

MARCA DA BOLLO
Ministero dell'Economia
e delle Finanze
€16,00
SEDICI/00



Ministero della Salute

DIPARTIMENTO DELLA PROGRAMMAZIONE E DELL'ORDINAMENTO DEL
SERVIZIO SANITARIO NAZIONALE
DIREZIONE GENERALE DEI DISPOSITIVI MEDICI, DEL SERVIZIO FARMACEUTICO E
DELLA SICUREZZA DELLE CURE

PRODOTTI DI INTERESSE SANITARIO DIVERSI DAI DISPOSITIVI MEDICI

UFFICIO 07 EX DGFDM

D.G.D.F.S.C./ I.5.i.d.2/2011/369

IT/2014/00.173/AUT

IL DIRETTORE

VISTA la direttiva 98/8/CE del Parlamento Europeo e del Consiglio del 16 febbraio 1998 relativa all'immissione sul mercato dei biocidi;

VISTO il D. Lgs. 25 febbraio 2000, n. 174 recante "Attuazione della direttiva 98/8/CE in materia di immissione sul mercato di biocidi";

VISTO, in particolare, l'articolo 3, riguardante l'autorizzazione all'immissione in commercio dei prodotti biocidi;

VISTO il Regolamento (UE) 528/2012 del Parlamento Europeo e del Consiglio del 22 maggio 2012 relativo alla messa a disposizione sul mercato e all'uso del biocidi;

VISTO il D. Lgs. 30 marzo 2001, n. 165 recante norme generali sull'ordinamento del lavoro alle dipendenze delle amministrazioni pubbliche e successive modifiche;

VISTA la legge 15 luglio 2002, n.145 riportante disposizioni per il riordino della dirigenza statale;

VISTA la legge 13 novembre 2009, n. 172, recante l'istituzione del Ministero della salute;

VISTO il D.M. 12 settembre 2003, recante l'individuazione degli uffici dirigenziali di livello non generale del Ministero della salute, come modificato dal D.M. 23 giugno 2004;

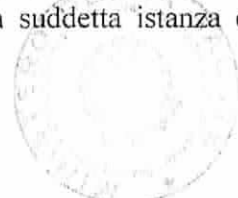
VISTO il D.P.R. 11 marzo 2011 n. 108 recante "Regolamento di organizzazione del Ministero della Salute";

VISTA la direttiva 2010/10/UE della Commissione del 9 febbraio 2010 recante modifica della direttiva 98/8/CE del Parlamento europeo e del Consiglio al fine di iscrivere il **BRODIFACOUM** come principio attivo nell'allegato 1 della direttiva;

VISTO il D.M. del 24 febbraio 2011., recante ad oggetto "Attuazione della direttiva 2010/10/CE., recante modifica della direttiva 98/8/CE del Parlamento europeo e del Consiglio del 16 febbraio 1998, relativa all'immissione sul mercato dei biocidi al fine di procedere all'inclusione della sostanza attiva **BRODIFACOUM** nell'allegato I della direttiva";

VISTA l'istanza, di cui alla nota del 26 gennaio 2012 (acquisita al prot. n. 5107 del 26 gennaio 2012), con cui la società **ACTIVA S.r.l.** Via Feltre 32 20132 Milano partita IVA n. 07509960154 ha chiesto l'autorizzazione del prodotto biocida denominato **ACTIPASTA - BROD** che sarà prodotto, confezionato e controllato presso l'officina I.N.D.I.A Industrie Chimiche S.p.a. Via Nona Strada 57 Z.I. Padova e distribuito da **ACTIVA S.r.l.** Via Feltre 32 20132 Milano con composizione di cui all'allegato 1 che forma parte integrante e sostanziale del presente decreto;

VISTA la documentazione presentata dalla società istante a sostegno della suddetta istanza di autorizzazione;



RITENUTA espletata l'istruttoria tecnica effettuata dall'ufficio competente, operante presso la Direzione generale dei dispositivi medici, del servizio farmaceutico e della sicurezza delle cure del Ministero della salute;

VISTO il parere dell'Istituto Superiore di Sanità, acquisito con prot. n. 0084688 del 22/11/2013;

VISTA la nota acquisita con prot. n. 7531 del 29/01/2014, con cui il Ministero dell'ambiente e della tutela del territorio e del mare ha espresso, per gli aspetti di competenza, parere favorevole all'autorizzazione, ai sensi dell'art. 3 del D. Lgs. 174/2000 e successive modifiche;

