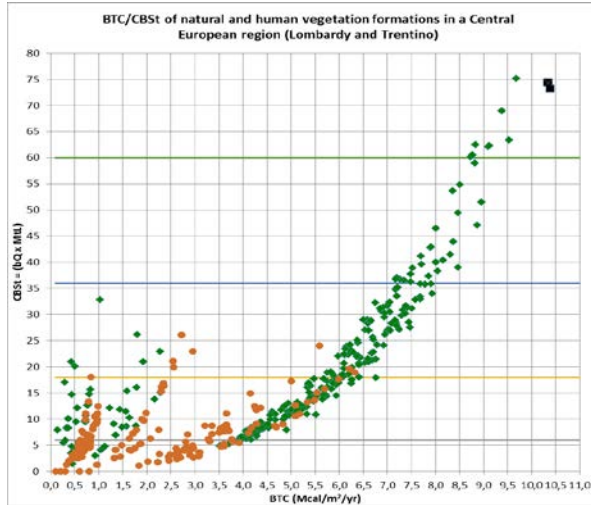








The previous two indexes allows to express systemic estimations *both of natural and human* vegetation types (Fig. 2) and, consequently, to better understand the concrete role played by the examined vegetation community on respect to other basilar landscape parameters, e.g. human health.



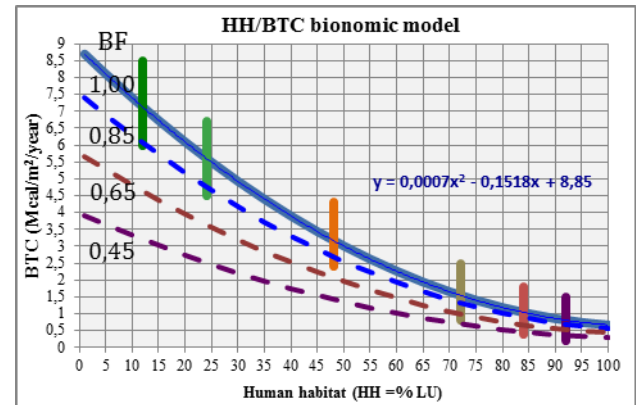
**Fig. 2.** Correlation between the BTC and the evaluation of its CBSt, from 430 samples of natural (green) and human (red dots) vegetation, from Lombardy and Trentino (Northern Italy), compared to two case studies from Bialowieza Forest (Poland). Vegetation communities with lower resistance strategy (hence lower BTC) reach a lower bionomic state and show less efficiency.

Focusing on the human component of the landscape, the basilar function is the HH, human habitat, which gives us a quantitative measure of the areas permanently managed by human population, adding subsidiary energy, so limiting the self-regulation capacity of natural systems (NH): thus, HH goes beyond a simple survey of the urbanised surface.

After the study of 45 landscape units (mainly in North Italy) an exceptional correlation between BTC and HH was found, with an  $R^2 = 0.95$  and a Pearson's correlation coefficient of 0.91 (about three time the minimum value of significance). From this experiment, it was possible to build the simplest mathematical model of bionomic normality (Fig. 3), available for a first framing even of the dysfunctions of landscape units (LU). Below a tolerance interval (0.15 from the curve of normality) we can register three levels of altered *bionomic functionality* (BF): normal BF (1.15-0.85), altered BF (0.85-0.65), dysfunctional BF (0.65-0.45) and degraded BF ( $< 0.45$ ).

Following bionomic principles the carrying capacity ( $\sigma$ ) of a LU, too, can be measured, as the ratio between the standard habitat per human capita

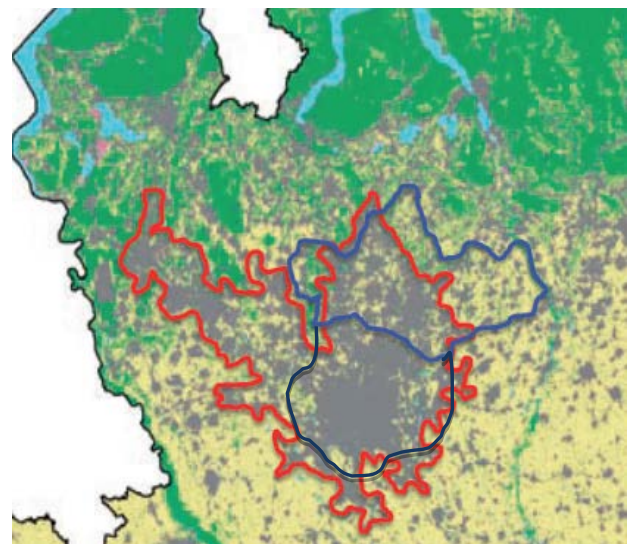
(SH) and the theoretical minimum SH\*. This SH index, deduced from nutritional and ecological parameters [6, 7] (available also for the fauna component, with proper values), gives also the level of heterotrophy of a landscape unit, whose minimum may be considered  $SH = 0.75-0.80$ .



