051	Akita P.	Akita Research Center for Public Health and Environment
061	Yamagata P.	Yamagata Prefectural Institute of Public Health
071	Fukushima P.	Fukushima Institute of Public Health
081	Ibaraki P.	Ibaraki Prefectural Institute of Public Health
091	Tochigi P.	Tochigi Prefectural Institute of Public Health and Environmental Science
101	Gunma P.	Gunma Prefectural Institute of Public Health and Environmental Sciences
111	Saitama P.	Saitama Institute of Public Health
121	Chiba P.	Chiba Prefectural Institute of Public Health
122	Chiba C.	Chiba City Institute of Health and Environment
131	Tokvo M.	Tokyo Metropolitan Institute of Public Health
141	Kanagawa P.	Kanagawa Prefectural Institute of Public Health
142	Yokohama C.	Yokohama City Institute of Health
143	Kawasaki C.	Kawasaki City Institute for Public Health
144	Yokosuka C.	Yokosuka City Institute of Public Health
145	Sagamihara C.	Sagamihara City Institute of Public Health
151	Niigata P.	Niigata Prefectural Institute of Public Health and Environmental Sciences
152	Niigata C.	Niigata City Institute of Public Health and Environment
161	Tovama P.	Tovama Institute of Health
171	Ishikawa P.	Ishikawa Prefectural Institute of Public Health and Environmental Science
181	Fukui P.	Fukui Prefectural Institute of Public Health and Environmental Science
191	Yamanashi P.	Yamanashi Institute for Public Health
201	Nagano P.	Nagano Environmental Conservation Research Institute
211	Gifu P.	Gifu Prefectural Institute of Health and Environmental Sciences
212	Gifu C.	Hygienic Laboratory of Gifu City
221	Shizuoka P.	Shizuoka Institute of Environment and Hygiene
222	Shizuoka C.	Shizuoka City Institute of Public Health
223	Hamamatsu C.	Hamamatsu City Institute of Public Health
231	Aichi P.	Aichi Prefectural Institute of Public Health
232	Nagoya C.	Nagoya City Public Health Research Institute
241	Mie P.	Public Health and Environment Research Division, Mie Prefectural Science
		and Technology Promotion Center
251	Shiga P.	Shiga Prefectural Institute of Public Health and Environmental Science
261	Kyoto P.	Kyoto Prefectural Institute of Public Health and Environment
262	Kyoto C.	Kyoto City Institute of Health and Environmental Sciences
271	Osaka P.	Osaka Prefectural Institute of Public Health
272	Osaka C.	Osaka City Institute of Public Health and Environmental Sciences
273	Sakai C.	Sakai City Institute of Public Health
281	Hyogo P.	Hyogo Prefectural Institute of Public Health and Environmental Sciences
282	Kobe C.	Kobe City Institute of Health
283	Himeji C.	Himeji City Institute of Environment and Health
284	Amagasaki C.	Amagasaki City Institute of Public Health
291	Nara P.	Nara Prefectural Institute for Hygiene and Environment
301	Wakayama P.	Wakayama Prefectural Research Center of Environment and Public Health
302	Wakayama C.	Wakayama City Institute of Public Health
311	Tottori P.	Tottori Prefectural Institute of Public Health and Environmental Science
321	Shimane P.	Shimane Prefectural Institute of Public Health and Environmental Science
331	Okayama P.	Okayama Prefectural Institute for Environmental Science and Public Health
341	Hiroshima P.	Hiroshima Prefectural Health Environment Center
342	Hiroshima C.	Hiroshima City Institute of Public Health
351	Yamaguchi P.	Yamaguchi Prefectural Research Institute of Public Health
361	Tokushima P.	Tokushima Prefectural Institute of Public Health and Environmental Sciences

371	Kagawa P.	Kagawa Prefectural Research Institute for Environmental Sciences and
		Public Health
381	Ehime P.	Ehime Prefectural Institute of Public Health and Environmental Science
391	Kochi P.	The Public Health Institute of Kochi Prefecture
401	Fukuoka P.	Fukuoka Institute of Health and Environmental Sciences
402	Fukuoka C.	Fukuoka City Institute for Hygiene and the Environment
403	Kitakyushu C.	Kitakyushu City Institute of Environmental Sciences
411	Saga P.	Saga Prefectural Institute of Public Health and Pharmaceutical Research
421	Nagasaki P.	Nagasaki Prefectural Institute of Public Health and Environmental Sciences
422	Nagasaki C.	Nagasaki Municipal Public Health and Environment Laboratory
431	Kumamoto P.	Kumamoto Prefectural Institute of Public Health and Environmental Science
432	Kumamoto C.	Kumamoto City Environmental Research Institute
441	Oita P.	Oita Prefectural Institute of Health and Environment
451	Miyazaki P.	Miyazaki Prefectural Institute for Public Health and Environment
461	Kagoshima P.	Kagoshima Prefectural Institute for Environmental Research and Public Health
471	Okinawa P.	Okinawa Prefectural Institute of Health and Environment

P.: Prefecture C.: City M.: Metropolitan

#### List of quarantine stations participating in the reporting system, 2004

Chitose Airport Branch Office, Otaru Quarantine Station Narita Airport Quarantine Station Chubu Airport Branch Office, Nagoya Quarantine Station Kansai Airport Quarantine Station Hiroshima Airport Branch Office, Hiroshima Quarantine Station Fukuoka Airport Branch Office, Fukuoka Quarantine Station Naha Airport Branch Office, Naha Quarantine Station



Fig. 1. Code number of prefectural and municipal public health institutes participating in the reporting system, 2004.

## Foreword

This is the 2004 Annual Report on Findings of Infectious Agents in Japan. Since the National Epidemiological Surveillance of Infectious Diseases was established in July 1981, it has been providing a valuable source of information necessary to take effective countermeasures against the prevention and control of infectious diseases in Japan. Under this program before April 1999, 27 infectious diseases were specified to be surveyed at designated sentinels in each prefecture, and the epidemiological data on both patients and pathogens has been gathered through nationwide surveillance network systems.

Infectious Disease Surveillance Center (IDSC) under National Institute of Infectious Diseases (NIID) serves as an information center where laboratory data on pathogens are collected, analyzed and distributed for public use in collaboration with various institutions such as prefectural and municipal public health institutes, quarantine stations, designated hospitals and health centers. Some departments other than IDSC of NIID function also as national reference laboratories for standardization of the diagnostic methods and reagents used at central and regional public health laboratories. It is noteworthy that such reference activities have strengthened cooperative relationships among many organizations participating in this surveillance program and that the quality of laboratory data obtained through the surveillance network has been much improved during the last decade.

In April 1999, the Law Concerning the Prevention of Infectious Diseases and Medical Care for Patients of Infections (the Infectious Diseases Control Law) was enacted (see p. 16-19 of this issue) and partially amended on November 5, 2003 (http://idsc.nih.go.jp/iasr/25/287/tpc287.html). Relatively important 58 infectious diseases newly specified to be notified by all physicians to nearby health centers (see p. 19 of this issue). The weekly numbers of cases with common infectious diseases at sentinel clinics has been also reported to the health centers, in continuation of the surveillance system started in 1981. Those data are sent to IDSC through a national computer network (WISH-NET) organized by Ministry of Health and Welfare in 1987.

In parallel with the disease surveillance, data on the infectious agents detected from some of the patients at prefectural and municipal public health institutes and quarantine stations are also sent to IDSC through another system on the WISH-NET installed in 1997. This on-line system has been improved since January 2000 in accordance with the above Law. This annual report deals with the data on the infectious agent surveillance in 2004.

NIID together with Tuberculosis and Infectious Diseases Control Division, Ministry of Health, Labour and Welfare publishes a monthly report, *Infectious Agents Surveillance Report (IASR)* from 1983. To address the need for wider distribution and utilization of the *IASR* information, the special review article on a selected topic has been translated into English since January 1993 and distributed to relevant institutions including major overseas institutions every month, together with the original article written in Japanese. The English articles, THE TOPIC OF THIS MONTH from the January issue of 2005 to the December issue of 2005 are contained in this 2004 annual report. We hope the publication of this annual report will provide useful information to clinical and research organizations and also to public health administrators.

Finally, I would like to express my sincere thanks to all the organizations and individuals participating in this surveillance program for their sustained efforts and enthusiastic cooperation which have made this publication possible.

Nobuhiko Okabe, M.D. & Ph.D. Director, Infectious Disease Surveillance Center Chairman, Editorial Committee of Infectious Agents Surveillance Report National Institute of Infectious Diseases

## Annual Report on Findings of Infectious Agents in Japan, 2004

#### 1. The surveillance system for collecting and distributing the information of infectious agents

#### History and Organization

Collection and distribution of the information on laboratory findings of infectious agents were initiated by the working group of the Research Project for Development of a Surveillance System of Pathogenic Microbes in Japan (1979-1982) (leader Hiromasa Inoue, Former Director, Aichi Prefectural Institute of Public Health). As a result of the cooperation among prefectural and municipal public health institutes (PHIs) and the National Institute of Infectious Diseases (NIID, former National Institute of Health), the infectious agent surveillance system was established.

In 1981, when the National Epidemiological Surveillance of Infectious Diseases (NESID) program was inaugurated, both the pathogen and patient information systems were considered to be inseparable like the two wheels of a cart (Fig. 2). However, the report of pathogen information from laboratory comes out usually after that of patient information, and technical and professional quality controls were needed for their reliability. Therefore, the function of this system, being independent of patient information, was undertaken by two committees, the Editorial Committee for Infectious Agents Surveillance Report (IASR), NIID, and the Committee for Laboratory-based Information, the Association of Public Health Laboratories for Microbiological Technology. The daily function of the information center has been undertaken by the Infectious Disease Surveillance Center (IDSC), NIID.

A computer network was installed for the patient information system in 1987. That for pathogen information was investigated by a working group of the Research Project for Development of a Surveillance System of Pathogenic Microbes in Japan (leader, Makoto Ohashi, Former Director, Tokyo Metropolitan Research Laboratory of Public Health) for three years (1988-1991), but the plan was not conducted for a while. After a three-year trial (1994-1996), the network started to function in January 1997 as the National Electronic Data Transfer System for Laboratory Findings of Infectious Agents utilizing the Wide-area Information-exchange System for Health and Welfare Administration (WISH) by the Ministry of Health, Labour and Welfare (MHLW, former Ministry of Health and Welfare) and the Value Added Network (VAN) (Fig. 2). As a result of the on-line system between PHIs/quarantine stations and NIID, rapid information exchange by electronic mail/board has become possible.

In compliance with the Law concerning the Prevention of Infectious Diseases and Medical Care for Patients of Infections (hereafter referred to as the Infectious Diseases Control Law) enacted in April 1999, the on-line system has revised in January 2000 (Version 4).

#### Contents and Collection of Information

The infectious agents surveillance information deals with such pathogenic agents as bacteria, fungi, spirochetes, protozoa, viruses, rickettsiae, and chlamydiae. The data are collected in two large categories for the convenience; bacterial pathogens (including fungi, spirochetes, and protozoa) and viral pathogens (including rickettsiae and chlamydiae).



Fig. 2. Network of National Epidemiological Surveillance of Infectious Diseases in Japan.