RITENUTA la conformità di detta documentazione alla normativa vigente in materia di immissione sul mercato di biocidi;

ATTESO che la società ha adempiuto agli obblighi previsti dal D.M. 16 aprile 2004 recante ad oggetto "Determinazione delle tariffe relative al programma di revisione ed all'immissione in commercio di biocidi";

DECRETA:

L'autorizzazione del prodotto biocida:

DENOMINAZIONE	ACTIPASTA - BROD
TITOLARE DELL'AUTORIZZAZIONE E RESPONSABILE DELL'IMMISSIONE SUL MERCATO ▪ Sede legale ▪ Direzione Commerciale	ACTIVA S.r.l. Via Feltre 32 20132 Milano
OFFICINE DI PRODUZIONE	I.N.D.I.A Industrie Chimimiche S.p.a. Via Nona Strada 57 Z.I. Padova
SOSTANZA ATTIVA	BRODIFACOUM (CAS N 56073-10-0)
PT	14 RODENTICIDA
DESCRIZIONE PRODOTTO	Esca rodenticida pronta all'uso in pasta fresca per uso professionale e non professionale
CONFEZIONI/TAGLIE	Per uso non professionale: 40-60-100-160-200-260-500 g (contenenti esche rodenticide predosate da 20g cadauna pronte all'uso in sacchetti di materiale ad uso alimentare) Per uso professionale 1-3-5-10-20-25 kg (contenenti esche rodenticide predosate da 20g cadauna pronte all'uso in sacchetti di materiale ad uso alimentare).
CATEGORIA DI UTILIZZATORI	Professionale e non professionale
DISTRIBUTORI	ACTIVA S.r.l. Via Feltre 32 20132 Milano
STABILITA' PRODOTTO	24 mesi
NUMERO DI AUTORIZZAZIONE	IT/2014/00 / 176 / Aut 5/10/2014
SCADENZA AUTORIZZAZIONE	31 gennaio 2017
COMPOSIZIONE DEL PRODOTTO	Vedi allegato 1 (dato confidenziale)

Sono approvate e fanno parte integrante del presente decreto le allegate etichette (allegato 2) con cui il citato biocida sarà immesso sul mercato.

L'esatta denominazione del biocida **ACTIPASTA - BROD** dovrà comunque essere individuabile attraverso una colorazione e un carattere unici in contrasto con le altre eventuali colorazioni e caratteri usati nel testo degli stampati autorizzati.

A far data dalla notifica del presente decreto, la ditta **ACTIVA S.r.l.** è tenuta a produrre e commercializzare il prodotto esclusivamente come biocida alle condizioni sopra riportate.

Il presente decreto viene redatto in duplice originale, di cui un esemplare è notificato in via amministrativa alla ditta interessata e l'altro è conservato agli atti di questo Ufficio.

Roma, li

5 FEB 2014

IL DIRETTORE
UFFICIO VII/DGFDM
(Dr.ssa Paola D'Alessandro)

MRL/AU

Annex 1: Summary of product characteristics

(a) Product trade name: ACTIPASTA-BROD

(b) (i) Qualitative and quantitative information on the composition of the biocidal product

NB: This information is confidential and should not be disclosed to third parties

Active substance(s) Common name	IUPAC name	CAS number	EC number	Concentration	Contents		Minimum purity (% w/w)	Same source as for Annex I inclusion
					Unit ⁸	w/w (%)		

Brodifacoum	3-[3-(4'-bromobiphenyl-4-yl)-1,2,3,4-tetrahydro-1-naphthyl]-4-hydroxycoumarin	56073-10-0	259-980-5	0.05	g/kg	0.005	99.2	<input type="checkbox"/> yes <input type="checkbox"/> no
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				Contents					
Co-formulants Common name	IUPAC name	Function	CAS number	EC number	Concentration	Unit	w/w (%)	Classification	Substance of concern
Confidential – see R4BP									

⁸ g/l, g/kg, other. For biological products, the concentration should state the number of activity units/units of potency (as appropriate) per defined unit of formulation (e.g. per gramme or per litre).