**Fig. 3.** The bionomics model HH/BTC (Human Habitat-Biological Territorial Capacity of vegetation) in which the BF curves (dotted lines) indicate the functionality of the examined LU.

### 2.3 The frame data for investigation

The choice of a land area of experimentation presenting an available gradient of 5-6 types of landscape (from dense urban to agricultural-forested) was facilitated by living in Milan, one of the largest metropolitan areas of Europe (close to 5 million inhabitants).



**Fig. 4.** The blue-grey lines indicate the land area of experimentation, consisting of Monza-Brianza province (North) and the city of Milan (South), within the great hinterland of the town of Milan (red line). This territory cover 655.9 Km<sup>2</sup> with a population of  $2.5 \times 10^6$  inhabitants and with a gradient of at least 5-6 landscape types. [26].

As plotted on Fig. 4, the blue-grey lines indicate a territory covered by the province of Monza-Brianza (North) and a portion of the province of Milan (South): in summary 655.8 Km<sup>2</sup>, 2.5 x 10<sup>6</sup> inhabitants with a gradient of at least 5-6 landscape types (Tab. 1).

Tab. 1. Six landscape types and their bionomic characters in the study area					
Landscape Type	HH	BTC	MR	PA	BF
Agricultural	72,00	1,68	7,11	42,38	1,08
Rural	78,16	1,25	8,33	42,66	0,98
Suburban-Rural	83,62	0,89	7,62	42,57	0,84
Suburban-Tech.	88,41	0,61	8,66	43,47	0,67
Urban	92,17	0,44	9,45	44,75	0,54
Dense urban	93,65	0,39	9,57	45,08	0,41
Mean	84,67	0,88	8,46	43,49	0,78

MR = mortality rate; PA = Population age; BF = bionomics Functionality

Note (Fig. 5) that not only Milan, but also Monza and Brianza are contained within one of the most air-polluted area of Europe [27], therefore pollution could be considered as homogeneous in our sample land area.

We have to note that the city of Milan was analysed by being divided into its 9 administrative sections, to have data comparable with the other municipalities, both in surface and population. Remembering the carrying capacity  $\sigma = SH/SH^*$ , few municipalities reach its minimum value in our territory, thus being heterotrophic.

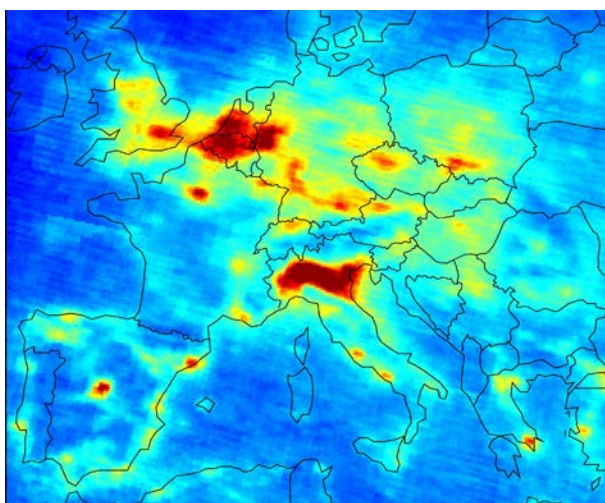


Fig.5. In the Po plain the distribution of air pollution is quite homogeneous and one of the most high in EU. Not only Milan but also Monza-Brianza are inserted in this wide polluted area (from E.S.A [27])

The biological territorial capacity of vegetation (BTC) was estimated both with field surveys and the registered statistical data on the main types of vegetation. The evaluation of concise bionomic state of vegetation (CBSt) better allows the assessment of the forest vegetation in the entire Monza-Brianza Province. vegetation". Fig. 6 exposes the most significant set of forest assessment surveyed on the field. The modest value of the mean territorial capacity, BTC = 5.84 Mcal/m<sup>2</sup>/year, is confirmed by the presence of 57.14 % of altered and weak forests, Vs only 19.05 % of good ones.

