

Toll Free Number for Rare Diseases

# Un numero raro, prezioso e gratuito.





## **Background**

The Minister of Health, the Honourable Senator Livia Turco:

- Has established the national committee of patients associations ("Consulta) for rare diseases on the 05 of June 2007
- Has designated The Istituto Superiore di Sanità as the institution with technical and scientific know-how to promote activities and meetings of the committee
- Has highlighted the importance of establishing in the Center of Rare Diseases-ISS a Toll Free number devoted to rare diseases



## The 6 workgroups of the Committee

3 workgroups have been activated on the 03 of July 2007:

- 1) To assess and certify disability and exemption
- 2) Continuity of care
- 3) Integration of social and medical care

Additional 3 groups have been activated on the 08 of September 2007:

- 4) Training of staff and uniformity of medical care nationally
- 5) Scientific Research
- 6) Information and Communication



### Critical aspects: lack of information

#### Possible solutions

- To standardise all the information supplied locally and nationally
- To elaborate specific technical and scientific forms about specific rare diseases
- To promote public campaigns on TV and radio and print leaflets about rare diseases
- To promote a national day devoted to rare diseases
- To collect and make valuable the informative material of local and regional associations and charities
- Ta make available throughout the web the material collected
- To ensure that help lines are available to patients, staff and citizens





## 29 February 2008 rare diseases day

The conference at the ISS and the other national activities

http://www.iss.it/site/cnmr/dina/29febbr2008/giornata.html





## **TVMR**

Toll Free Number for Rare Diseases

The service was launched in February 29, 2008: an active service to inform and help patients, citizens, operators and Associations



#### Telefono Verde Malattie Rare

The Toll Free Number for Rare Diseases (TVMR) is a public Service, managed by the National Centre for Rare Disease (CNMR).

TVMR is a **national service** giving the opportunity to spread information about rare diseases nationally.

The **service** is **anonymous** and **totally free** for all users.

Currently, TVMR is open from Monday to Friday from 9 a.m. to 1 p.m.

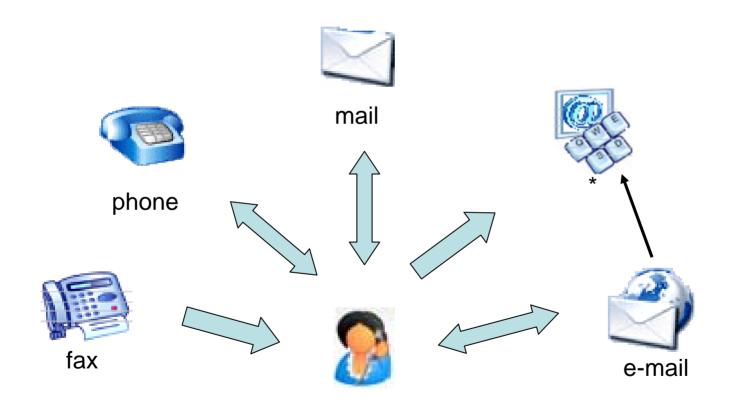


## **Objectives of TVMR**

- to inform about public health services caring and treating rare diseases nationally
- to give contacts and information about Patients
  Associations nationally
- to give support about legal and administrative questions related to rare diseases
- to give **scientific information** about rare diseases
- to support psychologically patients with rare diseases



## Means of communication utilised



## The methodology: Counselling

**Telephone counselling** is the methodology used by operators of TVMR. The counselling allows to:

<u>Provide scientific information</u> updated and personalised (useful for the specific need/request of the person calling): *informative counselling* 

<u>Contribute to identify the real problem</u> of the person calling in order to identify possible solutions: *problem solving counselling* 

Support the person during critical situation: crisis counselling

<u>Reduce anxiety and discomfort</u> in order to improve the communication and the understanding of the message/information provided facilitating behavior changes: *decision making counselling* 



## Information and communication













Ricerca scientifica finalizzata alla prevenzione, sorveglianza, diagnosi e trattamento delle malattie rare è la principale attività del CNMR dell'Istituto Superiore di Sanità.



Sono prodotti potenzialmente utili per trattare una malattia rara ma che non hanno un meruato sufficiente a ripagare le spese del loro sviluppo. Vengono definiti orfani proprio petthé è scarso l'interesse delle industrie farmaceutiche ad investire su farmaci destinati a pochi pazienti.





#### Workshops and meetings

- Workshops: on RD national register functioning, primary prevention, genetic testing
- International meeting: annual international meeting on orphan drugs and rare diseases