Through the on-line system, the information on bacterial pathogens is collected from (a) PHIs and health centers (HCs) and (b) quarantine stations, and that on viral pathogens from PHIs. In 2004, PHIs in 47 prefectures and 25 designated cities and eight quarantine stations listed on p. 4-5 cooperated.

The figures for bacterial pathogens from (a) and (b) are totaled separately as follows.

(a) PHIs/HCs: Results of tests conducted for administrative purposes on a public health standpoint are reported. Investigations mainly of infectious disease epidemics and food poisoning outbreaks, and tests for safety assessment of food, for environmental pollution, and for infectious agent surveillance under the NESID are included.

A PHI, playing two roles, the reference center and the infectious agents surveillance center in the district, not only performs the laboratory tests but also collects the results of laboratory tests conducted by HCs in the district. These results are concluded in the report form A from PHIs/HCs (Form A, Fig. 3 on p. 10-12) and are sent on a monthly basis to IDSC, NIID. The more detailed results of serotyping of Salmonella and group A Streptococcus has been included in form A to collect on a monthly basis through the on-line system since January 2000 (before 2000, these serotyping data were collected on a yearly basis and sent out by conventional mail). In addition, PHIs provide also information on individual bacterial isolation in the Individual Report (Form B, Fig. 4 on p. 13) including serotypes, toxin types, the date of specimen collection, specimens, and age, gender and clinical symptoms of EHEC/VTEC-infected cases and other important patients. They provide also the Outbreak Report (Form C, Fig. 5 on p. 14), Food Examination Report and Surveillance Report for Environmental Sources and for Animals (Form D, Fig. 6 on p. 15). On the other hand, form E categorized by source of specimen used to be sent from PHIs on a monthly basis to IDSC, NIID from1990 to collect the results of tests conducted for the purpose of diagnosis and treatment of infectious disease patients at general clinical institutions (GCIs), is tabulated at each PHI from 2000 for the analysis of the trend in each district.

(b) Quarantine stations: The data of examinations for the quarantine infectious diseases in compliance with the Quarantine Law, particularly of enteric infections, of overseas travelers (imported cases) are reported. Each quarantine station send individual bacterial information with patient's age, gender, clinical symptoms, and estimated districts of acquiring infection in the Individual Report (Form B, Fig. 4 on p. 13) and that of traveler's diarrhea outbreak in the Outbreak Report (Form C, Fig. 5 on p. 14).

The virus reports from PHIs are the results of etiological diagnosis (including those conducted under NESID), the National Epidemiological Surveillance of Vaccine-Preventable Diseases, and the periodic sentinel surveys and specified research projects of the individual PHIs. When examinations are completed, PHIs input the individual data of isolated/ detected viruses in the Individual Report for virus detections from human sources (Form B, Fig. 4 on p. 13) to be sent immediately. The virus report includes the patient's age, gender, and clinical symptoms, and the date of specimen collection, specimens and the method of examination. If a influenza virus isolated, PHI reports the strain name and the results of hemagglutination-inhibition test of the strain together with the virus report. The strain surveillance kits are supplied to PHIs by the Department of Virology III, NIID, the National and WHO Collaborating Center for Influenza.

When an incident of such viral disease as nonbacterial gastroenteritis involving two or more cases occurs in a community, its data are input in the Outbreak Report (Form C, Fig. 5 on p. 14), to be sent for rapid exchange of information. Moreover, the viruses isolated from food, animals, or the environment (river water and sewage) are input into the Reports of Food Examinations and Surveillance for Environmental Sources and for Animals (Form D, Fig. 6 on p. 15) to be provided.

Separately from the on-line system, the Laboratory of Enteric Infections II, the Department of Bacteriology, NIID submits phage typing results of *Salmonella* Typhi and *S*. Paratyphi A isolates (see p. 93-96). The isolates are sent from various institutions to NIID in accordance with the notice #788, November 16, 1966, "on the implementation of countermeasures for typhoid and paratyphoid fevers," issued by the director of the Bureau of Public Health, MHLW.

In addition, a total of 16 infectious diseases hospitals (IDHs) in 13 cities report the results of etiological diagnoses of all inpatients with diarrheal diseases together with detailed information on patient's ages, clinical symptoms, estimated districts of acquiring infection, and the drug susceptibilities fulfilled on individual cards (mark sheet) and send out by conventional mail. The results are shown on p. 97-101.

#### **Distribution of Information**

The collected infectious agent surveillance information is daily updated at IDSC, NIID and electrically published on the Infectious Agents Surveillance website constructing on WWW-WISH, the advanced WISH intra-network, including Forum for Bacteria and that for Virus. Overview of Individual Reports and Outbreak Reports are searchable with link to the disease diagnosed sorted by the bacterial/viral pathogen. At the end of every month, the primary information from the whole country is converted into the monthly-feed-back-data file, which is distributed to the source of information, PHIs and quarantine stations from the download site on WWW-WISH. Such secondary data of the whole country as the breakdown tables are published monthly in IASR, which is distributed to cooperating and other institutions concerned. In IASR, THE TOPIC OF THIS MONTH, an analysis of infectious agent and patient information, is published in both Japanese and English. This issue contains the articles from January 2005 to December 2005 (see p. 21-45). Since August 1996, the website of IDSC has been opened on the internet. To make the information public, the HTML version of *IASR* is retrievable from the *IASR* URL (http://idsc. nih.go.jp/iasr/). The homepage also includes many updated tables and graphic data on pathogen detection for easy comprehension of the infectious agent trend.

In addition, isolation of influenza viruses is reported to the WHO Influenza Center. The data is uploaded on the website of FluNet (http://www.who.int/flunet) installed by WHO.

#### Publication of Annual Reports

The infectious agents information is highly valued as a material in Japan and other countries. To keep the precious data, annual reports have been edited. During the four years of 1979-1982, yearly information have been published in *the Annual Report of Laboratory-Confirmed Agents for Infectious Diseases in Japan* (the Research Project for Development of a Surveillance System of Pathogenic Microbes in Japan). Since 1983, it has been published in *Annual Report of the NESID Program* in Japanese and an English version in

### REPORTED BY MONTH OF SPECIMEN COLLECTION

CODE	SPECIES, GROUPS, SEROVARS*	TOTAL NUMBER	IMPORTED
NUMBER		OF ISOLATES	CASES**
		FROM HUMAN SOURCES	
100-001	Verotoxin-producing <i>Escherichia coli</i> (EHEC/VTEC)		
100-002	Enterotoxigenic <i>Escherichia coli</i> (ETEC)		
100-003	Enteroinvasive Escherichia coli (EIEC)		
100-004	Enteropathogenic Escherichia coli (EPEC)		
100-005	Other diarrhegenic Escherichia coli		
134 001			
134-001			
134-002	Salmonella Paratyphi A		
104	Salmonella 04*		
105	Salmonella 07*		
106	Salmonella 08*		
107	Salmonella 09*		
108	Salmonella 09, 46 *		
109	Salmonella 03, 10 *		
110	Salmonella 01, 3,19 *		
111	Salmonella 011*		
112	Salmonella 013*		
113	Salmonella 06 14*		
114	Salmonella 016*		
115	Salmonolla 017*		
110	Salmonolla 011		
110			
118	Salmonella 028*		
119	Salmonella 030*		
120	Salmonella 035*		
121	Salmonella 038*		
122	Salmonella 039*		
123	Salmonella 040*		
124	Salmonella 041*		
125	Salmonella 042*		
126	Salmonella 043*		
127	Salmonella 045*		
128	Salmonella 047*		
120	Salmonella 047		
129			
130			
131	Salmonella 051*		
134-003	Salmonella other groups		
134-004	Salmonella group unknown		
201-001	Listeria monocytogenes		
202-001	Yersinia enterocolitica		
202-002	Yersinia pseudotuberculosis		
204-001	Vibrio cholerae 01: El Tor, Ogawa, CT (+)		
204-002	Vibrio cholerae 01: El Tor, Ogawa, CT (-)		
204-003	Vibrio cholerae 01: El Tor, Inaba, CT (+)		
204-004	Vibrio cholerae 01: El Tor, Inaba, CT (-)		
204-005	Vibrio cholerae 01: Classical Ogawa CT (+)		
204-005	Vibrio cholerae OT: Classical, Ogawa, CT (+)		
204-000	Vibria abalaraa O1, Classical, Oyawa, C1 (-)		
204-007			
204-008	Vibrio criolerae OT: Classical, Inaba, CT (-)		
204-011			
204-009	Vibrio cholerae 0139, CT (+)		
204-010	Vibrio cholerae 0139, CT (-)		
204-012	Vibrio cholerae non-01 & 0139		
204-013	Vibrio parahaemolyticus		
204-014	Vibrio fluvialis		
204-015	Vibrio mimicus		
219-001	Aeromonas hydrophila		
219-002	Aeromonas sobria		
219-002	Aeromonas hydronhila/sobria		
222 001	Plasiamanas chigallaidas		
222-001	ricsionionas singenoues		
223-001	campyiopacter jejuni		
223-002	campylobacter coll		
223-003	Campylobacter jejuni/coli		
226-001	Staphylococcus aureus		
227-001	Clostridium perfringens		
227-002	Clostridium botulinum E		
227-003	Clostridium botulinum non-E		
230-001	Bacillus cereus		
	1		

\*\* IMPORTED CASES INCLUDED IN THE TOTAL

Fig. 3. Reporting form A for isolation of pathogenic bacteria (prefectural and municipal public health institutes) from human sources.

CODE	SPECIES, GROUPS, SEROVARS*	TOTAL NUMBER	IMPORTED
NUMBER		OF ISOLATES	CASES**
		FROM HUMAN SOURCES	0,1020
301-001	Shigella dysenteriae 1		
301-002	Shigella dysenteriae 2		
301-003	Shigella dvsenteriae 3		
301-004	Shigella dysenteriae 4		
301-005	Shigella dysenteriae 5		
301-006	Shigella dysenteriae 6		
301-007	Shigella dysenteriae 7		
301-008	Shigella dysenteriae 8		
301-009	Shigella dysenteriae 9		
301-010	Shigella dysenteriae 10		
301-011	Shigella dysenteriae 11		
301-012	Shigella dysenteriae 12		
301-090	Shigella dysenteriae other serovars		
301-099	Shigella dysenteriae serovar unknown		
302-001	Shigella flexneri 1a		
302-002	Shiqella flexneri 1b		
302-003	Shigella flexneri 1		
302-004	Shiqella flexneri 2a		
302-005	Shigella flexneri 2b		
302-006	Shiqella flexneri 3a		
302-007	Shigella flexneri 3b		
302-008	Shigella flexneri 4a		
302-009	Shigella flexneri 4b		
302-010	Shigella flexneri 4		
302-011	Shigella flexneri 5a		
302-012	Shigella flexneri 5b		
302-013	Shigella flexneri 6		
302-014	Shigella flexneri var. X		
302-015	Shigella flexneri var. Y		
302-090	Shigella flexneri other serovars		
302-099	Shigella flexneri serovar unknown		
303-001	Shigella boydii 1		
303-002	Shigella boydii 2		
303-003	Shigella boydii 3		
303-004	Shigella boydii 4		
303-005	Shigella boydii 5		
303-006	Shigella boydii 6		
303-007	Shigella boydii 7		
303-008	Shigella boydii 8		
303-009	Shigella boydii 9		
303-010	Shigella boydii 10		
303-011	Shigella boydii 11		
303-012	Shigella boydii 12		
303-013	Shigella boydii 13		
303-014	Shigella boydii 14		
303-015	Shigella boydii 15		
303-016	Shigella boydii 16		
303-017	Shigella boydii 17		
303-018	Shigella boydii 18		
303-090	Shigella boydii other serovars		
303-099	Shigella boydii serovar unknown		
304-001	Shigella sonnei		
304-002	Shigella species unknown		
401-001	Entamoeba histolytica		
402-001	Cryptosporidium spp.		
403-001	Giardia lamblia		

\*\* IMPORTED CASES INCLUDED IN THE TOTAL

Fig. 3.-Continued-1.