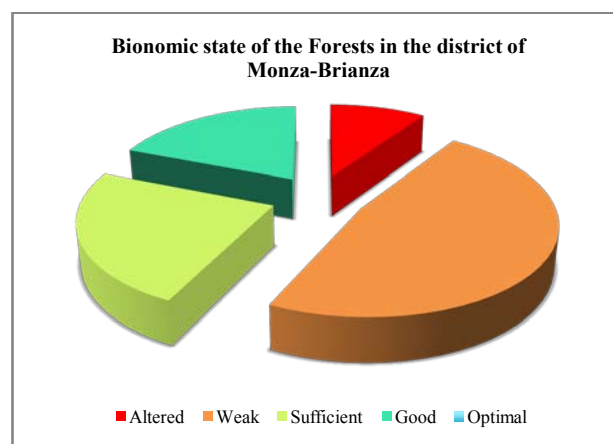


Fig. 6. The bionomic state of the forest formation on the Province of Monza-Brianza shows only 19.05 % of good conditions and no one in optimal one.

Considering our 72 LU, these range of values were registered: human habitat HH = 66-97 (% of LU); biological territorial capacity of vegetation BTC = 2,01-0.24 (Mcal/m<sup>2</sup>/year); mortality rate MR = 4.10-11.35 (x 1000 inhabitants); population age PA = 38.6-47.1 (years); level of alteration or degradation of functional bionomic state BF = 1.10-0.34.

Only 30.56% municipalities (22/72) were found in a normal condition (BF ranging between 1.15-0.85), while 34.72% of them (25/72) were in dysfunctional or degraded condition, the other being altered. These levels of BF were obtained comparing the mean BTC of each LU with the value given by the BTC model (see Fig. 3) corresponding to the LU value of HH.

### 3.Results

Plotting the results of the previous Tab.1 related to the unit, we reach a first evidence of the increase of mortality rate (MR) when the bionomic functionality (BF) is decreasing from normality (BF = 1.0)

Traditional ecology states that the increase of the mean age of a population (PA) indicates a very developed country and also less negative environmental factors. On the contrary, in figure 7 we see that the increase of population age is linked to an increase of landscape degradation: causes and effects need further investigations. However, the contrast between traditional and bionomic interpretation is not so strong as it may appear. We may observe that PA is correlated with the landscape dysfunction in a way very similar to MR (Fig. 7).

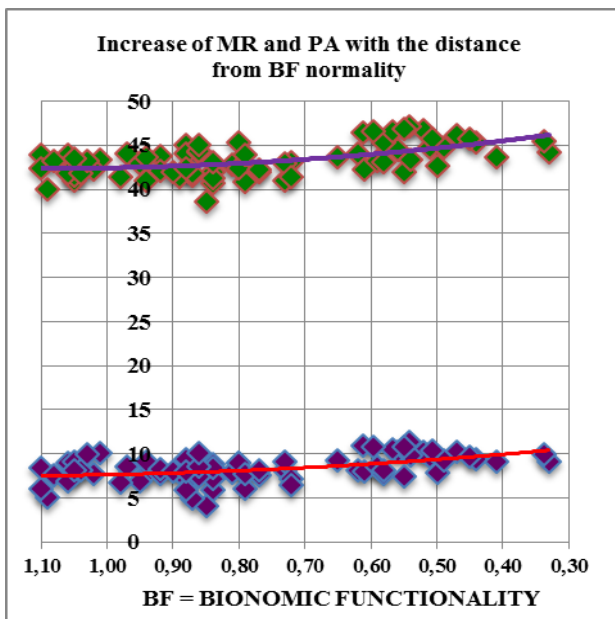


Fig. 7. MR (mortality rate, violet and red) and PA (population age, green) plotted in function of BF.

