Major Hazards and People with Disabilities

Their Involvement in Disaster Preparedness and Response

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European and Mediterranean Major Hazards Agreement (EUR-OPA)
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Introduction

People with disabilities constitute a very large minority that consists of between one sixth and one fifth of the general population of most countries. When major incidents and disasters occur, people with disabilities face hardship that is potentially greater than that of the majority population, and they can suffer additional forms of discrimination or neglect. The moral and ethical case for an inclusive approach that guarantees the right of people with disabilities to adequate care in disasters is unassailable. Thus, warning, evacuation, shelter, transitional housing and other emergency provisions are services that need to be fully accessible and usable by a wide range of people with disabilities. Whereas measures for the general population are generally created for groups, a certain number of persons with disabilities require individual assistance, which may involve a fundamental reorientation in the way that civil protection services are planned and delivered.

It is vitally important to understand the needs of people with disabilities during the exceptional circumstances created by major incidents and disasters. It is also essential not to subsume these needs among those pertaining to minorities and disadvantaged groups in general. Emergency measures should seek to preserve the dignity and (where possible) the autonomy of people with disabilities. Academic and practical studies of disability and disaster reveal that there is a significant shortfall between the recognition of these principles and their implementation in practical programmes of action. The shortfall includes failure to design programmes and plans, implement them and monitor their effectiveness.

Planning is an essential part of preparing for emergencies. In order to ensure that resources, manpower and organisation are in place, plans and preparations need to be made at the national level, which should also be the level at which plans and measures...
are promoted and harmonised at the intermediate and local levels of public administration. Healthcare institutions, social services, and voluntary organisations in the fields of disability and civil protection need to work together at both the planning and response modes to create viable programmes of emergency care for people with disabilities. Coordination by a single, responsible government entity should nevertheless involve all the organisations involved in responding to emergencies on behalf of people with disabilities. It is important to note that all plans to assist people with disabilities are local in their implementation and outcome, and hence attention needs to be devoted to this level. Plans must be consolidated by frequent updating and testing, which should be complemented by programmes of training designed to ensure that all emergency responders are fully familiar with their roles, responsibilities and the procedures they will need to employ in a crisis or disaster.

In conclusion, people with disabilities, and the organisations that represent them, need to be drawn into the civil protection preparedness process. Policies and plans need to be inclusive, but the particular needs of people with disabilities should not be subsumed in a “compromising manner” into wider amalgamations of disadvantaged groups.

Preparing for disaster with and on behalf of people with disabilities requires political commitment, national and local coordination, strategic planning, networking, knowledge management, optimisation of resources and the development of good communication strategies. With this in view the Council of Europe, through the EUR-OPA Major Hazards Agreement and its member countries1, has developed pertinent Guidelines and a Recommendation.

In Europe and the Mediterranean area, countries are striving to improve their emergency preparedness. However, little has been done to include people with disabilities into practical programmes of action in civil protection. However, some examples of good practice do exist. These include the creation of specific offices to run programmes for protecting people with disabilities in disaster, ensuring that the problem is adequately dealt with in national disaster response legislation, finding innovative and alternative ways of disseminating warnings to people with cognitive problems, hearing impairments, or who do not understand the local language (for example, tourists, visitors and workers from other countries).

Examples of good practice from around the world highlight the importance of translating it to new situations and ensuring that lessons are learned by implementing them into improved outcomes. For example, evacuation needs require attention to accessibility issues and forms of alerting that take account of people’s disabilities. It also requires accessible transportation and shelter. Occupant emergency plans (OEPs) should be written for key buildings, and such instruments should take account of the needs of people with disabilities.

1 The member States of EUR-OPA are: Albania, Armenia, Azerbaijan, Belgium, Bosnia and Herzegovina, Bulgaria, Croatia, Cyprus, Georgia, Greece, Luxembourg, Malta, Republic of Moldova, Monaco, Portugal, Romania, Russian Federation, San Marino, Serbia, Spain, “The former Yugoslav Republic of Macedonia”, Turkey, Ukraine, and three Mediterranean countries which are not member States of the Council of Europe: Algeria, Lebanon, Morocco.
Key Requirements

to the attention of European Policy Makers
taking into account the Guidelines for
Assisting People with Disabilities during
Emergencies, Crises and Disasters

1 People with disabilities should receive support that is as good as that enjoyed by the general population.

2 The support should be tailored to the whole range of potential individual needs, and it should be recognised that, as a wide variety of disabilities is involved, needs will vary considerably from one person to another.

3 Planning for the care of people with disabilities should involve political authorities, public administrators, civil protection authorities and civil society organisations.

4 Emergency plans should consider persons with disabilities individually rather than as groups or categories.

5 The locations and emergency needs of people with disabilities should be known and assessed before disaster strikes.

6 Special emergency planning provisions should be made for care homes, psychiatric hospitals and other centres where people with disabilities are likely to be concentrated.

7 Education programmes for all who are involved in planning for, managing, responding to or recovering from disasters should include information on how to improve provisions for people with disabilities.

8 Alert processes should be configured in a way that automatically includes the needs and capacities of persons with disabilities (the “Design for All” principle).

1 Cf. “Universal Design”, the term used in Council of Europe documents
Evacuation, emergency transportation, sheltering and rehabilitation processes should not discriminate against people with disabilities but should ensure that their needs are catered for.

Emergency responders should maintain a correct, professional and non-discriminatory attitude to people with disabilities.

Emergency responders who are required to lift and transport people with physical impairments should receive appropriate training and have appropriate equipment for these tasks.

In pre- or post-disaster evacuation, procedures should be in place to ensure that no one is left behind.

Rest centres and temporary dormitories should be equipped to accommodate people with disabilities who are expected to use them.

People with disabilities should not suffer discrimination in the assignment of temporary, post-disaster accommodation, which should be accessible to them and designed to meet their essential needs.

Procedures should be put in place to ensure that people with disabilities are not discriminated against during planning, warning, alert, evacuation, emergency response, respite, transitional shelter or recovery from disaster. Cases of discrimination should be dealt with promptly and fairly.
With the gradual development of more enlightened attitudes, and the growing popularity of Paralympic athletics, movie actors with learning disabilities or autism, or with sensorial impairments, persons with Down’s Syndrome working as teachers, and many other examples where persons with disabilities have reached international recognition, disability has begun to shed some of its stigma and people with disabilities are at last beginning to be more accepted into the mainstream of European society. Although this recognition has been very valuable for showing a new picture of disability, it must be admitted that the people involved are exceptional and not representative for the large majority of people with disabilities.

In parallel, thanks to the impressive lobby work of disability organisations worldwide, the process of overall inclusion of people with disabilities has been helped by legislation that requires property owners, the managers of public amenities and local administrations to provide better facilities and access for people with disabilities, who are no longer “invisible citizens” and fight for their right to be present at all levels of social life.