CODE	SPECIES, GROUPS, SEROTYPES*	TOTAL NUMBER	IMPORTED
NUMBER		OF ISOLATES	CASES**
		FROM HUMAN SOURCES	
501	Streptococcus A*		
502-001	Streptococcus B		
502-002	Streptococcus C		
502-003	Streptococcus G		
502-004	Streptococcus other groups		
502-005	Streptococcus group unknown		
502-006	Streptococcus pneumoniae		
508-001	Corynebacterium diphtheriae		
509-001	Bordetella pertussis		
510-001	Legionella pneumophia		
510-002	Legionella others		
512-001	Mycobacterium tuberculosis		
512-002	Mycobacterium bovis		
512-003	Mycobacterium avium- intracellulare complex		
515-001	Haemophilus influenzae type b		
515-002	Haemophilus influenzae other types		
517-001	Klebsiella pneumoniae		
518-001	Neisseria meningitidis		
518-002	Neisseria gonorrhoeae		
520-001	Leptospira spp.		
521-001	Borreria burgdorferi		
522-001	Mycoplasma pneumoniae		
	Total		

\*\* IMPORTED CASES INCLUDED IN THE TOTAL

Fig. 3.-Continued-2.

REPORTED BY CATEGORY OF DISEASE		_	DATE OF REPO	DRTING(YYMMD	D) □VIRUS □RICK	(ETTSIA/CHL	AMYDIA
ID No.		_					
INFECTIOUS AGENT*		_	DATE OF SPEC	CIMEN COLLECT	TON (YYMMDD)		
*TYPING RESULT OF BACTERIA				BIOCHEMICAL	FEATUER, etc		
SEROTYPE/SEROVAR							
TOXIN TYPE/PATHOGENICITY							
PATIENT DATA							
GENDER		AGE IN YEARS			AGE IN MONTHS		
DIAGNOSIS	HUN				COR ASYMPTOM	ATIC	
DATE OF ONSET (YYMMDD)				PROGNOSIS			
SOURCE OF SPECIMEN							
		ITOPST (ORGAN	:				
		PINAL FLUID			LE	LOID, etc)	
□EYE				GENITAL			
	□OTHER (		)				
		CASE					
	FADACHE	CASE	□SHOCK (← I	OW BLOOD PR	ESSURE, CIRCULA	ATORY DISTU	RBANCE)
HEALTHY/NO ILLNESS			□GASTRO-IN	TESTINAL DISE	ASE ( DIARRHEA		G)
□FEVER( ℃)				ARRHEA		PAIN	
□ FEBRILE CONVULSION			□KERATITIS		TIVITIS DKE	RATOCONJU	NCTIVITIS
DISEASE OF MUSCLES & JOINTS	5 (←ARTHRITIS)				□ ALTERED CON	ISCIOUSNESS	
UPPER RESPIRATORY TRACT INF	ECTION			TIS		ATHY	
LOWER RESP. TRACT INFECTION	N (□PNEUMONIA □BRO	NCHITIS)					
				CULAR DISORD	ER		
				SORDER			
SALIVARY GLANDULAR DISEAS	E		RENAL DISC	RDER			
□OTHER (	)			ARY DISEASE			
			YO (PASSAGE	)		OTHER	
SENSITIVE CELLS		(PASSAGE )	SENSITIVE CEI	LLS	(1	PASSAGE )	
ANTIGEN DETECTION	A □EIA	□RPHA	□LA	□PA		OTHER	-
GENE DETECTION NON							
			⊔PCK+HYBKII	JIZATION	LPCK+SEQUEN	SING	
□OTHER (	)						
	PIDEMIC O (DIACEOEDE		JIBREAK		(PLACE	)	
	ES COUNTRY/AF	REA		COUNTRY/A	REA		
	COUNTRY/AF	REA		COUNTRY/A	REA		
	TRAVEL PERI	OD FROM(YYM	MDD)	TO (YYMMD	D)		
	))	_	VACCINE				
DATE OF VACUNATION (TIMMDL	,,						
REMARKS							
-							

RECORD No.

Fig. 4. Reporting form B for individual case of pathogen isolation/detection from human sources.