Traduzione dall'italiano in inglese del decreto AUT
ACTIPASTA-BROD

Stamp duty € 16,00
Stamp: Ministry of
Health

Ministero dell'Economia
e delle Finanze
MARCA DA BOLLO
€16,00
SEDDICI/08
Entrate
00001895 000076ZF 001CE001
00024293 27/02/2014 18:50:20
4578-00088 ZFA04A7C33A688DD
IDENTIFICATIVO : 01121176264267



Ministry of Health

**DEPARTMENT OF PLANNING AND ORGANIZATION OF THE NATIONAL HEALTH
SERVICE**

General Directorate for Medical Devices, Pharmaceutical Services
and Safety in Care

OFFICE 07 FORMER DGFDM

D.G.D.F.S.C./I.5.i.d.2/2011/369

IT/2014/00179/AUT

THE ACTING DIRECTOR

GIVEN Directive 98/8/EC of the European Parliament and Council of February 16, 1998 concerning the placement of biocidal products on the market;
GIVEN Legislative Decree No. 174 of February 25, 2000 on "Implementation of Directive 98/8/EC concerning the placement of biocidal products on the market";
GIVEN, in particular, Article 3 concerning authorization to place biocidal products on the market;
GIVEN Regulation (EU) No. 528/2012 of the European Parliament and Council of May 22, 2012 concerning the availability and use of biocidal products on the market;
GIVEN Legislative Decree No. 165 of March 30, 2001, containing general rules on the order of government employment, as amended;
GIVEN Law No. 145 of July 15, 2002 providing instructions for the reorder of government management;
GIVEN Law No. 172 of November 13, 2009 concerning the establishment of the Ministry of Health;
GIVEN Ministerial Decree of September 12, 2003 concerning the identification of non-executive office levels of the Ministry of Health, as amended by the Ministerial Decree of June 23, 2004;
GIVEN Decree of the President of the Republic No. 108 of March 11, 2011 on "Organization Rules of the Ministry of Health";
GIVEN Directive 2010/10/EU of the Commission dated February 09, 2010 amending Directive 98/8/EC of the European Parliament and Council with the scope of registering **BRODIFACOU** as active substance in Annex 1 of the Directive itself;
GIVEN the Decree of the Ministry of Health dated April 24, 2011, on the subject "Implementation of Directive 2010/10/EU, amending Directive 98/8/EC of the European Parliament and Council of February 16, 1998, concerning the placement of biocidal products on the market in order to proceed with including the active substance **BRODIFACOU** in Annex 1 of the Directive itself;
GIVEN the request contained in note dated January 26, 2012 (acquired under Protocol No. 5107 of January 26, 2012), with which the company **ACTIVA S.R.L.** registered office in 20132 Milano

Via Feltre N°32 - Tax Code and VAT No. 07509960154, has requested authorization for the biocidal product called "**ACTIPASTA-BROD**", which will be produced, packaged and controlled at the workshop of I.N.D.I.A. Industrie Chimiche S.p.A., Nona Strada - Z.I. - Padova and distributed by ACTIVA S.R.L. Via Feltre N°32 20132 Milano, composed as per Annex 1 which forms an integral and substantial part of this Decree;

GIVEN the documentation submitted by the requesting company in support of the abovementioned request for authorization.

CONSIDERING that the technical appraisals have been carried out by the relevant office, at the General Directorate for medical devices, pharmaceutical services and safety in care of the Ministry of Health;

GIVEN the opinion of the Higher Institute of Health, acquired with protocol N° 0084688 of November 22, 2013;

GIVEN the note acquired with protocol N° 7531 of January 29, 2014 with which the Ministry of Environment and Protection of Land and Sea, for those aspects of its expertise, approved the authorization of the above mentioned biocide, in accordance with Article 3 of Legislative Decree 174/2000, as amended;

CONSIDERING that this documentation complies with regulations in force regarding the marketing of biocides;

SINCE the company has fulfilled the obligations foreseen under Ministerial Decree dated April 16, 2004 concerning the "Determination of fees for the review program and marketing of biocidal products";

DECREES:

The authorization of the biocidal product:

NAME	ACTIPASTA - BROD
HOLDER OF THE AUTHORIZATION AND RESPONSIBLE FOR PLACEMENT ON THE MARKET	ACTIVA S.r.l.
<ul style="list-style-type: none"> ▪ Registered Office ▪ Sales Management 	Via Feltre, 12 - 20132 Milan
PRODUCTION WORKSHOPS	I.N.D.I.A. Industrie Chimiche S.p.A. - Nona Strada 57 - Z.I. Padova
ACTIVE INGREDIENT	BRODIFACOUM (CAS N 56073-10-0)
PT	14 - Rodenticides
PRODUCT DESCRIPTION	Ready to use Rodenticide bait in fresh paste - For professional and non professional use.
PACKAGES/SIZES	For non professional use 40-60-100-160-200-260-500 gr. (containing rodenticide baits, 20 gr. each, in food wrapping paper). For professional use 1-3-5-10-20-25 kg. (containing rodenticide baits, 20 gr. each, in food wrapping paper).
USER CATEGORY	Professionals and non

	professionals
DISTRIBUTOR	ACTIVA S.r.l. Via Feltre, 12 - 20132 Milan
PRODUCT STABILITY	24 months
AUTHORIZATION NUMBER	IT2014/00179/AUT (stamped February 05, 2014)
AUTHORIZATION EXPIRES	January 31, 2017
PRODUCT COMPOSITION	See Annex 1 (Restricted information)

The attached labels (Annex 2) with which the biocide will be placed on the market are approved and are an integral part of this decree.

The exact name of the biocide "ACTIPASTA-BROD" must however be identifiable by a unique colour and font, different from any other colours and fonts used in the printed text of authorized printed material.

Effective on the date this decree is notified, the company ACTIVA S.r.l. is required to produce and market the product exclusively as a biocide under the above conditions.

This decree is prepared in two original copies, one copy of which is administratively notified to the company involved and the other of which is stored in the files of this Office.

Rome, February 04, 2014

THE ACTING DIRECTOR
OFFICE 07 DGFDM

Dr.ssa Paola D'Allessandro
Signature

ANNEX 1

Stamp: *Ministry of Health*

Annex 1: Summary of product characteristics

(a) Product trade name: ACTIPASTA-BROD

(b) (i) Qualitative and quantitative information on the composition of the biocidal product

NB: This information is confidential and should not be disclosed to third parties

Active substance(s) Common name	IUPAC name	CAS number	EC number	Contents			Same source as for Annex I inclusion
				Concentration	Unit ⁸	w/w (%)	

Brodifacoum	3-[3-(4'-bromobiphenyl-4-yl)-1,2,3,4-tetrahydro-1-naphthyl]-4-hydroxycoumarin	56073-10-0	259-980-5	0.05	g/kg	0.005	99.2	<input type="checkbox"/> yes <input type="checkbox"/> no
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Co-formulants				Contents				
Common name	IUPAC name	CAS number	EC number	Concentration	Unit	w/w (%)	Classification	Substance of concern
Confidential – see R4BP								

⁸ g/l, g/kg, other. For biological products, the concentration should state the number of activity units/units of potency (as appropriate) per defined unit of formulation (e.g. per gramme or per litre).

ACTIPASTA - BROD

**Ready to use rodenticide bait in fresh paste
For professional use
To use, carefully read the label instructions**



COMPOSITION

100 g of product contain:

Brodifacoum (N°CAS 56073-10-0)	0,005 g	
Denatonium Benzoate		0,001 g
Edible substances and coformulants:	up to g	100

BIOCIDAL PRODUCT (PT14)

Authorisation of the Italian Ministry of Health n.

IT/2014/00179/AUT

Authorisation Holder:

**ACTIVA S.r.l., Via Feltre, 32- 20132 Milano- Italy - Tel. +39
02.70637301**

Manufacturing, packing and quality control plant:

**I.N.D.I.A. Industrie Chimiche S.p.A. - Nona Strada 57 - Z.I.
Padova -Italy**

**Distributed by: ACTIVA S.r.l., Via Feltre, 32- 20132 Milano -
Italy- Tel. +39 02.70637301**

Packs: 1-3-5-10-20-25 Kg

**(containing rodenticide baits of 20 g of fresh paste in food
sachet, ready to use)**

Batch n°..... date

Validity: 24 months

SAFETY PHRASES

The packages should be stored under lock and keep out of the reach of children. Keep away from food, drink and animal feedstuff. When using do not eat, drink or smoke. Avoid contact with skin. If swallowed, seek medical advice immediately and show this container or label. Do not release in the environment. Refer to warnings label.