Discrimination is still common enough to be a serious problem, and physical barriers to accessibility are still a long way from being abolished. According to a report by the BBC, the 2012 London Paralympics had a positive effect on public attitudes in the United Kingdom to disability, but it remains difficult to assess whether the impact is profound and enduring. Hostility to people with disabilities remains a serious problem, including attacks on individuals and guide dogs. In a difficult economic context, people with disabilities enter into competition with other groups of potentially disadvantaged individuals who are looking for jobs or attempting to remain in work (such as people from difficult social backgrounds, workers of a mature age and the long-term unemployed). Moreover cuts to disability benefits associated with the recession have caused serious hardship among those who depend on state funding to survive.

People with disabilities are numerous enough in society to constitute a very large minority. They are probably in the region of 15 per cent of the population of most countries, although any assessment of numbers depends upon the system of definition, registration and counting. In fact, it may be that with the inclusion of people who are not registered the figure rises to one in five of the general population. The forms of disability, and the degrees to which a person may be disabled, are extremely varied, and hence a true count of the numbers involved is difficult. Nevertheless,
people with disabilities may suffer, not only widespread discrimination, but also
unnecessary restrictions in what they can do, amounting to a form of social exclusion.
In addition, they may be economically disadvantaged relative to other members of
society through the difficulty of finding adequate employment, assuming that they
are able to work at all. Moreover, they may find it more difficult to manage their daily
routine than do people who do not live with disability.

One field in which progress is still very slow is the care of people with disabilities
in disasters, and their protection against disaster risk. Emergencies, crises and other
civil contingencies tend to distort or interrupt the pattern of everyday life and hence
are fertile occasions for the resurgence of prejudice, discrimination and neglect.
The excuse that “resources are insufficient” (including time and manpower) is not
particularly valid as, with proper planning and organisation, it is perfectly possible
to redress the balance in favour of caring for and protecting people with disabilities
against disasters.

This publication provides an overview of the state of the art in emergency
preparedness and disaster risk reduction for people with disabilities. It asks whether
an adequate level of resilience has been achieved by and on behalf of people with
disabilities. The publication begins by defining disability and clarifying questions
of ethics regarding the right of people with disabilities to an acceptable level of
protection in situations of high disaster risk. In considering the institutional framework
for achieving such protection, the publication considersthe implementation shortfall,
in which established principles have not been sufficiently activated. Planning, training
and exercising need to be improved, and examples of good practice adapted to
new environments and circumstances. The publication shows that there is much
potential to improve emergency preparedness for people with disabilities, and the
arguments for doing so are indisputable.

PeoPle wIth dIsabIlItIes –
defInItIons, facts and fIgures

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he UN Convention on the Rights of Persons with Disabilities (UNCRPD)
promotes the following definition: [people] “who have long-term physical,
mental, intellectual or sensory impairments which in interaction with
various barriers may hinder their full and effective participation in society on an
equal basis with others”. Disability can be physical or mental, partial or well-nigh
total, temporary or permanent, and constant or fluctuating. Hence it covers
a multitude of different conditions which represent a set of restrictions that may
reduce a person’s ability to participate fully in the regular activities of normal life,
or at least to do so without significant help from equipment, medication or carers.

The World Health Organization’s International Classification of Functioning,
Disability and Health (ICF18) adopts a broad approach in which impairment or
restriction of functioning is seen in relation to contextual factors related to a person’s
life and environment:

“Functioning is an umbrella term encompassing all body functions, activities and
participation; similarly, disability serves as an umbrella term for impairments, activity
limitations or participation restrictions. ICF also lists environmental factors that interact
with all these constructs. In this way, it enables the user to record useful profiles of
individuals’ functioning, disability and health in various domains.”

The broadest possible range of human conditions is taken into account in the
classification, which assesses what a person can do in both a standard environment
and in that person’s specific circumstances. Nevertheless, Paralympic athletics and
other examples have shown that people with disabilities are not necessarily frail,
and many people with disabilities have excelled in a wide range of activities or have
taken important leadership roles. This leads to the conclusion that the “normal person”
does not really exist and neither does the typical person with a disability. Implicitly,
the same human diversity must be taken into consideration in strategies, concepts
and solutions to be developed in any field.

1 Cf. WHO 2001. International Classification of Functioning, Disability and Health (ICF18). World Health Organization,
Classification of Functioning, Disability and Health. WHO/EIP/GPE/CAS/01.3, World Health Organization, Geneva,
22 pp.
The following list illustrates the variety of forms of disability:

- restricted mobility
- blindness or partial sight (possibly requiring the use of guide dogs)
- deafness and hearing impairment
- difficulties of cognition, communication and expression
- medical problems
- use of support systems to maintain vital functions
- psychiatric disturbances and panic attacks
- infirmity associated with old age.

The categories are not necessarily comprehensive or mutually exclusive.

It should also be noted that major incidents and disasters can create disability. In most cases, paraplegia and permanent physical impairment represent, thankfully, a small proportion of disaster-related injuries, but earthquakes, structural collapses and bombings can all cause lasting physical damage to people that in a minority of cases cannot be healed. In addition, although disaster does not usually increase the number of serious, long-term mental health cases, it does cause large rises in short- to medium-term psychological suffering that can disable people with respect to their normal mental state.

One common problem with institutional arrangements for dealing with people with disabilities is that as a category they may be lumped together with other disadvantaged, or potentially disadvantaged, groups. These may include pregnant women, children, elderly persons, the sick, the wounded, indigenous populations, environmentally displaced persons, and ethnic and religious minorities.¹ These groups may be the recipients of neglect, discrimination, racism, hostility, violence or repression or they may simply be more at risk of these things than are members of the general population. While it is important to ensure that measures are taken to fight prejudice, unfairness or victimisation in society, it is also important that the specific needs of people with disabilities as a very heterogeneous, group are not neglected in the process.

It is a general ethical principal that people with disabilities should be encouraged and helped to live independently and to play a full a role in mainstream society. While severe forms of mental or physical disability might preclude that, many people with disabilities have a good ability to participate strongly in a wide range of activities in regular society, and some are able to live autonomously. They should not be discouraged or prevented from doing so. The UN Convention on the Rights of Persons with Disabilities goes further and stresses the importance of respecting human rights.¹ It points out the responsibility of society to support the inclusion of people with disabilities.

People who live with severe disabilities have an ethical right to assistance from the rest of society that enables them to live their lives as fully as possible. In close-knit societies in which family sizes are large, most people with disabilities are cared for en famille. However, many others are not the beneficiaries of sufficient family resources and so require to be cared for by the state and civil society. State or charitable assistance may need to be financial, administrative, logistical, medical and social. Moreover, they need to respect the increasing wish and right of people with disabilities for autonomy and self-determination.

When disaster strikes, or a major incident occurs, people with disabilities have a right to receive the same level of protection as is given to the rest of society. In other words, disaster should not be the occasion to suspend fundamental human rights. This may require that people with disabilities receive a greater level of assistance than at other times. The help that is given should ensure that they are not put into a position of disadvantage with respect to vulnerability reduction, warning, evacuation, shelter, recovery, reconstruction, or any of the other fundamental actions associated with risk reduction and disaster response. Their basic human rights to privacy, courtesy, impartiality and measures to ensure their safety should not be infringed because they live with disability.