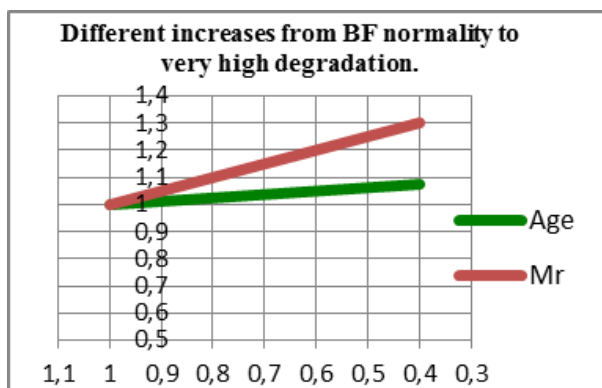


Fig. 8. Different increases of MR and PA related to the decrease of BF (from 1 to 0,4)

Note that the increase of  $PA = f(BF)$  cannot be read as the reason for the parallel increase of MR. If we examine our two comparing functions,  $MR = f(BF)$  and  $PA = f(BF)$  (Fig. 8), we note a sharp difference: e.g. going from the point of full normality ( $BF = 1,0$ ) to a point of very high

degradation ( $BF = 0,4$ ), PA increases only 7.3 %, while the MR increases 30.3 %. This signifies that merely 24 % of the MR increase can be attributed to the PA, therefore 76 % being due to other causes as, remember, all the area presents the same level of pollution. Being very contained, the increase of PA seems to imply limited negative environmental factors.

Detailing the 72 LU, the correlation MR/BF (mortality rate/bionomic functionality) emerges (Fig. 9). The result of  $MR = f(BF)$  presents the  $R^2 = 0,252$ , but the Pearson's correlation coefficient is (-0,438) that is twice the minimum value of significance: therefore MR increases with BF diminishing. At full normality BF is pair to 1.00 and  $MR = 7.64$ , becoming  $MR = 7.95$  at  $BF = 0,85$  but  $MR = 10.27$  at  $BF = 0.35$ , enlightening deep dysfunction.

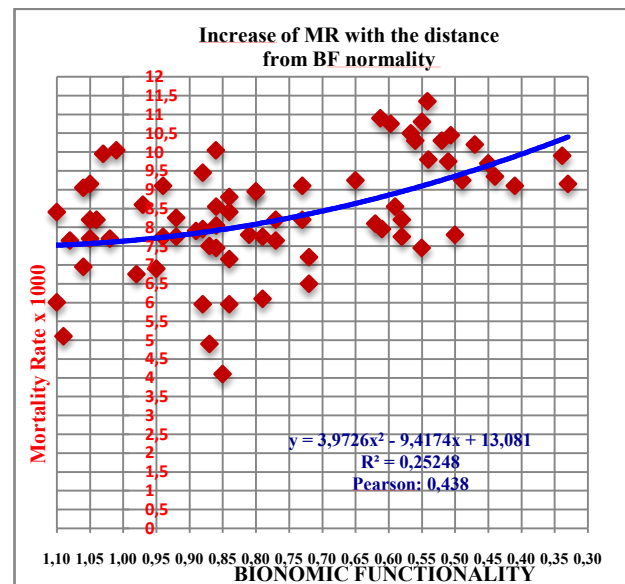


Fig. 9. Correlation between Mortality Rate and Bionomic dysfunction in 72 Landscape Units of the Milano-Monza (MI-MB) Area. The Pearson coefficient is twice the minimum values of significance.

A first base to estimate a R risk Factor can be deduced from Tab. 1: the mean BF is 0.78, therefore

$$\Delta MR_{BF} = (MR_{BF} - MR_{BF=1}) \times 76\% = (8.34 - 7.64) \times 0.76 = 0.532 \times 10^{-3}$$

Consequently, the estimation of premature death (PD) probability due to BF ( $PD_{BF}$ ) can be measured remembering that the total population in the study area is 2.524 millions:



$$PD_{BF} = (2,524 \times 10^6) \times (0.532 \times 10^{-3}) = 1,342.7/\text{year}$$

This huge number of premature death/year can be compared with the total death number (TD) in this area, that is  $TD = 21,050$  people/year. The rate

$$PD_{BF}/TD = 6.4\%$$

This high Mortality Rate results similar to the fine dust premature death ( $PD_{FD}$ ) = 7.0 % related on the Milan Metropolitan Area.