CATEGRA OR VERSAGE       IACTERIA OR VERS         DAGONS       IDAGON         DAGONS       IDAGON         DAGONS       IDAGON         DAGONS       IDAGON         PERIOD       RIGAL TERIA OR VERSAGE         JUSPECTED PLACE OF INFECTION/EATING FOOD       IDAGON         SUSPECTED PLACE OF INFECTION/EATING FOOD       IDAGON         SUSPECTED PLACE OF INFECTION/EATING FOOD       IDAGON         NUMBER OF CONSUMERS       NUMBER OF POSITIVE CASE         NUMBER OF POSITIVE CASE       IBOCHEMICAL FEATURE, etc         SECONDARY OF DAGONAR       IDON TYPE/CARING CASE         SECONDARY OF TOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS       IBOCHEMICAL FEATURE, etc         SECOND TYPE/SERIOVAR       IDON TYPE/SERIOVAR         TORN TYPE/PATHOGENCITY       IBOCHEMICAL FEATURE, etc         SECOND TYPE/SERIOVAR       IDON TYPE/SERIOVAR         TORN TYPE/PATHOGENCITY       IDON TYPE/SERIOVAR         MEECTIDUS AGENT       IDON TYPE/SERIOVAR	REPORTED BY		DATE OF REP	ORTING(YYMMD	D)	
SUSPECTED ROUTE OF PRECTOR       D.No.         PLACE OF NOIDENT OCCURED       DARANGES         DARADISS       TO (YYMADD)         SUSPECTED PLACE OF PREMANG FOOD       TO (YYMADD)         SUSPECTED PLACE OF PREMANG FOOD       SUSPECTED PLACE OF PREMANG FOOD         SUSPECTED PLACE OF PREMANG FOOD       MARE         SUSPECTED PLACE OF PRETCONVERTING FOOD       MARE         SUSPECTED PLACE OF PRETCONVERTING FOOD       MARE         SUSPECTED PLACE OF PRETCONVERTING FOOD       MARE         SUSPECTED CAUSE       NUMBER OF PATIENTS         NUMBER OF CONSUMERS       NUMBER OF PATIENTS         SUSPECTED CAUSE       FROM TO         NUMBER OF EXAMINED       Int         SECONDARY INFECTION       YES or No         BACTERIA/PROTOZOA SOLATED FROM HIMAN       SECONDARY INFECTION         INTECTIOUS AGENT       BIOCHEMICAL FEATURE, etc         SEROTYPE/SEROVAR       INTEL OCUS AGENT         SEROTYPE/SEROVAR       INTECTIOUS AGENT         SEROTYPE/SEROVAR       INTECTOUS AGENT         INTR	CATEGORY OF DISEAS	Ē	BACTERIA OR	VIRUS		
DIAGONESS DI DI DI CLUBAR DI NO. PARAGO PEROPERANKO COURSED PERODO FROM (YYMMOD) TO (YYMMOD) PERODO FROM (YYMMOD) TO (YYMMOD) SUSPECTED PLACE OF PEROPENKO FOOD SUSPECTED PLACE OF PEROPENKO FOOD SUSPECTED PLACE OF PEROPENKO FOOD SUSPECTED PLACE OF PEROPENKO FOOD NAME SUSPECTED PLACE OF PEROPENKO NAME SUSPECTED PLACE OF PEROPENKO NAME SUSPECTED CAUSE INCREMENT NAME SUSPECTED CAUSE INCREMENT SECONDARY INFECTION ANALE SUSPECTED CAUSE INCREMENT SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTIONS AGENT SECONDARY SECONDARY INFECTIONS AGENT SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTIONS AGENT SECONDARY INFECTIONS AGENT SECONDARY INFECTION AGENT SECONDARY INFECTION NON-AMELIER INFECTION INFERO INFECTION NON-AMELIER INFECTION NON-AMELIER INFECTION INFERO INFECTION NON-AMELIER INFECTION INFERO INFECTION NON-AMELIER INFECTION AGENT SECONDARY INFECTION NON-AMELIER INFECTION NON-AMELIER INFECTION NON-AMELIER INFECTION INFERO INFECTION INFORMARY INFECTION INFECTION INFERO INFECTION INFERO INFECTION INFERO INFECTION INFERO INFECTION INFERO INFECTION INFERO INFECTION INFERO INFECTION INFECTION INFECTION INFERO INFECTION INFEC						
Diversion         Diversion         Diversion           Diversion         Diversion         Diversion           SUSPEctor PLACE OF PERFONCEMENTAGE FOOD         TO (YYMMOD)           SUSPEctor PLACE OF PERFONCEMENTAGE FOOD         Diversion           SUSPEctor PLACE OF PERFONCEMENT REGION         Diversion           NAME         Diversion           SUSPEctor PLACE OF PERFONCEMENT REGION         Diversion           NAME         Diversion           SUSPEctor PLACE OF PERFONCEMENT REGION         Diversion           NUMBER OF CONSUMERS         NUMBER OF POSITIVE CASE           NUMBER OF EXAMINED         NUMBER OF POSITIVE CASE           SECONDARY INFECTION         SECONDARY INFECTION           SECONDARY INFECTION         SECONDARY INFECTION           SECONDARY INFECTION         SECONDARY INFECTION           SECONDARY INFECTION         SECONDARY           INFECTIONS AGENT         SECONDARY           SECONDARY INFECTION         SEC		INFECTION	ID No			
Second Seco						)
LINECTUD PLACE OF INDERANGE TOOD       TO (1199000)         SUSPECTED LACKS OF INFECTION/ACTING FOOD       MAME         SUSPECTED LACKS       NUMBER OF PATIENTS         SUSPECTED LACKS       NUMBER OF POSTIVE CASE         NUMBER OF CONSUMERS       NUMBER OF POSTIVE CASE         SUSPECTED LOCAUSE       NUMBER OF POSTIVE CASE         NUMBER OF EXAMINED       NUMBER OF POSTIVE CASE         SECONDARY INFECTION       SECONDARY INFECTION         SECONDARY INFECTION       SECONDARY INFECTION         SECONDARY INFECTION       SECONDARY INFECTION         SECONDARY INFECTION       SECONDARY INFECTION         SECONDARY INFECTION       SECONDARY         SECONDARY       SECONDARY         SECONDARY       SECONDARY         SECONDARY       SECONDARY         SECONDARY       SECONDARY         SECONDARY       SECONDARY         NEECTED/ARCAL FEATURE, etc       SECONDARY         SECONDARY       SECONDARY         SECONDARY       SECONDARY         SECONDARY </td <td></td> <td></td> <td></td> <td></td> <td>(COUNTRI/AREA</td> <td><u>)</u></td>					(COUNTRI/AREA	<u>)</u>
BUCKETED JALE OF NETERINSTOLUCE NORMARIZE OF CATEGORY AND FOOD ANAME SUSPECTED CAUSE MAME SUSPECTED CAUSE SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY INFECTION SECONDARY SEC						
AUDELDUSCED OF DECEMPONENT OF ATTENDED OF DECEMPONENT OF DECEMPONE						
INTEGRINATION DO CONSTRUCTION       INTEGRINATION OF CONSTRUCTION         SUSPECTED CAUSE       INTEGRINATION OF CONSTRUCTION         INTEGRICAL FEATURES       INTEGRICAL FEATURE, etc.         SECONDARY INFECTION       YES or No         BACTERIA/PROTOZOA ISOLATED FROM HUMAN       INTEGRICAL FEATURE, etc.         SEROTYPE/SEROVAR       BIOCHEMICAL FEATURE, etc.         SEROTYPE/SEROVAR       CONTARY						
SUSPECTED CAUSE           NUMBER OF CASUMERS         NUMBER OF PATIENTS           NAMER OF CAMMED         INDUSTRIAL PROM           NAMER OF CAMMED         INDUSTRIAL PROTOZOA ISOLATED FROM HUMAN           INFECTIONS AGENT         BIOCHEMICAL FEATURE, etc           SECONDARY INFECTION         BIOCHEMICAL FEATURE, etc           SECONDAR         CONDUCTECTON           NETECTORS AGENT         BIOCHEMICAL FEATURE, etc           SECONDAR         CONDUCTECTON           NETECTORS AGENT         BIOCHEMICAL FEATURE, etc           SECONDAR         CONDUCTECTON           MECTIONS AGENT         CONDUCTECTON           SENSTIVE CEL	INCRIMINATED FOOD	NAME			-	
SUBJECTION CODE       INUMBER OF PATIENTS         TO BERO F CONSUMERS       TO BERO F POSITIVE CASE         NUMBER OF CONSUMERS       NUMBER OF POSITIVE CASE         INDUCATION PERDO       Ins.         SECONDARY INFECTION       YES or No         BACTERIA/PROTOZOA ISOLATED FROM HUMAN       BIOCHEMICAL FEATURE, etc.         SEROTYFE/SEROVAR       CODEL         SEROTYFE/SEROVAR       CODEL         SEROTYFE/SEROVAR       CODEL         SEROTYFE/SEROVAR       CODEL         SECONTYFE/SEROVAR       CODEL         SECONTYFE/SEROVA						
NUMBER OF CONSUMERS INTO A CONSUMERS INTO A CONSUMERS OF PATIENTS TO TO THE PATHENTS AGE INCLUSATION PERIOD INTS. SECONDARY INFECTION YES OF NO.  ANTRECTORS AGENT SECONDARY INFECTION YES OF NO.  BIOCHEMICAL FEATURE, etc  SECONDARY INFECTION AGENT  SECONDARY INFECTION AGENT  BIOCHEMICAL FEATURE, etc  SECONDARY INFECTION AGENT  SECONDARY  SECONDARY SECONDARY  SECONDARY  SECONDARY SECONDARY  SECONDARY SECONDARY  SECONDARY  SECONDARY S	SUSPECTED CAUSE					
NUMBER OF CANSUMERS       NUMBER OF PARTNERS         NUMBER OF EXAMINED       NUMBER OF POSITIVE CASE         NUMBER OF EXAMINED       NUMBER OF POSITIVE CASE         BACTERIA/PROTOZOA ISOLATED FROM HUMAN       SECONDARY INFECTION         SECONDARY INFECTION       YES or No         BACTERIA/PROTOZOA ISOLATED FROM HUMAN       BIOCHEMICAL FEATURE, etc         SECONDARY TYPE/PATHOGENICITY       BIOCHEMICAL FEATURE, etc         SECONDARY TYPE/PATHOGENICITY       BIOCHEMICAL FEATURE, etc         SEROTYPE/SEROVAR       INFECTIOUS AGENT         MINECTIOUS AGENT       INFECTIOUS AGENT         MINECTIOUS AGENT       INFECTIOUS AGENT         SEROTYPE/SEROVAR       INFECTIOUS AGENT         INFECTIOUS AGENT       ICOLL IMMOUM         INFECTIOUS AGENT       ICOLATED/OPTECTON         INFECTIOUS AGENT       ICOLATED/OPTECTION         ICOLATED/DETECTION       ICOLATED/OPTECTION         ICOLATED/DETECTION						
A LIEUX JAGE A MUNER OF EXAMINED INUMER OF POSITIVE CASE INUMER OF ROM HUMAN INFECTION ACRAT SECONDARY INFECTION BACTERIA/PROTOZOA ISOLATED FROM HUMAN INFECTIONS AGENT BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER SECONDA BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SECONDA BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SECONDA BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SECONDA SECONDA VIRUS SOLATED/DETECTED FROM HUMAN INFECTOUS AGENT SECONTOR OF SOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SECONDA VIRUS SOLATED/DETECTED FROM HUMAN INFECTOUS AGENT SECONDA VIRUS SOLATED/DETECTED FROM HUMAN INFECTOUS AGENT METHOD OF ISOLATED FROM HUMAN INFECTOUS AGENT VIEUS SOLATED/DETECTED FROM HUMAN INFECTOUS AGENT METHOD OF ISOLATED FROM HUMAN INFECTOUS AGENT VIEUS SOLATED/DETECTED FROM HUMAN INFECTION AGENT VIEUS SOLATED/DETECTED FROM HUMAN INFECTION AGENT VIEUS SOLATED/DETECTED FROM HUMAN INFECTION AGENT VIEUS SOLATED/DETECTION INFECTION AGENT VIEUS SOLATED/	DATIENT'S ACE	TROM	TO		-	
NUMBER OF EXAMINED NUMBER OF POSITIVE CASE SECONDARY INFECTION VES or No ACTERIA/PROTOZOA ISOLATED FROM HUMAN INFECTIOUS AGENT SERVITYE/PATHOGENCITY BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER SECOMEN INFECTIOUS AGENT SERVITYE/PATHOGENCITY BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SERVITYE/SERVIAR SERVITYE/SERVIAR TOXIN TYPE/PATHOGENCITY URUS SOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN INFECTIOUS AGENT VIRUS AGENT VIRUS SOLATED FROM HUMAN POSITIVE INFECTIOUS AGENT VIRUS SOLATED FROM HUMAN POSITIVE VIRUS SOLATED/DETECTED FROM HUMAN POSITIVE P		FROM			-	
INCURATION PERGU       ITS.       SELUMUART INFECTION       TES OF NO         BACTERIA/PROTOZOA ISOLATED FROM HUMAN       INFECTIONS ACENT       BIOCHEMICAL FEATURE, etc         SEROTYPE/SEROVAR       INFECTIONS ACENT       BIOCHEMICAL FEATURE, etc         VIRUS ISOLATED/RETECTED FROM HUMAN       IPOSITIVE       INFECTIONS ACENT         VIRUS ISOLATED/RETECTED FROM HUMAN       IPOSITIVE       INFECTIONS ACENT         VIRUS ISOLATED/RETECTED FROM HUMAN       IPOSITIVE       INFECTIONS ACENT         INFECTIONS ACENT       IPOR       IPOR       IPOR         SENSITIVE CELLS       IPORAL       IPOR       IPOR         INFECTIONS ACENT       IPOR       IPOR       IPOR       IPOR         INFECTION ACECSCOPY       IMERGOD       IPHRIDIZATION       IPOR-SEQUENCING       IOTHER         INFECTION ACENT       IPOR       IPOR       IPOR-SEQUENCING       IOTHER <td></td> <td></td> <td>NUMBER OF POSITIVE CASE</td> <td>VEC 11</td> <td></td> <td></td>			NUMBER OF POSITIVE CASE	VEC 11		
ACTERIA/PROTOZOA ISOLATED FROM HUMAN INFECTOUS AGENT SEROTYPE/PATHOGENICITY  BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER SECOMEN INFECTOUS AGENT SECOMEN SECOMEN INFECTOUS AGENT SECOMEN INFECTON SEC	INCUBATION PERIOD	nrs.	SECONDARY INFECTION	YES or No		
BACTERIA/PROTOZOA ISOLATED FROM HUMAN INFECTIOUS AGENT SEROTYPE/SEROVAR TOXIN TYPE/PATHOGENICITY BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN INFECTIOUS AGENT BACTERIA/PROTOZOA ISOLATED FROM HUMAN POSITIVE BACTERIA/PROTOZOA ISOLATED FROM HUMAN BACTERIA/PROTOZOA ISOLATED FROM HUMAN BACTERIA/PROTOZOA ISOLATED BACT						