In many countries, these ethical goals are far from being achieved: however, they remain essential goals and no government, society, organisation or community

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should be deterred from striving to attain them. Disasters and emergencies represent a particular class of adversity. This should not be used as an excuse to reduce the level of attention, the magnitude of resources, or the strength of safeguards regarding the position and care of people with disabilities in society. In other words, there is no justification for relaxing either ethical or moral standards, or human rights, during a major incident or disaster.

THE IMPLEMENTATION SHORTFALL

Failure to make any provision for people with disabilities, including failure to appreciate the nature of their special needs, is still endemic in the modern world, including in parts of Europe. Lack of a registration system for people with disabilities is one of the most basic shortcomings and it is still widespread. This is a question of understanding the magnitude of the task, and it is of fundamental importance, because knowing who needs to be assisted, where they are located, what disabilities are involved and how many individuals will require helping is essential basic information that will underpin any viable programme of assistance.

- There is a significant number of academic studies of disability and there are some academic centres that have developed expertise in the study of people with disabilities in relation to disaster, notably at the Universities of Kansas in the USA and Leeds in the UK. However, systematic data on people with disabilities are lacking, academic theory has seldom been applied to the problem of protecting people with disabilities against disaster, and it is not common to include people with disabilities in evaluation methodologies and procedures. Hence, the implementation shortfall begins with a deficiency in data and analysis, or in other words in attaining a precise understanding of the issues at hand.

- The wide range of political systems, cultural differences and standards of living, combined with geographic and climatic differences, and diverse likelihoods for particular kinds of disasters, makes it very hard to develop a common set of criteria. Hence the approach must be based on a common means of identifying the challenges that must be faced in each context. The very first step is to collect reliable data, which at present are largely absent. Article 31 of the UNCRDP, which deals with statistics and data collection, stresses the importance of respecting the privacy of people with disabilities.

- Many countries do not have a register of people with disabilities. This requires that a formal definition of disability be adhered to and people who fall within it be required to register with health and social security authorities, and to maintain a record of their home addresses. The register needs to be available to local authorities and to be kept up to date. In normal times, registers are used to determine who receives living allowances from the state, if such exist. They also serve to identify needs that health authorities and social services can satisfy. If such provisions are lacking among state provisions, there is little incentive to compile the register. However, it
is something that can be used to good effect when disaster strikes, and during the formulation of emergency plans, as it will represent an inventory of special needs and the location of people who may be in need of assistance.

Unfortunately, allowances or similar benefits are very often provided by different kinds of administrative unit and are based on different definitions of who are the potential beneficiaries. The result can be a different definition of disability for the same person, depending on the kind of allowance for which he or she may apply.

The WHO International Classification of Functioning, Disability and Health (ICF18), which is endorsed by many states, endeavours to distinguish between the medical and social approaches to disability. However, it is a complicated tool that has not gained much popularity among people and organisations that work in the disability field. In contrast, the “Design for All” approach is becoming more popular, and it evolves in a completely opposite direction. Instead on disability, it focusses on human diversity, but this could make it even more difficult to ascertain who is to be considered a person with a disability. Moreover, in some countries the mobility limitations of citizens over 65 are linked purely to their age, and these persons are thus not defined as living with disabilities.

People with disabilities are not mentioned in a number of key legal instruments and policy declarations. For example, they are absent from the UN Millennium Declaration. In many countries they are not specifically mentioned in national disaster management laws. Even the landmark Americans with Disabilities Act (ADA, 1991) does not include any legislation regarding the evacuation of people with disabilities. Moreover, the 2004 ADA Guide for Local Governments, which is supposed to help city authorities to establish evacuation plans for individuals with disabilities, lacks enforcement.

The following are common aspects of the ‘implementation shortfall’:

- There is a tendency to subsume provisions for people with disabilities in an inefficient or compromising way into other forms of legislation, such as disaster response acts, health and safety legislation and general legislation on social welfare provisions. In order to honour the provision of European and UN conventions, arrangements for assisting people with disabilities in emergencies and disasters should not disappear in the general legislation - i.e. they should be specific in each country. However, specific measures should not over-concentrate responsibility in single organisation, leading others to relinquish their roles in caring for people with disabilities. The approach must remain holistic and must involve full collaboration between organisations.

- There may be failure to implement legislation, guidelines and organisational arrangements. This is usually attributed to lack of financial resources, but it can also mean a lack of political or administrative motivation to find and use the resources, or indeed to face up to the problem at all. Decentralised organisation of competencies has the merit of bringing services closer to the beneficiaries, but it may result in incomplete responses to need, as well as the division of those responses into highly separate sectors, the enemy of a holistic approach. The effect of such systems may be lack of completeness in the calculation of what resources are needed in order to help people with disabilities.

- Failure to monitor and develop programmes for the protection of people with disabilities in disaster is still common. This includes failure to inform and train emergency responders. This problem is increased by decentralisation of responsibility and lack of concerted action among services.

- Failure to apportion responsibilities among the various organisations and institutions involved in the care of people with disabilities during disasters is also a widespread deficiency. Either the responsibilities remain undefined, or they are not allotted in an efficient and functional manner. Hence, it is common for planning to assist people with disabilities in disaster to “fall down the cracks” and disappear into bureaucratic limbo. This risk looms even larger when large institutions that house people with disabilities set up their own emergency programs, unless these are competently and rigorously formulated and practised. However, demographic change and the trend to deinstitutionalise people with disabilities mean that such people increasingly do not live in institutions but in their own homes.

- Failure to ensure capillarity in national programmes is another problem. The legislation and organisational arrangements may be exemplary at the national level, but people with disabilities need to be assisted at the local level: hence, arrangements need to be securely in place at the intermediate and local tiers of government. Another case concerns of particular relevance in Europe is that of people who live in border regions, with a high probability of being dependent on foreign emergency system and on communicating their needs in a foreign language.

- Conflicts may arise from different interpretations of disability policies. While the organisations that work on behalf of people with disabilities insist on policies of inclusion, with the right to access any level of any building, fire fighters could usefully promote the view that, in order to avoid evacuation problems, wheelchair users should be allowed to work only at the lower levels of tall buildings.

- Lastly, there may be a funding shortfall. Disability organisations may lack access to mainstream funding connected with disaster mitigation, response and recovery. Civil protection agencies and local governments may be reluctant to devote funds destined for emergency preparedness to a single sector of the population, no matter how demonstrable are its needs. This problem can only be remedied by a serious rethink of priorities when defining the policies which govern the allocation of public or civil society (NGO) funds.

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During the recovery phase after a disaster, people with disabilities may endure discrimination, which may be intensified by scarcity of resources, deprivation or deterioration of life choices. Those who previously lived independently may find themselves moved to nursing homes, and more generally their degree of choice about living arrangements may be curtailed. Moreover, major disasters kill or injure the carers of people with disabilities. On the other hand, recovery and reconstruction can offer the opportunity to improve disability standards and inaugurate more inclusive forms of emergency planning. Article 32 of the UN Convention on the Rights of Persons with Disabilities promotes cooperation at the international level and can stimulate exchange of expertise in this field.