MEDICAL INFORMATION

MECHANISM OF ACTION: The active substance in the product is an antagonist competitor of vitamin K and decreases the hepatic synthesis of K-dependent factors.

SYMPTOMS: In case of ingestion of large quantities, severe poisoning inhibits vitamin K, causing bleeding in the skin and mucosas and at level of organs and parenchymas.

TREATMENT: In case of ingestion of large quantities, induce vomiting, perform gastrolavage and administer activated carbon. The prothrombin activity must be monitored immediately after ingestion and in the following days. If prothrombin activity is

reduced, administered vitamin K. Follow proper medical protocol in agreement with a Poison Control Centre.

CONTRAINDICATION: anticoagulants

WARNINGS

Bait boxes must be securely deposited in order to avoid tampering and accidental ingestion by children or non-target animals. Always ensure that bait boxes are secured properly and that the bait cannot be dragged away by rodents. Avoid treatments in the presence of not protected food or feed. Not to use in agriculture. Do not reuse the empty pack and not release in the environment, but dispose in accordance with current rules. In case of contamination wash hands thoroughly with soap and plenty of water. In case of suspected ingestion consult a Poison Control Center. The product can be dangerous if swallowed by pets or other non-target animals. If swallowed, induce vomiting. Contact your veterinarian immediately and show the container or label.

CHARACTERISTICS OF THE PRODUCT

ACTIPASTA-BROD is a ready to use rodenticide bait based on an anticoagulant active substance known as Brodifacoum, effective against House mouse (*Mus musculus*), Brown rat (*Rattus norvegicus*), Black rat (*Rattus rattus*) after a single ingestion. The formulation consists in an attractive "fresh paste" for all the above mentioned rodent species. The product is not a warning for all the rest of rodent population

ACTIPASTA-BROD contains a bittering agent to prevent accidental ingestion by children.

ACTIPASTA-BROD can be used in and around civilian and industrial buildings (warehouses and holds included), in cellars, garages, closets and gardens of property, farms and houses.

INSTRUCTIONS AND DOSES FOR USE

Product is ready to use and must be used according doses written below. Avoid touching the product with bare hands, use protective gloves. Inner packs must be handled carefully and placed, as they are, into the bait box, according the doses written below, protected from the atmospheric agents, the ingestion of non-target species and in order to avoid dispersion in the environment. Observe the area of infestation and place bait boxes with baits at the dens, along rodents routes and places of major attendance.

Recommended doses are:

House mouse: 2 baits (40g) x 100 sqm

Brown rat: 3-5 baits (60-100g) x 100 sqm

Black rat: 3-5 baits (60-100g) x 100 sqm

Mark the area of the treatment and place a notice

Check weekly bait boxes and displace baits eaten. Search for and remove dead rodents at frequent intervals disposing according current rules. Do not dispose dead animals in waste or dumps. Use proper gloves to manipulate dead animals. The product is not intended for permanent use, organize treatments lasting up to 6

weeks. At the end of treatment remove bait boxes and eliminate the bait remained according to current rules. Carefully read the label instructions. Read carefully the Safety Data Sheet.
PRODUCT INTENDED ONLY FOR PROFESSIONAL USE

ACTIPASTA - BROD

**Ready to use rodenticide bait in fresh paste
For non professional use
To use, carefully read the label instructions**

COMPOSITION

100 g of product contain:

Brodifacoum (N°CAS 56073-10-0)	0,005 g	
Denatonium Benzoate		0,001 g
Edible substances and coformulants:	up to g	100

BIOCIDAL PRODUCT (PT14)

Authorisation of the Italian Ministry of Health no.

IT/2014/00179/AUT

Authorisation Holder

**ACTIVA S.r.l., Via Feltre, 32- 20132 Milano -Italy- Tel. +39
02.70637301**

Manufacturing, packing and quality control plant:

**I.N.D.I.A. Industrie Chimiche S.p.A. - Nona Strada 57 - Z.I.
Padova-Italy**

**Distributed by: ACTIVA S.r.l., Via Feltre, 32- 20132 Milano-Italy
- Tel. +39 02.70637301**

**Packs: 40-60-100-160-200-260-500g
containing rodenticide baits of 20 g of fresh paste (in food
sachet, ready to use)**

Batch n° date Validity: 24 months

SAFETY PHRASES

The packages should be stored under lock and keep out of the reach of children. Keep away from food, drink and animal feedstuff. When using do not eat, drink or smoke. Avoid contact with skin. If swallowed, seek medical advice immediately and show this container or label. Do not release in the environment. Refer to warnings label.