INFECTIOUS AGENT SERVIYE/SERVIAR TOXIN TYPE/PATHOGENICITY BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER SECOMEN INFECTIOUS AGENT SERVIYE/SERVIAR TOXIN TYPE/PATHOGENICITY BIOCHEMICAL FEATURE, etc SERVIYE/SERVIAR SECOMEN INFECTIOUS AGENT SERVIYE/SERVIAR SECOMEN INFECTIOUS AGENT UNECTIOUS AGENT UNECTION CLLL FEATURE, etc SERVIYE SERVIY	BACTERIA/PROTOZOA	ISOLATED FROM HUMAN				
SERUTYPE/SEROVAR         TOXIN TYPE/PATHOGENICITY         BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER         SPECIMEN         INFECTIOUS AGENT         SEROTYPE/SEROVAR         TOXIN TYPE/PATHOGENICITY         BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS         SPECIMEN         INFECTIOUS AGENT         BIOCHEMICAL FEATURE, etc         SEROTYPE/SEROVAR         TOXIN TYPE/PATHOGENICITY         INFECTIOUS AGENT         INFECTION         INFECTION         INFECTION         ICLL         INFECTION         INFELFICENT	INFECTIOUS AGENT			BIOCHEMICAL	FEATURE, etc	
TOXIN TYPE/PATHOGENICITY         BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER         SPECOMEN         INFECTIOUS AGENT         BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS         SPECOMEN         BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS         SPECOMEN         INFECTIOUS AGENT         BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS         SPECOMEN         INFECTIOUS AGENT         TOXIN TYPE/PATHOGENICITY         INFECTIOUS AGENT         INFECTION MICROSCOPY         INFECTION ICRO MULTION </td <td>SEROTYPE/SERO</td> <td>VAR</td> <td></td> <td></td> <td></td> <td></td>	SEROTYPE/SERO	VAR				
BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER SPECIMEN  BIOCHEMICAL FEATURE, etc BIOCHEMI	TOXIN TYPE/PAT	HOGENICITY				
BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER SECOMEN INFECTOUS AGENT SEROTYPE/SEROVAR TOXIN TYPE/PATHOGENICITY BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SECOMEN BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SECOMEN INFECTOUS AGENT SEROTYPE/SEROVAR BIOCHEMICAL FEATURE, etc SECOMEN INFECTOUS AGENT INFECTON NON-AMPLIFIED AMPUIFED AMPU						
BACTERIA/PROTOZOA ISOLATED FROM FOOD, DRINKING WATER SPECIMEN  SPECTAUS AGENT  SEROTYPE/SEROVAR  TOXIN TYPE/PATHOGENICITY  BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN  INFECTIOUS AGENT  SEROTYPE/SEROVAR  SEROTYPE/SEROVAR  SEROTYPE/SEROVAR  OUTHER SEROTYPE/SEROVAR  OUTHER SEROTYPE/SEROVAR  SEROTYPE/SEROTYPE/SEROVAR  SEROTYPE/SEROVAR  SEROTYP						
INFECTIOUS AGENT INFECTIOUS AGENT SECOMEN INFECTIOUS AGENT SECOMEN BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN INFECTIOUS AGENT BIOCHEMICAL FEATURE, etc INFECTIOUS AGENT VIRUS ISOLATED/DETECTED FROM HUMAN POSITIVE BIOCHEMICAL FEATURE, etc INFECTIOUS AGENT ICAL MEDIUM POSITIVE BIOCHEMICAL FEATURE, etc INFECTIOUS AGENT BIOCHEMICAL FEATURE, etc INFECTIOUS AGENT INFECTIOUS AGENT ICAL MEDIUM POSITIVE BIOCHEMICAL FEATURE, etc INFECTION SEGNITYPE/PATHOGENICITY BIOCHEMICAL FEATURE, etc INFECTION SEGNITYPE/PATHOGENICITY BIOCHEMICAL FEATURE, etc INFECTION SEGNITYPE/PATHOGENICITY BIOCHEMICAL FEATURE, etc INFECTION SEGNITYPE CELLS POSITIVE BIOCHEMICAL FEATURE, etc INFECTION SEGNITYPE BIOCHEMICAL FEATURE, etc INFECTION SEGNITYPE CELLS POSITIVE BIOCHEMICAL FEATURE, etc INFECTION POSITIVE BIOCHEMICAL FEATURE POSITIVE BIOCHEMICAL FEATURE, etc INFECTION POSITIVE BIOCHEMICAL FEATURE, etc INFECTION POSITIVE BIOCHEMICAL FEATURE BIOCHEMICAL FEATURE, etc INFECTION POSITIVE BIOCHEMICAL FEATURE, etc INFECTION POSITIVE BIOCHEMICAL FEATURE BIOCHEMI			2 WATER			
JFECTIOUS AGENT       BIOCHEMICAL FEATURE, etc         SERDTYPE/PATHOGENICITY       BIOCHEMICAL FEATURE, etc         BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS       BIOCHEMICAL FEATURE, etc         SPECIMEN       SEROTYPE/SEROVAR         TOXIN TYPE/PATHOGENICITY       BIOCHEMICAL FEATURE, etc         VIRUS ISOLATED/DETECTED FROM HUMAN       IPOSITIVE         INFECTIOUS AGENT       BIOCHEMICAL FEATURE, etc         VIRUS ISOLATED/DETECTED FROM HUMAN       IPOSITIVE         INFECTIOUS AGENT       INEGATIVE         METHOD OF ISOLATION/DETECTION       MEDIUM         CHELL       MEDIUM         CHELC TON NICHARDY CELLS       (PASSAGE)         SENSITIVE CELLS       (PASSAGE)         GENE DETECTION       FA         GENE DETECTION       FA         GENE DETECTION MICROSCOPY       INCROSCOPY         OTHER       OPCR         MIRECTION SAGENT       (PASSAGE)         VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER         SPECIMEN       INCENTIVE CELLS         (PASSAGE)       SENSITIVE CELL	SDECIMEN	ISOLATED FROM FOOD, DRINKING	JWAIER			
INPECTIOUS AGENT       BOULHEMICAL PEATURE, etc         SEROTYPE/SEROVAR						
SERVITYP2/SERVOVAK         TOXIN TYP2/PATHOGENICITY         BACTERIA/PROTOZOA ISOLATED FROM K/ITCHEN, COOKWARE, FOOD HANDLERS         SPECIMEN         INFECTIOUS AGENT         SEROTYP2/SEROVAR         TOXIN TYP2/PATHOGENICITY         VIRUS ISOLATED/DETECTED FROM HUMAN         OF ISOLATED/DETECTED FROM HUMAN         OCELL         METHOD OF ISOLATION/DETECTION         GENE DETECTION         JURUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER         SECOMEN         INFECTIOUS AGENT         INFECTIOUS AGENT         METHODO FISOLATION/DETECTION         GENE DETECTION NOLADRUPIED         ANTIGEN DETECTION NOLADRUPIED         GENE DETECTION NOLADRUPIED         GENENDETECTION NOLADRUPIED	INFECTIOUS AGENT	VAD		BIOCHEMICAL	FEATURE, etc	
IDXIN TYPE/PATHOGENICITY  BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN INFECTIOUS AGENT  SEROTYPE/PATHOGENICITY  URUS ISOLATED/DETECTED FROM HUMAN OPOSITIVE INFECTIOUS AGENT  CCELL IMEDIUM CHICK EMERYO (PASSAGE) ANIMAL OTHER SENSITIVE CELLS (PASSAGE) SENSITIVE CELLS (PASSAGE) ANUPLIFED OPCR PCR PCR+HYBRIDIZATION PAGE OTHER AMPLIFED OPCR PCR PCR+HYBRIDIZATION PAGE OTHER (PASSAGE) ANIMAL A	SERUT TPE/SERU	VAR				
BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN  SEROTYPE/SEROVAR TOXIN TYPE/PATHOGENICITY  VIRUS ISOLATED/DETECTED FROM HUMAN OFISOLATION/DETECTION CELL MEDIUM CHICK EMBRYO (PASSAGE ANTIGEN DETECTION FA ELA CPASHITIVE CELLS (PASSAGE ANTIGEN DETECTION AMPLIFIED AMPLIFIED CELL CPASHITIVE CELLS (PASSAGE COTHER SPECIMEN CELL CPASHITIVE CELLS (PASSAGE COTHER CPCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER SPECIMEN CELL CPASHITIVE CELLS (PASSAGE COTHER CPCR PCR+HYBRIDIZATION PCR+SEQUENCING COTHER CPCR PCR+HYBRIDIZATION PCR+SEQUENCING COTHER CENC AMPLIFIED CELL CPASHITIVE CELLS (PASSAGE COTHER CPASSAGE COTHER CPCR PCR+HYBRIDIZATION PCR+SEQUENCING COTHER CPCR PCR+HYBRIDIZATION PCR+SEQUENCING COTHER CPCR AMPLIFIED ANTIGEN DETECTION CELL CPCR AMPLIFIED PCR PCR+HYBRIDIZATION PCR+SEQUENCING COTHER CPCR AMPLIFIED ANTIGEN DETECTION CELLS CPASSAGE COTHER CPCR AMPLIFIED CPCR CPCR+HYBRIDIZATION COTHER CPCR AMPLIFIED CPCR CPCR+HYBRIDIZATION COTHER CPCR AMPLIFIED CPCR AMPLIFIED CPCR COTHER CPCR COTHER CPCR COTHER CPCR COTHER CPCR COTHER CPCR COTHER CPCR CPCR CPCR+SEQUENCING COTHER CPCR CPCR CPCR CPCR CPCR CPCR CPCR CP	TOXIN TYPE/PAT	HOGENICITY				
BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN INFECTIOUS AGENT VIRUS ISOLATED/DETECTED FROM HUMAN OF ISOLATION/DETECTION CCELL SENSITIVE CELLS (PASSAGE ) CANIMAL OTHER SENSITIVE CELLS (PASSAGE ) CANIMAL OTHER ( COTHER AMPLIFED PCR PCR PCR+HYBRIDIZATION PCC+SEQUENCING OTHER SENSITIVE CELLS (PASSAGE ) CANIMAL OTHER SENSITIVE CELLS (PASSAGE ) CANIMAL OTHER ( COTHER C						
BACTERIA/PROTOZOA ISOLATED FROM KITCHEN, COOKWARE, FOOD HANDLERS SPECIMEN INFECTIOUS AGENT SEROTYPE/SEROVAR TOXIN TYPE/PATHOGENICITY  VIRUS ISOLATED/DETECTEO FROM HUMAN OF ISOLATION/DETECTION CELL MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ) CONTHER AMPLIFED OF ISOLATION/DETECTION CELL MEDIUM CHICK EMBRYO (PASSAGE ) CONTHER AMPLIFED OF ISOLATION/DETECTION CELL MEDIUM CHICK EMBRYO (PASSAGE ) CONTHER SPECIMEN CELL MEDIUM CHICK EMBRYO (PASSAGE ) CONTHER CONTHER CELL CRON MICROSCOPY OF ISOLATION/DETECTION CELL MEDIUM CHICK EMBRYO (PASSAGE ) CONTHER SPECIMEN CELL MEDIUM CHICK EMBRYO (PASSAGE ) CONTHER CELL CRON MICROSCOPY OF ISOLATION/DETECTION CELL MEDIUM CHICK EMBRYO (PASSAGE ) CONTHER SPECIMEN CELL CRON MICROSCOPY CELL MEDIUM CHICK EMBRYO (PASSAGE ) CONTHER CELL CRON MICROSCOPY CELL CRON MIC						
SPECIMEN INFECTIOUS AGENT SEROTYPE/SEROVAR TOXIN TYPE/PATHOGENICITY  VIRUS ISOLATED/DETECTED FROM HUMAN OPOSITIVE INFECTIOUS AGENT SENSITIVE CELLS (PASSAGE) ANIMAL OTHER SENSITIVE CELLS (PASSAGE) ANIMAL OTHER ( AMPLIFED OPCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER SENSITIVE CELLS (PASSAGE) (PASSA	BACTERIA/PROTOZOA	ISOLATED FROM KITCHEN, COOK	WARE, FOOD HANDLERS			
INFECTIOUS AGENT  SEROTYPE/SEROVAR  TOXIN TYPE/PATHOGENCITY  VIRUS ISOLATED/DETECTED FROM HUMAN  VIRUS ISOLATED/DETECTION  CELL  METHOD OF ISOLATION/DETECTION  CELLS  CANTIGEN DETECTION  AMPLIFIED  AMPLIFIED  VIRUS ISOLATED/DETECTEO FROM FOOD, DRINKING WATER  SPECIMEN  VIRUS ISOLATED/DETECTION  CELLS  (PASSAGE )  ANIMAL OTHER  VIRUS ISOLATED/DETECTION  CELLS  (PASSAGE )  ANIMAL OTHER  (PASSAGE )  (PAS	SPECIMEN					
SEROTYPE/SEROVAR         TOXIN TYPE/PATHOGENICITY         VIRUS ISOLATED/DETECTED FROM HUMAN       POSITIVE         INFECTIOUS AGENT         CELL       MEDIUM         CELL       MEDIUM         CELL       (PASSAGE)         SENSITIVE CELLS       (PASSAGE)         GENE DETECTION       CANTIGEN DETECTION         GENE DETECTION NON-AMPLIFIED       HYBRIDIZATION         HYBRIDIZATION       PAGE         OTHER       AMPLIFIED         AMPLIFIED       PCR         INFECTIOUS AGENT         METHOD OF ISOLATION/DETECTION         CHECKTRON MICROSCOPY         OTHER (         NINFECTIOUS AGENT         METHOD OF ISOLATION/DETECTION         CELL       MEDIUM         CHICK EMBRYO (PASSAGE )         METHOD OF ISOLATION/DETECTION         INFECTIOUS AGENT         METHOD OF ISOLATION/DETECTION         CELL       MEDIUM         CHICK EMBRYO (PASSAGE )       ANIMAL         OTHER       SENSITIVE CELLS         SENSITIVE CELLS       (PASSAGE )         SENSITIVE CELLS       (PASSAGE )         CHECTION NON-AMPLIFIED       HYBRIDIZATION         METHOD OF ISOLATION       FA	INFECTIOUS AGENT			BIOCHEMICAL	FEATURE, etc	
TOXIN TYPE/PATHOGENICITY         VIRUS ISOLATED/DETECTED FROM HUMAN       POSITIVE         INFECTIOUS AGENT         METHOD OF ISOLATION/DETECTION         INFLORD OF ISOLATION ON-AMPLIFIED         INFLORD OF ISOLATION/DETECTION         INFECTIOUS AGENT         INFECTIOUS AGENT         INFECTIOUS AGENT         INFECTIOUS AGENT         INFECTION/DETECTION         INFECTIOUS AGENT         INFECTIOUS AGENT         INFECTION/DETECTION         INFECTIOUS AGENT         INFECTIOUS AGENT         INFECTION/DETECTION         INFECTIOUS AGENT         INFECTION/DETECTION	SEROTYPE/SERO	VAR				
VIRUS ISOLATED/DETECTED FROM HUMAN POSITIVE INFECTIOUS AGENT METHOD OF ISOLATION/DETECTION CCELL MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ANTIGEN DETECTION NOR-AMPLIFED HYBRIDIZATION PAGE OTHER AMPLIFIED PCR PCR +HYBRIDIZATION PCR+SEQUENCING OTHER ELECTRON MICROSCOPY MICROSCOPY OTHER ( ) ) VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT METHOD OF ISOLATION/DETECTION CCELL MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ANTIGEN DETECTION NOR-AMPLIFIED HYBRIDIZATION PCR+SEQUENCING OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ANTIGEN DETECTION NOR-AMPLIFIED HYBRIDIZATION PAGE OTHER CELL MEDIUM CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ANTIGEN DETECTION NOR-AMPLIFIED HYBRIDIZATION PAGE OTHER CELL MEDIUM CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE AMPLIFIED OF PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER CELLCTRON NOR-AMPLIFIED HYBRIDIZATION PCR+SEQUENCING OTHER AMPLIFIED OFCR PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER AMPLIFIED PCR PCR PCR+HYBRIDIZATION PCR+SEQUENCING PC	TOXIN TYPE/PAT	HOGENICITY				