**INSTITUTIONAL FRAMEWORK**

In order to ensure fairness and equity governments must have policies that:

- guarantee and safeguard the fundamental rights of people with disabilities
- promote the inclusion of people with disabilities into mainstream society
- seek actively to prevent discrimination against people with disabilities
- create legal and administrative mechanisms to achieve these goals
- identify exactly which organisations are responsible for implementing and enforcing the policy
- are subject to impartial scrutiny and are regularly monitored with respect to their ethics, effectiveness and level of implementation.

Once again, the particular needs of people with disabilities should not be subsumed as some kind of compromise into general policies designed to ensure the rights of disadvantaged groups.

The deficiencies and failures listed in the previous section may add up to a failure to connect organs of central government, and tiers of government. Registries of people with disabilities are usually managed by a Ministry of Health or Social...
Services. Emergency planning and management may be a jurisdiction of, for example, the Ministry of the Interior (or Home Office). In countries (such as Sweden, Italy and the UK) in which emergency responses are a dependency of the national Cabinet, there may be more opportunity to connect the different competencies, but there is no inherent reason why that should occur automatically. Nor does such an arrangement guarantee liaison with outside organisations such as NGOs and voluntary associations, which may be necessary at the operational level in order to provide services for people with disabilities.

Answering the needs of people with disabilities in disaster requires a concerted effort by government at all levels, together with civil society in the form of families and voluntary associations. Institutionally there need to be strong links in several directions, as follows:

- between the national, intermediate and local tiers of government
- between healthcare, social services and other ministries, such as those that deal with emergency response (usually a Ministry of the Interior or Home Office)
- between the forms of public administration listed above and voluntary associations, and
- between civil protection organisations and the associations that care for people with disabilities or defend their rights.

Ideally, government provisions for people with disabilities will be coordinated and managed by a single entity that has a department or unit which deals with planning for emergencies. This agency would need to ensure capillarity throughout the system of public administration, or in other words that arrangements are implemented at the local level and fully backed by local administrations. As envisaged in article 33, concerning national implementation and monitoring of the UN Convention on the Rights of People with Disabilities, the organisational format includes focal points and monitoring entities that promulgate clear rules for coordinating, reporting and controlling. It seems to be a model with considerable promise for sharing and exchanging expertise at both the national and international levels.

One very important aspect of the institutional arrangements is to encourage civil protection forces to work together with associations that work on behalf of people with disabilities (see figure opposite). This is one way of connecting two sets of institutions with different competencies and agendas, and also of connecting public administration with the civil society organisations on which it depends to carry out tasks that are either outside the domain of the state or have been ceded to the voluntary sector, usually through lack of public resources.

In addition to fostering bilateral relationships, for example, between a civil protection agency and a voluntary organisation, there are other key players that need to be drawn into the process of preparing to assist people with disabilities in disaster. These include disability advocates and legal rights lawyers, experts in disability access, assisting technology experts and the managers of personal assistance services. All have a role to play and all need to know how that role will change during the straightened circumstances of a major incident or disaster.

The relationships described in this section should be implemented at the emergency planning stage, which is described in the next section of this publication.
PLANNING

In responding to public emergency situations (major incidents, disasters, contingencies, crises, etc.), there are three elements to programmed activities: plans, procedures and improvisation. Procedures or protocols are a form of guidance for conducting particular activities. Planning is the orchestration of procedures and activities into a concerted, multi-disciplinary approach to the process of emergency response (or, in other phases, mitigating risks or recovering from disaster). Most successful planning is based on one or more master documents that are periodically updated and are shared between the people who will use them. The ‘bedrock’ level of emergency planning is the local level, as this is always the “theatre of operations” when an emergency occurs. The essence of emergency planning is to predispose things (personnel, supplies, vehicles, communications, fuel, equipment, and so on) so that urgent needs can be met in the most efficient way possible.

Hence, planning should continue after the start of the emergency, and into the recovery from it, as the changing pattern of resources requires flexibility in how they can be applied to emerging needs. Throughout this process it will be necessary to improvise when particular contingencies have not been foreseen. However, avoidable improvisation is a form of inefficiency that, at its worst, can lead to unnecessary loss of life or destruction of property. Thus, the purpose of planning is to reduce improvisation to a necessary minimum by foreseeing needs that can be fulfilled in advance of the moment when they become imperative. Many forms of emergency intervention cannot easily be improvised as they require the acquisition of equipment and supplies, the training of personnel and the organisation of forces to carry out specific tasks. An additional element in the scheme can be information and training activities addressed directly to people with disabilities and the staff members who care for them.

This, then, is the matrix in which planning to cater for the emergency needs of people with disabilities should take place. The main difference between ‘standard’ emergency planning and that required for people with disabilities is that the former can be carried out on behalf of undifferentiated groups of people, while the latter has to take into consideration a range of supplemental and diverse needs. This adds up to a need for individual attention in each case.

It is a reasonable assumption that if emergency plans for the general population are inadequate or missing, planning for people with disabilities is unlikely to be successful. On the other hand, well-thought emergency plans based on the diverse needs of people with disabilities will work with the general population as well and will add better preparedness for unexpected situations. Hence, the one is a prerequisite for the other. However, emergency planning arrangements for the general population should not be allowed to subsume those for people with disabilities.

The first stage of all emergency planning is to collect data on hazards and vulnerability. Notwithstanding the growing number of persons with disabilities who exercise their right to autonomy and self-determination, and thus do not appear in any official data, wherever possible, planners should acquire adequate information on people with disabilities in the planning jurisdiction, including their addresses, ages, types of disability, and probable requirements during an emergency. At the same time, an assessment needs to be made of the nature of hazards that threaten the community and the probable local impacts of an extreme event. Obviously it will be impossible to have absolute numbers or absolutely complete information: therefore local emergency plans should not be designed in a rigid way, and all stakeholders must learn to cope with unforeseen situations.

The next stage is to design measures to safeguard the community, including people with disabilities, in relation to the inventory of resources (personnel, vehicles, equipment, supplies, communications, buildings and expertise) that will be available at any given time during an emergency. Where hazard impacts can be forecast with enough time to react, plans should make provision for warning and evacuating people with disabilities, with careful consideration as to what is involved in these processes. Where shelter is designated in advance, care should be taken to ensure that it is accessible to people with disabilities and meets their needs for privacy and functionality. For those people with disabilities who depend on medicines or medical equipment, arrangements need to be made to ensure continuity of supply and availability during an emergency and its aftermath. Meticulous planning may be required if there is any likelihood that electricity supplies will be interrupted, and there are people with disabilities who depend on medical or living aids that require electricity. Provisions may also need to be made to ensure that working animals such as guide dogs are properly accommodated and cared for in an emergency. The prevailing policy must be one of inclusion, not exclusion, of people with disabilities in emergency arrangements.