MEDICAL INFORMATION

MECHANISM OF ACTION: The active substance in the product is an antagonist competitor of vitamin K and decreases the hepatic synthesis of K-dependent factors.

SYMPTOMS: severe poisoning from ingestion. Poisoning inhibits vitamin K, causing bleeding in the skin and mucosas and at level of organs and parenchymas.

TREATMENT: In case of ingestion of large quantities, induce vomiting, perform gastrolavage or administer activated carbon.

The prothrombin activity must be monitored immediately after ingestion and in the following days. If prothrombin activity is reduced, administered vitamin K. Follow proper medical protocol in agreement with a Poison Control Centre.

CONTRAINDICATION: anticoagulants

WARNINGS

Bait boxes must be securely deposited in order to avoid tampering and accidental ingestion by children or non-target animals. Always ensure that bait boxes are secured properly and that the bait cannot be dragged away by rodents. Avoid treatments in the presence of not protected food or feed. Not to use in agriculture. Do not reuse the empty pack and not release in the environment, but dispose in accordance with current rules. In case of contamination wash hands thoroughly with soap and plenty of water. In case of suspected ingestion consult a Poison Control Center. The product can be dangerous if swallowed by pets or other non-target animals. If swallowed, induce vomiting. Contact your veterinarian immediately and show the container or label.

CHARACTERISTICS OF THE PRODUCT

ACTIPASTA-BROD is a ready to use rodenticide bait based on an anticoagulant active substance known as Brodifacoum, effective against House mouse (*Mus musculus*), Brown rat (*Rattus norvegicus*), Black rat (*Rattus rattus*) after a single ingestion. The formulation consists in an attractive "fresh paste" for all the above mentioned rodent species. The product is not a warning for all the rest of rodent population

ACTIPASTA-BROD contains a bittering agent to prevent accidental ingestion by children.

ACTIPASTA-BROD can be used in and around buildings including industrial buildings, in cellars, garages, closets and gardens of property, farms and houses.

INSTRUCTIONS AND DOSES FOR USE

Product is ready to use and must be used according doses written below. Avoid touching the product with bare hands, use protective gloves. Inner packs must be handled carefully and placed, as they are, into the bait box, according the doses written below. The rodenticide baits must be placed in an appropriate bait box, tamper resistant and marked, available on the market. Observe the area of infestation and place bait boxes with baits at the dens, along rodents routes and places of major attendance.

Recommended doses are:

House mouse: 2 baits (40g) x 100 sqm

Brown rat: 3-5 baits (60-100g) x 100 sqm

Black rat: 3-5 baits (60-100g) x 100 sqm

Check weekly bait boxes and displace baits eaten. Search for and remove dead rodents at frequent intervals disposing according current rules. Do not dispose dead animals in waste or dumps. Use proper gloves to manipulate dead animals. The product is not intended for permanent use, organize treatments lasting up to 6

weeks. At the end of treatment remove bait boxes and eliminate the bait remained according to current rules. Carefully read the label instructions.

Monza, 27 febbraio 2014

Il traduttore giurato



Maria Luisa Bertoli



TRIBUNALE CIVILE E PENALE DI MONZA

VERBALE DI GIURAMENTO

N. cronologico: 512

In data 28 febbraio 2014, nella Cancelleria del Tribunale di Monza avanti al sottoscritto Cancelliere è personalmente comparsa la Dott.ssa Bertoli Maria Luisa nata a Seregno il 03-03-1972, residente a Seregno in Via Bologna, 31 ed iscritta nell'albo dei Traduttori del Tribunale di Monza al n. **1971/2012** per le lingue inglese e tedesco, la quale presenta la traduzione che precede e chiede di asseverarla con giuramento. Ammonita ai sensi di legge, la comparente presta il giuramento di rito ripetendo le parole: "Giuro di aver bene e fedelmente adempiuto le funzioni affidatemi al solo scopo di far conoscere la verità ai giudici".

Detto, confermato e sottoscritto



Maria Luisa Bertoli

[Signature]
IL FUNZIONARIO GIUDIZIARIO
Giacoma CIMINO