VIRUS ISOLATED/DETECTED FROM HUMAN       POSITIVE       INEGATIVE         INFECTIOUS AGENT       INEGATIVE       INEGATIVE         METHOD OF ISOLATION/DETECTION       IC       INEGATIVE         SENSITIVE CELLS       (PASSAGE)       SENSITIVE CELLS       (PASSAGE)         INFECTION       IFA       IEIA       IRPHA       ILA       IC       OTHER         GENE DETECTION       IFA       IEIA       IPARA       IC       OTHER         GENE DETECTION       NON-AMPLIFIED       IPVBRIDIZATION       IPAGE       OTHER         MICROSCOPY       MICROSCOPY       MICROSCOPY       OTHER       OTHER         VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER       SPECIMEN       INFECTIOUS AGENT       INFECTIOUS AGENT         METHOD OF ISOLATION/DETECTION       ICCELLS       (PASSAGE)       SENSITIVE CELLS       (PASSAGE)         OTHER       SENSITIVE CELLS       (PASSAGE)       SENSITIVE CELLS       (PASSAGE)         INFECTIOUS AGENT       IFTHOD OF ISOLATION/DETECTION       ICCELL       ICCELLS       (PASSAGE)       ICCINC (PASSAGE)       IANIMAL       OTHER         SENSITIVE CELLS       (PASSAGE)       SENSITIVE CELLS       (PASSAGE)       IANIMAL       OTHER         GENE DETECTION       IFA       I						
VIRUS ISOLATED/DETECTED FROM HUMAN POSITIVE NEGATIVE NEGATIVE NEGATIVE NEGATIVE CELLS POSITIVE CELLS PASAGE ANTIGEN DETECTION FA EIA PA CONTRACTOR OF ISOLATION/DETECTION FA EIA PA CONTRACTOR OF ISOLATION NON-AMPLIFIED HYBRIDIZATION PAGE OTHER AMPLIFIED PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER OTHER ( )  VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ) CONTRACTOR OF ISOLATION/DETECTION CONTRACTOR OF ISOLATION CONTRACTOR OF ISOLATION/DETECTION CONTRACTOR OF ISOLATION/DETECTION CONTRACTOR OF ISOLATION/DETECTION CONTRACTOR OF ISOLATION CONTRACTOR OF ISOLATION CONTRACTOR OF ISOLATION OF ISO						
VIRUS ISOLATED/DETECTED FROM HUMAN POSITIVE INFECTIOUS AGENT METHOD OF ISOLATION/DETECTION FRA CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ) ANTIGEN DETECTION ON-AMPLIFIED CHYBRIDIZATION CAR COTHER AMPLIFIED PCR PCR + HYBRIDIZATION PCR+SEQUENCING OTHER VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT METHOD OF ISOLATION/DETECTION FAA CHICK EMBRYO (PASSAGE ) ANIMAL OTHER CELL CRON MICROSCOPY MICROSCOPY OTHER ( ) MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SPECIMEN INFECTIOUS AGENT METHOD OF ISOLATION/DETECTION FAA CHICK EMBRYO (PASSAGE ) ANIMAL OTHER CELL CRON MICROSCOPY CHICK EMBRYO (PASSAGE ) CANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER AMPLIFIED CHICK EMBRYO (PASSAGE ) SENSITIVE CELLS (PASSAGE ) COTHER (PASSAGE ) COTHER (PASSAGE ) COTHER (PASSAGE ) COTHER (						
INFECTIOUS AGENT  METHOD OF ISOLATION/DETECTION  CELL SENSITIVE CELLS (PASSAGE ANTIGEN DETECTION  CHICK EMBRYO (PASSAGE ) ANIMAL OTHER  CENCE DETECTION  FA ELA PCR	VIRUS ISOLATED/DETE	CTED FROM HUMAN				
INTEGRIGUES ACTION / DETECTION METHOD OF ISOLATION/DETECTION CCELL MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) CONTHER ANTIGEN DETECTION ON-AMPLIFIED HYBRIDIZATION PAGE OTHER AMPLIFIED PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER ELECTRON MICROSCOPY MICROSCOPY OTHER ( ) VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT METHOD OF ISOLATION/DETECTION CELL MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) ANIMAL OTHER GENE DETECTION OF A EIA RPHA LA PA IC OTHER AMPLIFIED PCR PCR+HYBRIDIZATION PAGE OTHER ANTIGEN DETECTION OF A EIA RPHA CLA PA IC OTHER AMPLIFIED PCR PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER AMPLIFIED PCR PCR PCR PCR PCR PCR PCR SEQUENCING OTHER AMPLIFIED PCR PCR PCR PCR PCR SEQUENCING OTHER AMPLIFIED PCR						
MELTIOD OF ISOLATION SENSITIVE CELLS (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ANTIGEN DETECTION OFA EIA ORPHA LA PA CONTRER GENE DETECTION NON-AMPLIFIED OPCR OPCR-HYBRIDIZATION OPCR-SEQUENCING OTHER AMPLIFIED OPCR OPCR-HYBRIDIZATION OPCR-SEQUENCING OTHER ELECTRON MICROSCOPY MICROSCOPY OTHER ( ) VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT CELL OMEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) CANIMAL OTHER GENE DETECTION OF FA EIA ORPHA LA PA CIC OTHER AMPLIFIED OPCR OPCR+HYBRIDIZATION OPCR+SEQUENCING OTHER REMARKS				_		
SENSITIVE CELLS       (PASSAGE)       SENSITIVE CELLS       (PASSAGE)         ANTIGEN DETECTION       FA       EIA       RPHA       LA       PA       IC       OTHER         GENE DETECTION       NON-AMPLIFIED       HYBRIDIZATION       PAGE       OTHER       AMPLIFIED       OTHER         AMPLIFIED       PCR       PCR       PCR+HYBRIDIZATION       PCR+SEQUENCING       OTHER         ELECTRON MICROSCOPY       MICROSCOPY       OMICROSCOPY       OTHER       OTHER       OTHER         VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER       SPECIMEN       INFECTIOUS AGENT       OTHER       OTHER         METHOD OF ISOLATION/DETECTION       CELL       MEDIUM       CHICK EMBRYO (PASSAGE)       ANIMAL       OTHER         GENE DETECTION       IMEDIUM       CHICK EMBRYO (PASSAGE)       SENSITIVE CELLS       (PASSAGE)         METHOD OF ISOLATION/DETECTION       IMEDIUM       CHICK EMBRYO (PASSAGE)       SENSITIVE CELLS       (PASSAGE)         GENE DETECTION       IMEDIUM       CHICK EMBRYO (PASSAGE)       SENSITIVE CELLS       (PASSAGE)         GENE DETECTION       IMEDIUM       CHICK EMBRYO (PASSAGE)       SENSITIVE CELLS       (PASSAGE)         GENE DETECTION       IMEDIUM       CHICK EMBRYO (PASSAGE)       SENSITIVE CELLS	METHOD OF ISOLATION			)		
ANTIGEN DETECTION OFA CLLLS (PASSAGE ) SENSITIVE CLLLS (PASSAGE ) SENSITIVE CLLLS (PASSAGE ) SENSITIVE CLLLS (PASSAGE ) COTHER AMPLIFIED PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER COTHER ( ) (PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER ) (PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER ) (PASSAGE ) COTHER ( ) (PCR PCR+HYBRIDIZATION PCR+SEQUENCING PCR+ENDER ) (PASSAGE ) (PCR PCR+HYBRIDIZATION PCR+SEQUENCING PCR PCR+HYBRIDIZATION PCR+SEQUENCING PCHER PCR PCR PCR PCR PCR PCR PCR PCR PCR PC						(DASSACE
ANTIGEN DETECTION ON-AMPLIFIED HYBRIDIZATION OPAGE OTHER GENE DETECTION NON-AMPLIFIED OPCR OPCR+HYBRIDIZATION OPAGE OTHER ELECTRON MICROSCOPY MICROSCOPY OTHER ( ) VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT METHOD OF ISOLATION/DETECTION CELLS (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ANTIGEN DETECTION OFA OPCR OPCR OTHER GENE DETECTION NON-AMPLIFIED OPCR OPCR OPCR OTHER AMPLIFIED OPCR OPCR OPCR OTHER AMPLIFIED OPCR OPCR OPCR OTHER AMPLIFIED OPCR OPCR OPCR OPCR OPCR OPCR OPCR OPCR						
Image: Internation of the image: I						
AMPLIFIED       IPCR       IPCR       IPCR+HYBRIDIZATION       IPCR+SEQUENCING       IOTHER         Implified       Internation       Implified	GENE DETECTI	JN NON-AMPLIFIED				
ELECTRON MICROSCOPY       IMICROSCOPY         OTHER (       )         VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER         SPECIMEN         INFECTIOUS AGENT         INFECTIOUS AGENT         ICELL       IMEDIUM         CHICK EMBRYO (PASSAGE)       IANIMAL         OT HER       SENSITIVE CELLS         INFECTION       IFA         ELECTRON NOR-AMPLIFIED       HYBRIDIZATION         IELECTRON MICROSCOPY       MICROSCOPY         OTHER (       )		AMPLIFIED	□PCR □PCR+HYBRI	DIZATION	□PCR+SEQUENCING	
OTHER (       )         VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER         SPECIMEN         INFECTIOUS AGENT         METHOD OF ISOLATION/DETECTION         CELL       MEDIUM         CELLS       (PASSAGE)         ANTIGEN DETECTION       FA         GENE DETECTION       FA         GENE DETECTION       FA         METHOD OF ISOLATION/DETECTION       PAGE         SENSITIVE CELLS       (PASSAGE)         SENSITIVE CELLS       (PASSAGE)         GENE DETECTION       FA         AMPLIFIED       HYBRIDIZATION         AMPLIFIED       PCR         BELECTRON MICROSCOPY       MICROSCOPY         OTHER (       )	ELECTRON MIC	ROSCOPY DICROSCOPY				
VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT CELL MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ) ANTIGEN DETECTION FA EIA RPHA LA PA IC OTHER GENE DETECTION NON-AMPLIFIED HYBRIDIZATION PAGE OTHER AMPLIFIED PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER ELECTRON MICROSCOPY MICROSCOPY OTHER ( )	□OTHER (	)				
VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT CELL IMEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAG ANTIGEN DETECTION FA EIA RPHA LA PA IC OTHER GENE DETECTION NON-AMPLIFIED HYBRIDIZATION PAGE OTHER AMPLIFIED PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER ELECTRON MICROSCOPY MICROSCOPY OTHER ( )						
VIRUS ISOLATED/DETECTED FROM FOOD, DRINKING WATER SPECIMEN INFECTIOUS AGENT METHOD OF ISOLATION/DETECTION CELL  MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ) CONTHER CONT						
SPECIMEN         INFECTIOUS AGENT         METHOD OF ISOLATION/DETECTION         CELL       MEDIUM         CHICK EMBRYO (PASSAGE)       CANIMAL         SENSITIVE CELLS       (PASSAGE)         ANTIGEN DETECTION       FA         GENE DETECTION       NON-AMPLIFIED         AMPLIFIED       HYBRIDIZATION         PCR       PCR+HYBRIDIZATION         PCR       PCR         PCR       PCR+HYBRIDIZATION         PCR       PCR         PCR	VIRUS ISOLATED/DETE	CTED FROM FOOD. DRINKING WA	TER			
INFECTIOUS AGENT  METHOD OF ISOLATION/DETECTION  CCELL MEDIUM CHICK EMBRYO (PASSAGE ) ANIMAL OTHER  SENSITIVE CELLS (PASSAGE ) SENSITIVE CELLS (PASSAGE ANTIGEN DETECTION FA EIA RPHA LA PA IC OTHER  GENE DETECTION NON-AMPLIFIED HYBRIDIZATION PAGE OTHER  AMPLIFIED PCR PCR+HYBRIDIZATION PCR+SEQUENCING OTHER  ELECTRON MICROSCOPY OTHER ( )  REMARKS	SPECIMEN	,				
METHOD OF ISOLATION/DETECTION	INFECTIOUS AGENT			_		
INETTIOD OF ISOLATION THE INTERTION OF A SOLATION OF A SOLATION OF A SENSITIVE CELLS (PASSAGE) SENSITIVE SENSITY S				-		
Sensitive cells     (PASSAGE)     Sensitive cells     (PASSAGE)       ANTIGEN DETECTION     FA     EIA     RPHA     LA     PA     IC     OTHER       GENE DETECTION     NON-AMPLIFIED     HYBRIDIZATION     PAGE     OTHER     OTHER       AMPLIFIED     PCR     PCR+HYBRIDIZATION     PCR+SEQUENCING     OTHER       ELECTRON MICROSCOPY     MICROSCOPY     OTHER     OTHER       OTHER     )     REMARKS     REMARKS	METHOD OF ISOLATION		CHICK EMBRYO (PASSAGE	)		
ANTIGEN DETECTION     FA     EIA     RPHA     LA     PA     IC     OTHER       GENE DETECTION     NON-AMPLIFIED     HYBRIDIZATION     PAGE     OTHER       AMPLIFIED     PCR     PCR+HYBRIDIZATION     PCR+SEQUENCING     OTHER       ELECTRON MICROSCOPY     MICROSCOPY     OTHER     OTHER       OTHER     )     REMARKS     REMARKS		SENSITIVE CELLS	DACCACE )			(PASSACE
Image: Section of the section of t						
AMPLIFIED OF CR OF CRAFT ON OF A MPLIFIED OF CRAFT ON OF CRAFT.						
AMPLIFIED UPCK UPCK+HYBRIDIZATION UPCR+SEQUENCING OTHER ELECTRON MICROSCOPY OTHER ()						
□ELECTRON MICROSCOPY □OTHER ( ) REMARKS			⊔PCR+HYBRI	DIZATION	□PCK+SEQUENCING	
DOTHER ( )	LIELECTRON MIC	RUSCUPY DMICROSCOPY				
REMARKS	□OTHER (	)				
REMARKS						
REMARKS						
	REMARKS					
RECORD No.					RECORD No	