The rights of people with disabilities to be rescued and cared for after disaster are explained and codified in the Verona Charter1.

Ill emergency personnel need to be sensitive to the needs of people with disabilities. Hence, it is a good idea to provide a general training course that explains the issues. In most cases, courses should be preceded with some general diversity training. Once that has taken place, the relevant issues include the following:

- the prevalence of disability in modern society and the diversity of forms that it takes
- the moral and ethical case for inclusiveness
- legal and jurisdictional responsibilities of care
- planning provisions for people with disabilities
- scenarios for action when incidents occur or disaster strikes
- practical matters connected with first response to an incident or disaster.

The last of these issues may require its own specific training course for first responders in the emergency services, relevant branches of public administration and voluntary organisations. These courses should teach a combination of general principles and matters that are specific to local circumstances. Among the former are how to work with people with disabilities, how to lift and transfer bedridden individuals, how to ensure that warnings are disseminated to people who cannot hear or see adequately, and how to ensure that people with disabilities are able to maintain their dignity when evacuated to unfamiliar surroundings. Persons with disabilities and their organisations should be involved as teachers in the practical organisation of such training activities. On the one hand, this would allow participants to understand better the challenges and, on the other hand, it would enhance the level of commitment within disability communities.

Simulation exercises are an important extension of both planning and training. They can be carried out at three levels: by discussion (table-top exercise), as a command-post or command centre exercise, or in the field. Simulation of emergency and disaster conditions is one of the ways in which preparedness to meet civil contingencies can be increased. A simulation should be backed by adequate planning, both of the simulation event itself and of the emergency responses required, i.e. by creating and using viable emergency plans that remain as “standing orders” for future contingencies. In many countries such exercises are mandatory in institutions, and they allow one to “test” the effectiveness of practical co-operation with local emergency services.

The advantages of simulation exercises in the field are as follows:

- they sensitise the local population and participants to the need to prepare for disaster
- they help familiarise participants with the procedures they need to use during a major emergency
- they help organisations to work together under unfamiliar circumstances and under pressure
- they may reveal deficiencies of organisation, resources or preparation for a real emergency
- they can offer an opportunity to introduce the question of how to care for and assist people with disabilities during emergencies.
The Council of Europe’s Disability Action Plan 2006-2015 promotes the rights of people with disabilities and aims to help them improve their quality of life by “meeting country-specific conditions as well as transition processes that are taking place in various member states [...] and is intended to serve as a roadmap for policy makers, to enable them to design, adjust, refocus and implement appropriate plans, programmes and innovative strategies.”

In 2013 the Council of Europe began an initiative which involved sending a questionnaire on disaster risk reduction and emergency preparedness for people with disabilities to all 26 member countries of the European and Mediterranean Major Hazards Agreement (EUR-OPA) and other members of the Council of Europe. Twenty countries and two organisations responded. The result is a snapshot of the level of preparedness in European and Mediterranean countries.

Some responses were remarkably candid. Although it can be said that the responding nations are striving to improve their emergency preparedness, the level of provision for people with disabilities is relatively low and is also uneven from one country to another. There is little or no uniformity in the measures adopted and these are highly variable in their reach, effectiveness and level of implementation. Although the questionnaire did not enquire into the level of enforcement of government provisions, it can be assumed that there were many cases in which this was low as well.

Overall, there is a tendency to subsume measures for people with disabilities into general provisions for managing emergencies. There is also a tendency for responsibilities to be split between ministries and agencies, and for there to be no guarantee that communication and collaboration will be sufficient to produce viable measures.

Nevertheless, there are some examples of good practice that may serve as a beacon for other countries to emulate and adapt to their own circumstances and needs.

Various countries have organised psychological care in disasters at the national level, notably Austria, Bulgaria, Croatia, the Czech Republic, Italy, Latvia, Slovakia and Ukraine. Examples of emergencies in which the psychologists have intervened in an organised manner include floods, earthquakes, transportation crashes, and episodes of mass violence. Moreover, a European Commission-funded project, EUNAD, is designed to promote the integration of psycho-social care into disaster management on behalf of people with disabilities. This is very necessary, as, although the services are potentially very valuable to people with psychologically-related disabilities, they would benefit from being part of a comprehensive strategy that includes psychological support and other related services.

Examples of Good Practice

**Exam Ples of Good PractiCe**

However, there are also drawbacks to simulation exercises:

- they tend to be expensive and require considerable organisation, a process that generally takes 6-12 months of preparation, meetings and planning
- simulations tend to be artificial situations and to lack the urgency, spontaneity and realism of genuine emergencies
- the more realistic a simulation is, the more disruptive to normal life and the more it incurs potential safety risks
- usually, only part of the emergency arrangements can be tested in a simulation and
- there is a tendency not to repeat exercises, or not to do so on a regular basis or with any degree of frequency: hence, the experience acquired may gradually be lost afterwards.

Nevertheless, simulation exercises are valuable means of testing elements of the emergency response system, highlighting areas that need improvement, and raising awareness of issues. They present the opportunity to mobilise disability organisations in the cause of civil protection, train emergency responders in dealing with people with disabilities, and identify deficiencies in arrangements for rescuing, accommodating and assisting them when disaster strikes. The wise use of simulation is coupled with substantial efforts to observe and record processes in the field, and to match the experience with training initiatives. This “lessons-learning” approach can furnish valuable information about difficulties that would be encountered in a real emergency and how to anticipate them by designing an a priori solution.


have seldom been used in this context and have mostly been treated as services for the general population.

Bulgaria has identified a senior member of government who is responsible for coordinating policy, plans and measures in favour of people with disabilities in emergency and disaster situations. Measures for people with disabilities are an extension of the national Disaster Protection Act, which covers risk assessment, planning, risk reduction, resources, warning, emergency response and rescue procedures. The Bulgarian Government has a policy on the Integration of People with Disabilities, which is designed to take account of the special needs of people with disabilities and assist them actively. Measures for people with disabilities in the workplace are an extension of the national Health and Safety at Work Act, which, however, is primarily designed to reduce the incidents of accidents at work. Although the structure of disaster planning is well developed in Bulgaria, the provisions do not make specific provision for people with disabilities.

Miscellaneous limited examples of good practice in Europe include the following. In Belgium and other countries, text messages have been used to alert deaf people to emergency situations. In Norway, hospitals, nursing homes, and home care assistants have a general obligation to evacuate persons with reduced mobility in an emergency. In Greece people with special needs in schools are given an e-lesson under the programme “E-learning about earthquake protection for people with disabilities” of the European Centre on Prevention and Forecasting of Earthquakes. The Republic of Serbia has designed a pilot project to enable people with hearing and speech impairments to contact the emergency services on emergency numbers in case of need. The project was inaugurated in Belgrade in September 2013, and will gradually be expanded to the rest of the country.

Meanwhile, practical research is being conducted in Europe. A Centre for Disability Studies exists at the University of Leeds in the United Kingdom, and its researchers have conducted studies related to disaster preparedness. In the European Union, an Academic Network of European Disability Experts (ANED) has been created. With regard to specific disabilities, the “European Network for Psychosocial Crisis Management - Assisting Disabled in Case of Disaster” (EUNAD) project aims to help the disabled survivors of disasters. The project is designed to evaluate networks of associations for people with disabilities in terms of their levels of preparedness for disasters, to conduct further studies and to organise workshops. EUNAD will produce recommendations, a taskforce, and pilot training courses for different groups.

There have been significant developments outside Europe from which member countries of the EUR-OPA Agreement could derive inspiration. In New Zealand, for example, disaster services such as emergency call centres have been made accessible to persons with disabilities. Firefighters have created a unit in which officers speak sign language. During the aftermath of the Christchurch earthquake of February 2011, sign language interpreters were used in all television information sessions. A specific call centre was set up for persons with disabilities with the ability to address their needs or refer them to appropriate services. In the United States, the Federal Emergency Management Agency (FEMA) has dedicated part of its website to information resources for people with disabilities. The documentation offers advice and know-how to such people and explains projects connected with improving the access and support for disabled survivors of Hurricane Katrina.

The theme of the UNISDR 2013 International Day for Disaster Reduction was “living with disabilities and disasters.” This global initiative, which is centred on 13th October each year, is designed to promote resilience. UNISDR argues that solutions to the disasters problem must be fully inclusive. Moreover, decisions and policies to reduce disaster risks must reflect the needs of persons living with disabilities. Finally, investment in disaster risk reduction must satisfy the needs of persons who live with disabilities.

An EUR-OPA meeting also recorded the views of experts in the field. This enabled examples of good practice to be noted from further afield. For example, fire services in New Zealand have created a unit consisting of sign language interpreters. American practice recommends tailoring emergency access and egress routes of each building to its occupants, with the following provisions:

- shelter in place (SIP) plans
- buddy systems (although these are not to be relied upon solely, they are a good starting point)
- descent devices that will continue to operate in an emergency, including one in which electrical power is lost
- elevator (lift) policies
- the inclusion of service animals in emergency plans and drills
- multiple forms of communication and alert systems for the visually and hearing impaired.


For the safe, successful evacuation of people with disabilities, more attention should be given to the installation and use of guidance systems, to planning alternative accessible exits (when lifts are inoperative), to the installation of visual and acoustic alarms, to special signage for visually impaired persons and to ensuring that obstacles are not present along evacuation routes. Moreover, these provisions need to be robust so that they function during the early stages of a developing disaster impact. In countries such as the USA, these provisions are mandated by Occupational Health and Safety requirements for local governments, as these require an Occupant Emergency Plan (OEP) to be drawn up for major buildings.

In this regard, it is important to seek standardisation at the international level. Unfortunately, accessibility standards, guidance systems, pictograms, and other such measures are not yet harmonised, at either the European or the international levels. In view of this lack of standardisation, training activities can unfortunately be based only on local schemes.
CONCLUSIONS

The degree of civility in a society can be gauged by the way in which it treats disadvantaged people, including those who have disabilities. In this publication we have argued that, although people with disabilities are part of a larger constituency of disadvantaged groups and individuals, they are one that is remarkably heterogeneous in terms of both the range of disability and the needs it generates during emergencies and disasters.

■ Providing an adequate level of protection is a matter of taking the problem seriously, ensuring that the available structures, organisation and resources are adequate to cope with it, and vigorously promoting an active approach to the promotion of programmes designed to assist people with disabilities with reducing disaster risk and coping with emergencies. There also needs to be a process of monitoring and improvement, in order to adapt programmes to changes in society and to take advantage of potential innovations.

■ One key to the success of programmes is the inclusion of people with disabilities, and their representatives, in the decision-making process. Another is the inclusion of the programmes as an essential component of government policy-making and legislative processes. Providing greater safety for people with disabilities should be neither optional nor something that can be “left until later”. It is a matter of basic rights. Planning, training and the provision of information are activities that are essential in order to ensure that those rights are guaranteed. So are the processes of sharing information, learning from good practice and standardising approaches between European countries. With sufficient motivation and collaboration, the large minority of people in society who have disabilities can be protected in line with human rights obligations and basic ethical imperatives.

Guidelines for Assisting People with Disabilities during Emergencies, Crises and Disasters
In the words of the UN Convention on the Rights of Persons with Disabilities (UNCRPD), Article 11: Situations of risk and humanitarian emergencies:
“States Parties shall take, in accordance with their obligations under international law, including international humanitarian law and international human rights law, all necessary measures to ensure the protection and safety of persons with disabilities in situations of risk, including situations of armed conflict, humanitarian emergencies and the occurrence of natural disasters.”

In preparing for, reacting to and recovering from emergencies, crises and disasters, every effort should be made to ensure that people with disabilities are not discriminated against. The aim should be to ensure that the treatment and services they receive are as effective a form of support as that given to the able-bodied population. The Council of Europe Disability Action Plan 2006-2015 stresses the importance of “mainstreaming ... moving away from policies which support segregation.”

Leaders and decision makers in public administration, businesses and civil society who have responsibility for services that affect people with disabilities should accept that they may be more at risk than the general public, and may be more vulnerable to hazards, the consequences of a disaster, and unfair treatment during the event or its aftermath.

To ensure that people with disabilities receive adequate care is a matter of equity, fairness and justice, as well as an important affirmation of the values of civility. As the proportion of people with disabilities in society may be anywhere between 9 and 20 per cent, the problem is not one that can be ignored, nor should it be.

Whereas emergency preparedness for the non-disabled is usually provided to groups, it is necessary to consider the particular needs of people with disabilities more in detail. This is likely to be a resource-intensive process, and hence the provision of services should involve a wide variety of participants: official emergency managers and responders, volunteer groups in both emergency response and the care of people with disabilities, local and national social services, and others. On the other hand, including the needs of people with disabilities in traditional rescue-schemes will improve the quality of those schemes and will thus benefit many kinds of stakeholders.

Disability is not limited to restrictions on personal mobility such as those of people in wheelchairs. Instead it covers a very wide range of physical, sensorial, mental and emotional conditions, including the effects of old age or illness, and forms of dependency on medical drugs or equipment. With respect to the general
population, people with disabilities may be less able to perceive hazards and risks, or less mobile, or dependent upon assistance in order to be able to react to crisis situations.

Assisting people with disabilities during public emergencies requires planning, foresight and concerted action before disaster strikes, so that programmes and procedures are in place when the situation becomes critical. Improvisation is the enemy of good procedure and hence it needs to be reduced to a minimum by application of consultation and planning processes that predispose resources for use when they are needed.

This set of guidelines is intended to ensure that national governments, and their counterparts at regional and local level, civil society organisations and relevant offices in both the public and private sector obtain a clear idea of how to proceed with the provision of disaster risk reduction for people with disabilities. It begins with a set of working definitions and then considers the requirements of good preparedness during all the phases of crisis management: mitigation and planning (disaster risk reduction), alert, emergency action, and recovery. The care of people with disabilities needs to be considered with respect to all of these phases.