\_\_\_\_\_

Fig. 5. Reporting form C for outbreak.

REPORTED BY	DATE OF REPORTING(YYMMDD)
BACTERIA OR VIRUS	
ID No.	DATE OF SPECIMEN COLLECTION (YYMMDD)
SPECIMEN	
DOMESTIC or IMPORETED	
NUMBER OF SPECIMEN	NUMBER OF POSITIVE SPECIMEN
	BIOCHEMICAL FEATURE, etc
	EMBRYO (PASSAGE) 🗆 🗆 ANIMAL 🗆 OTHER
	(PASSAGE) SENSITIVE CELLS (PASSAGE)
AMPLIFIED PCR	$\square PCR+HYBRIDIZATION$ $\square PCR+SEQUENCING$ $\square OTHER$
$\Box$ OTHER ( )	
REMARKS	
	RECORD No.

Fig. 6. Reporting form D for pathogen isolation/detection from food, environment and animal.

this supplement of *Japanese Journal of Infectious Diseases* (formerly *Japanese Journal of Medical Science and Biology*). This is the eighth annual report of infectious agent surveillance based upon the data collected through the on-line system.

#### 2. Notes on the use of the infectious agent information

The information given in this report is a summary of the infectious agents isolated/detected by etiological diagnosis or pathogen surveillance conducted for a public health purpose.

In analyzing and citing the information of the infectious agents published in this report, the following should be considered.

The report includes exclusively the positive results.

The information system collects exclusively the positive results of pathogen isolation/detection, without recording the number of specimens, having failed to yield any pathogen. The number of positive results may or may not reflect that of all the examinations. This must be taken into consideration, particularly when comparison is made by district.

The results may not necessarily be correlated to the diseases or clinical symptoms.

As a common problem in laboratory diagnosis, the infectious agent can not always be identified as the direct cause of the disease or the clinical symptoms.

When a pathogen is isolated/detected from specimens of lesions, such as cerebrospinal fluid, blood, vesicles, biopsy, and autopsy, it must be correlated to the disease. However, when stool, pharyngeal swabs, or urine is examined, a pathogen is isolated from a case of latent or inapparent infection or the one transiently present might be isolated or detected\*. Therefore, the correlation between the isolated/detected pathogen and the illness or clinical symptoms must be considered individually after checking up with epidemic conditions, specimens, examination methods and the results of other examinations

\*Since 1997, the results of detection by PCR can be reported. However, the above problems in PCR have not been solved, the viruses detected only by PCR are separately shown on the tables (see p. 121-133).

Isolation/detection from the same person may sometimes be reported from more than two sources.

Of the reports of examinations provided by PHI or quarantine station, particularly those concerned with legally designated notifiable infectious diseases (cholera and shigellosis, etc.), those of the pathogenic agent isolated/detected from the same case may be duplicated. There is no process for elimination of such duplication. It is improper, therefore, to total the reported numbers from PHIs and those from quarantine stations, if there is such duplication.

The numbers of isolated/detected pathogens reported in previous Annual Reports have been updated.

The previous data files, if there are new reports or corrections, are updated on a case basis at IDSC, NIID. The annual data for 1999-2003 on p. 54-55, 91-92, 100-101, 118-119 and 132-133 in this report may not necessarily coincide with the figures in the previously published annual reports. Such disagreement is rare in the reports of bacterial pathogens, whereas may be seen almost every year in viral isolation/ detection. Nevertheless, the difference is not so large as to influence the overall trend.

The numbers of reports of isolation/detection in this report were based on the laboratory data submitted to IDSC, NIID before October 2, 2007.

#### 3. The National Epidemiological Surveillance of Infectious Diseases in compliance with the enforcement of the Infectious Diseases Control Law

The NESID in Japan started in 1981 consists of 1) sentinel surveillance for occurrence of patients of 27 kinds of infectious diseases other than legally notifiable diseases, and 2) infectious agents surveillance. The surveillance, however, has not been based upon any legal basis.

In April 1999, the Communicable Disease Prevention Law in effect since 1897, the Venereal Disease Prevention Law since 1948, and the AIDS Prevention Law since 1989 have been abrogated and the Infectious Diseases Control Law is being enacted. In this new law, the NESID program is defined as one of the main objects. Intensifying the surveillance system based on notification from physicians, collection, comprehension and analysis of the incidence and the trend of infectious diseases, and feedback of such information are proposed; moreover active surveillance has been introduced for epidemiological investigations. Since the information of infectious agents is essential for providing adequate medical care to patients and important to prevent and control the spread of infectious diseases, it is necessary to collect, analyze, and publish the information on infectious agents. It is required that these information published benefit the general public as well as those working in medical fields.

The purpose of the new NESID program is settled in compliance with the Infectious Diseases Control Law to promote effective and accurate infectious disease control measures by reinforcing and expanding the conventional surveillance system, restructuring the computer-network system to obtain, analyze, and quickly publish the information on a nation-wide scale and conducting active surveillance. Government and local governments (prefectures and cities including special wards having health centers) are responsible for conducting the surveillance.

<u>Target diseases</u>: In the Infectious Diseases Control Law, all infectious diseases of Category I-V are designated as the targets of the NESID (see p. 19 Table 1).

Organization of the surveillance system: As the organization to play a central role in the national surveillance, the national infectious disease surveillance center has been organized. IDSC, NIID fulfills the function. A district infectious disease surveillance center has been organized by each local government, and it is placed mainly in the PHI to conduct the surveillance within the district. In each prefecture, one of district infectious disease surveillance centers is assigned for the key district infectious disease surveillance center, which collects and analyzes the information from the whole area of the prefecture and forwards the results to the rest of district infectious disease surveillance centers. With participation of experts on infectious diseases in different fields, a national committee of infectious disease surveillance is organized in MHLW and a district committee of infectious disease surveillance in the local government.