SUCCESSFUL IMPLEMENTATION

Disasters and their impacts can vary considerably from one place to another around the world and emergency response systems are strongly influenced by their political and cultural backgrounds. However, the conceptual approach to disaster risk management can be summarised in terms of a set of common factors.

1. Political commitment

Governments must make clear decisions and include in their political agenda the commitment to make a serious effort to develop effective disaster risk management for people with disabilities. As part of the more general endeavour to ensure the safety of their constituent populations, they must consistently pay attention to such people’s needs.

2. Co-ordination and continuity

In order to guarantee the effective development, application and monitoring of emergency systems for people with disabilities, one particular body of governmental administration must be responsible for co-ordination and the continuity of initiatives. In close cooperation with all relevant stakeholders, it will be the task of the co-ordinating body to make sure that all relevant information is collected and centralised.

3. Networking

At least one network should exist that allows stakeholders to meet and exchange information about the challenges to be met if risks are to be identified and solutions are to be found. These networks should always be open to new members and should take full account of evolutionary changes in technology, habits and expectations.

4. Strategic planning

A master plan should be set up and constantly updated. The organisation of training activities and the evaluation of emergency exercises should be part of a constant process of adaptation of the master plan.
5. Knowledge management

A coherent programme of knowledge management should be used to ensure the transfer of acquired know-how to those who can benefit from it. This knowledge would facilitate the organisation of training activities and allow emergency schemes constantly to be improved. Specific added value will be provided by the involvement of people with disabilities and their organisations.

6. Identification and optimisation of resources

The evaluation of a master plan and constant updating of its capacities, and the general level of knowledge, should allow stakeholders to estimate needs regarding financial, organisational and human resources. At the same time, the best possible use of existing or new resources may allow the action plan to be improved.

7. Communication

In order to ensure that everyone is kept informed about the state of preparedness, a good communication policy is needed. Energetic dissemination of information will ensure that more and more relevant stakeholders are contacted and involved in the preparedness process.

PHASES OF TRANSITION

Obviously the level of implementation for emergency strategies is very different at international level. In order to identify the level of preparedness, different phases can be identified, namely: awareness, inception, development and consolidation. These phases are summarised in the following table.

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Assuming that the political commitment is confirmed, the coordinating body has been identified and is operational, and the network has been set up, the elements of an action plan can now be defined. They include a preventative phase that emphasises the concept of disaster risk reduction, an operational phase that describes the emergency action and the follow-up phase after disasters, when rehabilitation must be organised.

**Prevention: Disaster Risk Reduction**

- Before disaster strikes, there should be a general process of mitigation, risk reduction, preparedness and planning. During the interval between disasters, it should be recognised and firmly established that in emergency situations, people with disabilities have a fundamental right to as good a level of protection as the rest of the population.

- When it comes to identifying the practical problems to be anticipated, the counter-disaster organisations of government and public administration should insist on involving civil protection voluntary organisations and organisations of people with disabilities, or that represent and assist such people, in a multilateral dialogue that is intended to promote planning and action.

- The overall goal of this phase is to ensure that people with disabilities are as resilient against disaster as the rest of the population, and that this level of resilience is satisfactory for all parties.

- Governments, public administrations and preparedness organisations should ensure that the rights and needs of people with disabilities form an integral part of the emergency planning process. This involves the following:
  - Knowing about people with disabilities by compiling records of addresses and needs for assistance, and where they are likely to be located. This may require using census data (with appropriate reference to legal requirements for privacy) or in the absence of registration systems for people with disabilities, conducting a survey of the local area. Hence, all beneficiaries with special needs during an emergency should be identified, quantified and registered with the civil protection authorities.
  - Common and specific hazards and risks should be considered in terms of how they affect people with disabilities, not merely how they impinge upon the general population.
  - Special needs are associated with care homes for the elderly, psychiatric hospitals and rehabilitation centres, as well as other special institutions that cater for people with severe disabilities who are unable to live in the community. These institutions represent concentrations of vulnerable people who may require special assistance during an emergency and should not be missed when designing preventative activities.
  - The needs of people with disabilities in an emergency must be estimated and resources found to cater for them. This process must recognise the individuality of particular needs resulting from disability and not overgeneralise them. It must be recognised that the needs of people with disabilities will be highly varied according to the types of disabilities involved, the living arrangements and the care and support services utilised by the individuals concerned.
  - Preventative emergency planning for people with disabilities should consider whether and how individuals are able to summon assistance, whether and to what extent rescuers are trained to deal with them, whether there are barriers to processes such as evacuation, and whether such processes are adequately endowed with resources, and whether appropriate temporary accommodation can be provided to people with disabilities if long-term evacuation is required.
  - Finally, planning should include measures to monitor, evaluate, and deal with discrimination against people with disabilities if it occurs during emergency, disaster or crisis situations.

- The approach to and respect for people with disabilities should be incorporated, as principles and as prescriptions for action, into training programmes for people who deal with disaster as planners, managers, decision makers or responders. Education for the contact with people with disabilities should extend to all phases of disaster: mitigation, alert, emergency action and recovery.

**Protection: Emergency Action**

This phase refers to an emergency or crisis situation in which responses such as rescue, evacuation and care are required. Those responders who are required to lift and transport physically people with disabilities, and the frail elderly, should be trained and equipped to carry out such actions in the proper, professional manner with minimal risk to the giver or receiver of assistance. All equipment and technical devices should be well maintained in order to be fully operational in case of an emergency.
Evacuation

Evacuation is one of the principal means of avoiding harm to people during threatening or crisis situations. It can be divided broadly into the pre-impact (preventative) kind and that which is practised during or after the impact (for rescue or the maintenance of public safety).

When evacuation is needed, civil protection authorities should have pre-existing procedures to ensure no one is left behind.

Arrangements should be made to ensure that people with disabilities are able to follow evacuation orders when these are given out by the authorities. This involves ensuring that departure, the journey and the arrival at destination can be conducted efficiently and in safety without undue delays or impediments—and under the same criteria of efficiency and safety as are applied to the general population. There should be no physical barriers to these three phases of movement: this involves checking for the presence of steps, that corridors are wide enough to permit passage, that manpower and transport are available and are suitably equipped, and that arrangements are in place for accommodating each type of disability.

Bedridden people who are unable to move themselves should be raised, dressed (if necessary) and transported by carers or responders who are trained in how to carry out this kind of work and who will use the proper procedures.

If people with disabilities are taken to rest centres, these should be planned and equipped so that they are accessible and able to accommodate such people as far as possible without additional hardship. The person with a disability should have access to any equipment that is essential to the normal maintenance of his or her health and safety, including, where necessary, medications and life-support machinery.

Rehabilitation

This phase refers to the aftermath of a disaster, crisis or emergency in which the emphasis is placed on restoring conditions to normal and recovering from damage and disruption. This may be a slow process that lasts years and requires a lengthy period of living with temporary arrangements.

Governments and public administrators should seek to ensure that people with disabilities are not discriminated against in the planning, design or assignment of temporary post-disaster shelter, which must be accessible and functional according to their needs. Moreover, people with disabilities should not be discriminated against in the provision of post-disaster employment opportunities, or in the assignment of permanently rebuilt housing.

Emergency responders and other carers must be required to maintain a correct, professional and non-discriminatory attitude to people with disabilities at all times.

On the basis of detailed knowledge of the people with disabilities who are likely to be present in the local area, detailed studies should be made of how each individual will perceive danger or receive an alert. In order to ensure that it is effective, the process of sending out an alarm should be studied in relation to the cognitive and sensorial capacities of each person to be alerted, or the needs of his or her carers.
As in the emergency phase, every effort should be made to accommodate working animals, such as guide dogs for the blind.

People with disabilities should not suffer higher levels of post-disaster risk than do the general population.

The presence of discrimination in any of the ways outlined here should be monitored regularly and, if it occurs, measures should be taken promptly to stop it and discipline or re-educate any staff who are guilty of exhibiting discriminatory attitudes or behaviours, or making decisions that cause discrimination.

Recommendation

on the Inclusion of People with Disabilities in Disaster Preparedness and Response
European and Mediterranean Major Hazards Agreement (EUR-OPA)

Recommendation 2013 - 1 of the Committee of Permanent Correspondents on the inclusion of people with disabilities in disaster preparedness and response, adopted at the 64th meeting of the Committee of Permanent Correspondents of the European and Mediterranean Major Hazards Agreement (EUR-OPA).
Paris, France, 24-25 October 2013

THE COMMITTEE OF PERMANENT CORRESPONDENTS OF THE EUROPEAN AND MEDITERRANEAN MAJOR HAZARDS AGREEMENT (EUR-OPA),

A Recognising that people with disabilities have a fundamental right to a degree of protection against disasters that is comparable with that enjoyed by the rest of the population and in other terms offers a satisfactory level of preparedness;

B Mindful that in most European countries awareness, planning, assistance, training and specialised emergency assistance to people with disabilities would benefit from significant improvement;

C Desirous to promote a more integrated, efficient and effective approach to the problem of ensuring that people with disabilities in Europe are protected against emergencies and disasters;

D Taking note with satisfaction of the report “Including People with Disabilities in Disaster Preparedness and Response” by Prof. David Alexander (Institute for Risk and Disaster Reduction, University College London) and Mr Silvio Sagramola (Director of the Centre National Disability Information and Meeting Centre, Luxembourg), and of the “Guidelines for Assisting People with Disabilities during Emergencies, Crises and Disasters” by the same authors.

RECOMMEND THAT MEMBER STATES OF THE EUROPEAN AND MEDITERRANEAN MAJOR HAZARDS AGREEMENT (EUR-OPA):

1 Integrate specialised measures for people with disabilities into national disaster risk reduction policies, planning processes, training curricula and emergency response practice, favouring, as appropriate, investment in longterm strategies that would reduce the vulnerability and exposure to disaster of people with disabilities;

2 Design and promote national programmes and standards for the protection, rescue and care of people with disabilities when and where they are at risk of disaster;

3 Design and promote measures at the community level through local administrations and civil protection services, making use of local organisations that provide care and representation to people with disabilities;

4 Ensure that people with disabilities are included in the entire disaster risk reduction process and that, wherever possible, their viewpoints are taken fully into account;

5 Ensure that education and training for the protection and assistance of people with disabilities are firmly in the mainstream curricula for incident, crisis and disaster management and response;

6 Support the efforts of the Agreement to promote inclusive disaster risk reduction for people with disabilities in its member States mainly through the promotion of adequate regulations, the sharing of expertise, the organisation of training programmes and the promotion of good practice.

And invites the Secretariat to submit the above Recommendation to the Committee of Ministers for information and possible distribution to all member States of the Council of Europe.
Bibliography


Yuba County 2009. Emergency Plan Workbook for People with Disabilities and Their Families. Yuba County Health and Human Services Department, California, 14 pp.
In both academic and professional publications relating to emergencies, crises and disasters there is a wide variety of definitions of key terms, and some of them conflict with one another. However, definitions are necessary as a means of establishing the nature of the phenomena and processes that are under consideration. Putting aside differences of opinion on meanings, this section provides some simple working definitions of terms. When documents of international interest are being translated into national languages, there is always a risk of misunderstanding, and it is therefore important to make sure that translations do not distort the message to be transmitted.

**Handicap**
A physical, sensorial or mental condition that impairs a person’s ability to perceive or react to events around himself or herself and, in interaction with various barriers, may hinder the person’s full and effective participation in society on an equal basis with others. Disabilities may be permanent or temporary, but these guidelines refer specifically to those that are permanent or affect a person in the long term.

**Hazard**
A condition that threatens the safety and well-being of people. In origin it may be natural (e.g. earthquakes, floods), technological (e.g. transportation crashes, toxic releases), social (e.g. crowd crushes, demonstrations) or intentional (e.g. terrorism, politically-inspired violence).

**Vulnerability** (in the present context)
A person’s susceptibility to harm as a result of external adverse events such as natural disasters, public emergencies, technological incidents or political violence.

**Risk** (in the present context)
The product of hazard and vulnerability leading to a probability of harm, expressed as physical or psychological injury, damage, destruction or interruption of productive and essential activities.
**Crisis** (in the present context)
A major incident that interrupts normal activities for a significant number of people and causes, or threatens to cause, harm to themselves or their property.

**Disaster**
A major destructive event that involves a large number of people and causes widespread damage and probably significant physical injury, possibly with a number, more or less large, of fatalities. Attempts to quantify the threshold at which an incident becomes a disaster have not generally been successful, but disasters have a profound effect on society and communities and tend to be larger, or more serious, than incidents.

**Disaster risk reduction**
The process of preparing for, reducing the risk of and planning to face disaster when it happens.

**Resilience**
For individuals, groups of people and society as a whole, the state (and process) of being robust in the face of disaster risks. This means being able to reduce the impact of disaster, manage its effects with efficiency and recover rapidly from it, hopefully to a state of greater resistance than existed before (the “bounce forward” strategy).

**Civil protection** (sometimes known as civil defence or civil security)
The provision of services to the general population that enable them to face the risk of, survive and reduce the damaging effects of disasters and crises.
Created in 1987, the European and Mediterranean Major Hazards Agreement (EUR-OPA) is a platform for co-operation between European and Southern Mediterranean countries in the field of major natural and technological disasters. Its field of action covers the knowledge of hazards, risk prevention, risk management, post-crisis analysis and rehabilitation. It has up to date 26 member States.

www.coe.int/europarisks

The Council of Europe is the continent’s leading human rights organisation. It includes 47 member states, 28 of which are members of the European Union. All Council of Europe member states have signed up to the European Convention on Human Rights, a treaty designed to protect human rights, democracy and the rule of law. The European Court of Human Rights oversees the implementation of the Convention in the member states.

www.coe.int