Surveillance for Category I-IV infectious diseases:

**Physicians diagnosing target infectious diseases:** The physician who has diagnosed any of the target infectious diseases must report immediately the name, age and sex of the patients and other information on the reporting form to the nearby HC. When HC requires tests for the etiological agent, the physician will send the available specimens and/or the information on the infectious agent to the PHI.

**HCs**: HC must immediately forward the patient information to the health department of the local government (local health department) and the district infectious disease surveillance center via computer-network system. When necessary, the HC will ask the physician to send the specimens and/or information on the etiology of the infection to PHI.

HC must also distribute the information on the incidence of target diseases and their infectious agents obtained from the district infectious disease surveillance center to the municipalities, the medical institutions concerned, the Medical Association, the Board of Education, etc. through weekly and monthly reports or other media.

HC receiving any notification on Category I-IV infectious diseases must inform the incident (except for the information on the patient's privacy) to the above-described organizations.

**PHIs:** PHI conducts the laboratory tests requested and sends the results to the physician through HC. The information on the infectious agents sent by the physician and the results of the laboratory tests must be sent to HC, the local health department, and the district infectious disease surveillance center.

Any tests difficult to conduct at PHI are transferred to the NIID.

**NIID**: The NIID conducts the laboratory test requested and reports the results to the PHI and the national infectious disease surveillance center.

Local health departments: Upon receiving the patient information from HC by electronic telecommunication, the local health department must send the information to the national infectious disease surveillance center through the computer-network system. The information on the infectious agents including the results of the tests sent from the PHI should also be reported immediately to the national infectious disease surveillance center.

**District (key district) infectious disease surveillance centers**: The district infectious disease surveillance center should collect and analyze the information on incidence of target diseases and their infectious agents (including that on the results of the tests conducted by PHI) and convey the information to HCs and other institutions concerned through weekly report or other media together with the published information on the whole country obtained from the national infectious disease surveillance center. The key district infectious disease surveillance center must furnish the information in the prefecture to the district infectious disease surveillance centers and other organizations together with the information on the whole country.

**National infectious disease surveillance center**: The national infectious disease surveillance center must immediately compile, analyze and evaluate the patient information received from the local health departments and send the information on incidence of target diseases in the whole country to the local health departments through weekly report or other media together with that of Category V infectious diseases. The information on infectious agents are analyzed and evaluated, and the results are to be sent immediately to

the local health departments and, if necessary, published in weekly report or other media.

Surveillance for Category V infectious diseases (required notifying all the cases):

**Physicians diagnosing target infectious diseases**: The physician who has diagnosed any of the target infectious diseases must report within 7 days the age and sex of the patient and other information on the reporting form to the nearby HC. Infectious agent surveillance will be conducted by request from the health center in the same way as for the infectious diseases in Category I-IV.

**HCs**: HC must forward as soon as possible, at the latest within 7 days the information to the local health department and the district infectious disease surveillance center through the computer-network system. Concerning the target diseases for infectious agent surveillance among Category V infectious diseases marked with \* in the Table 1 on p. 19, the health center will ask the physician to send specimens for the microbiological tests and/or the information on the infectious agent to PHI, if necessary.

HC must also regularly distribute in the same way as for the Category I-IV infectious diseases information on incidence of target diseases and their infectious agents retrieved from the district infectious disease surveillance center.

PHIs: Similar to the Category I-IV infectious diseases.

NIID: Similar to the Category I-IV infectious diseases.

**Local health departments**: Within 7 days after HC has received information, the local health department must send it to the national infectious disease surveillance center through the computer-network system. The information on the infectious agents sent from PHI should also be transmitted immediately to the national infectious disease surveillance center.

**District (key district) infectious disease surveillance centers**: Similar to the Category I-IV infectious diseases.

**National infectious disease surveillance center**: The national infectious disease surveillance center must immediately compile, analyze and evaluate the patient information sent from the local health departments and send the information on incidence of target diseases in the whole country to the local health departments through weekly reports or other media together with that on Category I-IV infectious diseases and Category V infectious diseases to be reported by the sentinel clinics and hospitals. The information on infectious agents is treated in the same way as for that on the Category I-IV infectious diseases.

Surveillance for Category V infectious diseases (required to be reported by the sentinel clinics and hospitals):

**Sentinels**: The number of sentinel clinics and hospitals is decided depending on the relative population of the jurisdiction of each HC and on consideration enabling comprehending the incidents in the whole area of the prefecture. The sentinel clinics comprise those for pediatric diseases (about 3,000 pediatrics in the whole country), those for influenza (3,000 pediatrics as mentioned above plus about 2,000 internal medicine), those for eye diseases (about 600 ophthalmology in the whole country), those for sexually transmitted diseases (about 900 STD clinics including gynecology, obstetrics, urology, and dermatology in the whole country). The sentinel hospitals primarily target inpatients (about 500 hospitals having more than 300 beds providing medical care in pediatrics and internal medicine in the whole country).

About 10% of sentinel clinics for pediatric diseases, influ-

enza, eye diseases and all the sentinel hospitals serve as sentinels for infectious agent surveillance. The target diseases for infectious agent surveillance are shown with a mark \* in the Table 1 on p. 19.

Incidence of the pediatric diseases, influenza and eye diseases found at sentinels are reported every week to health centers. Those of the target diseases at sentinel hospitals except for drug-resistant bacterial infections are also reported every week. Those of STD at sentinel clinics and drugresistant bacterial infections at sentinel hospitals are reported every month to health centers. Specimens for etiological tests are sent from sentinels for infectious agent surveillance to PHI.

**HCs**: HC must send the information on incidence of infectious diseases obtained from sentinels to the local health department and the district infectious disease surveillance center via computer-network system (weekly reports are sent by Tuesday of the next week and monthly reports by the 3rd of the next month).

HC must also feed back the information on incidence of target diseases and their infectious agents obtained from the district infectious disease surveillance center in the same way as for Category I-IV infectious diseases.

**PHIs**: Similar to the Category I-IV infectious diseases. **NIID**: Similar to the Category I-IV infectious diseases.

Local health departments: As soon as the information is received from HC, the local health department must forward it to the national infectious disease surveillance center through the computer-network system. The information on the infectious agents received from PHI should also be reported immediately to the national infectious disease surveillance center.

#### **District (key district) infectious disease surveillance centers**: Similar to the Category I-IV infectious diseases.

**National infectious disease surveillance center**: The national infectious disease surveillance center must immediately compile, analyze and evaluate the patient information received from the local health departments and send the information on incidence of target diseases in the whole country to the local health departments by weekly report or other media together with that on the Category I-IV infectious diseases and the Category V infectious diseases required to comprehend all the cases. The information on infectious agents is treated in the same way as that for the Category I-IV infectious diseases.

Active surveillance: Active surveillance for epidemiological investigation is introduced so that the governor of the local government can operate when any of the Category I-IV infectious diseases occurs or the incidence of Category V infectious diseases show an unusually different trend. Understanding and cooperation with close connection of people concerned may be necessary for operation of active surveillance. The Field Epidemiology Training Program to educate experts participating in active surveillance is being held at the NIID from the fiscal year of 1999.

For adequate treatment, prevention and control of infectious diseases on personal and district levels to the national level, high quality surveillance accurately comprehending the trend of the diseases and the infectious agents is essential. To realize this purpose, the understanding and cooperation of people in many different fields is required in operating the NESID program.

# Table 1. Target diseases of the Infectious Diseases Control Law, Japan (Nov. 5, 2003-)

(Reportable infectious diseases of categories I-V under the National Epidemiological Surveillance of Infectious Diseases)

Respirator	y Syndrome (SARS)*, Smallpox*
Category II	(to be notified all the cases promptly after diagnosis)
Acute pol	iomyelitis*, Cholera*, Diphtheria*, Paratyphoid fever*, Shigellosis*, Typhoid fever*
Category II	(to be notified all the cases promptly after diagnosis)
Enterohen	norrhagic <i>Escherichia coli</i> infection*
New catego Anthrax*, Epidemic Herpes B Lyssavirus Relapsing Yellow fer	<b>y IV</b> (to be notified all the cases promptly after diagnosis) Avian influenza virus infection*, Botulism*, Brucellosis*, Coccidioidomycosis*, Dengue fever*, Echinococco typhus*, Hantavirus pulmonary syndrome*, Hemorrhagic fever with renal syndrome*, Hepatitis A, Hepatitis virus infection*, Japanese encephalitis*, Japanese spotted fever*, Legionellosis*, Leptospirosis*, Lyme disea s infection (excluding rabies)*, Malaria, Monkeypox*, Nipah virus infection*, Psittacosis*, Q fever*, Rabies fever*, Scrub typhus (Tsutsugamushi disease)*, Tularemia*, West Nile fever (including West Nile encephalitiver*
New categor	y V
a. Diseases t	o be notified all the cases by all physicians within 7 days after diagnosis
Acquired	immunodeficiency syndrome*, Amebiasis*, <u>Acute encephalitis (excluding Japanese encephalitis and West Ni</u>
encephalit	is)*, Congenital rubella syndrome*, Creutzfeldt-Jakob disease*, Cryptosporidiosis, Giardiasis, Severe invasi
streptococ	cal infections (Streptococcal toxic shock-like syndrome)*, Syphilis, Meningococcal meningitis*, Tetanus*,
Vancomyo	cin-resistant <i>Enterococcus</i> infection*, <u>Vancomycin-resistant <i>Staphylococcus aureus</i> infection*</u> , Viral hepatitis
(excluding	g hepatitis A and E)*,
b. Diseases t	o be reported by the sentinel clinics and hospitals
<influenza s<="" td=""><td>entinel&gt; (weekly report)</td></influenza>	entinel> (weekly report)
Influenza	(excluding avian influenza virus infection)*
<pediatric di<="" td=""><td>sease sentinel&gt; (weekly report)</td></pediatric>	sease sentinel> (weekly report)
Chickenpo	xx, Erythema infectiosum, Exanthem subitum, Group A streptococcal pharyngitis*, Hand, foot and mouth dise
Herpangir	a*, Infectious gastroenteritis*, Measles (excluding adult)*, Mumps*, Pertussis*, Pharyngoconjunctival fever
Respirator	<u>y syncytial virus infection</u> *, Rubella,
<eye disease<="" td=""><td>e sentinel&gt; (weekly report)</td></eye>	e sentinel> (weekly report)
Acute hen	norrhagic conjunctivitis*, Epidemic keratoconjunctivitis*
<sexually td="" tr<=""><td>ansmitted disease (STD) sentinel&gt; (monthly report)</td></sexually>	ansmitted disease (STD) sentinel> (monthly report)
Condylor	a acuminatum, Genital chlamydial infection, Genital herpes, Gonorrhea
<target dise<="" td=""><td>ases at sentinel hospital&gt;</td></target>	ases at sentinel hospital>
(weekly rep	ort)
Aseptic m	eningitis*, Bacterial meningitis*, Chlamydial pneumonia (excluding psittacosis), Measles in adults*, Mycopla
pneumoni	a
(monthly re	port)
Methicilli	n-resistant Staphylococcus aureus infection, Multi-drug-resistant Pseudomonas aeruginosa infection,
Penicillin	resistant Strentococcus meumoniae infection