## 4. Interpretation

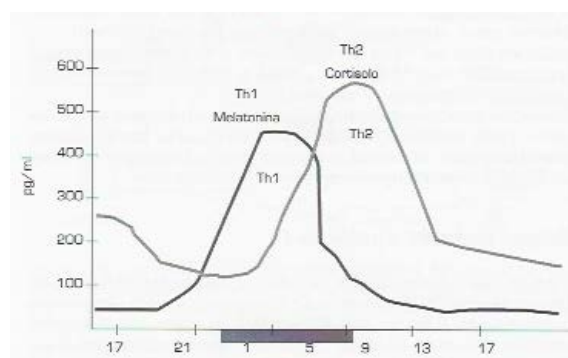
### 4.1 Theory explaining the results

First of all, there is a basilar ethological question, linked to information, meaning and survival, therefore to the concept of “values-judgment” [29]. This implies a correspondence between the perception of values and the passage from the most probable and disordered to the most ordered and improbable. The scale of values derived from this fundamental process can be applied to judge the health of an organism or of a landscape. This ethological process has an alarming meaning [7]: so it registers all the environmental alterations, which remains memorized within our cells. To reach this goal the ethological process have to be innate and unconscious, because to de-codify environmental signals we must reduce emotions. Remember that subconscious mind is able to process  $20 \times 10^6$  environmental stimuli/sec, Vs. only 40 stimuli/sec. of the conscious mind.

The ethological information on the state of our environment can be elaborated following the principles of psychoneuroendocrinimmunology [15, 30]. For instance, an emotional or alarm signal registered in dACC is correlated with an increase of plasmatic levels of a basilar inflammatory cytokine (type II of TNF). So, a direct relation between psychic stress and inflammation is demonstrated [31]. Moreover, the effects of hormones on immune system depend mainly on nervous systems and the hemato-encephalic barrier is today considered as a mistake.

Berne and Levy within their book “Principles of Physiology” [32] section VIII “Endocrine System” (written by Saul. M. Genuth) [33], discuss the integrated responses of the body to stress. The sympathetic nervous system and the hypothalamus-pituitary-adrenal axis mediate the integrated responses of the human organism to stress. The main factors that cause stress (stressors) simultaneously activates neurons in the hypothalamus, which secrete CRH and adrenergic

neurons [34].



**Fig. 10.** The complementary circadian rhythm of melatonin and cortisol is strongly influenced by chronicle stress. Note the relations with the type Th1 and Th2 of immune circuit [35].

These responses potentiate each other, both at the central and the peripheral level. The final effect of the activation of neurons that secrete CRH is the increase in plasma levels of cortisol, while the net effect of adrenergic stimulation is to increase plasma levels of catecholamine. The negative feedback exerted by 17-hydroxycorticosterone (cortisol) has the function to limit an excessive reaction, which is dangerous for the organism. But when the stress became chronicle the circadian rhythm melatonin/cortisol (Fig. 10) is altered. Consequently, plasma levels of cortisol brings to a dominance of the Th2 immune circuit, which a production of typical catecholamine (e.g. IL-4, IL-5, IL-13) and of the circuit Th17 [18].

Remember the Th2 immune response is not available to counteract viral infections, neo-plastic cells, auto-immune syndromes, which should need a Th1 response. We can deduce that the majority of illness is linked with chronicle stress conditions, confirming the intuitions of Selye [15]. A strong increase of morbidity inevitably brings to an increase of mortality rate (MR).

In his book “Landscape ecology, a widening foundation” [7] within section 5.5 “Landscape pathology and human health”, Ingegnoli noted that the abovementioned stressors can be frequently due to landscape degradations. In facts, psychic disorders, aggressiveness and high morbidity are evident in many degraded urban peripheries [36] or in many rural areas altered by monoculture, destruction of trees and hedgerows, highways, industrial buildings. In the case of landscape disorders due to structure degradation and the alteration of normal bionomic processes (even in the absence of pollution), the stressors tend to be chronic and the physiologic mediation of cortisol

may become insufficient. Therefore, landscape dysfunctions, even in absence of pollution, may attempt our health bringing to an excess of cortisol, which reduces our hormonal, immune and nervous system defences. The increase of the most common syndromes depend also on this process: first of all cardiovascular diseases, cancer, but even minor diseases such as allergies.

Today we may complete this theoretical interpretation of the increase of illness, therefore of MR, in dysfunctional landscapes. Landscape bionomics allows to assess and measure the level of environmental dysfunction, as exposed in the second paragraph (method).

Theoretically, even in absence of pollutants the environmental stress can be alarming, because an excess of cortisol can reduce our hormonal, immune, nervous and mental system defences. Anyway, it assumes a wider importance in case of high pollution, when our defence systems have to fight strongly to avoid the effects of toxins and degenerative diseases. Moreover, the increase of population age (PA) implies a decrease in birth rate, that traditional ecology asserts to be caused by social and economic reasons. But landscape bionomics observes that birth decline is due also by environmental degradation which may inhibit sexual fertility and reproduction [32].

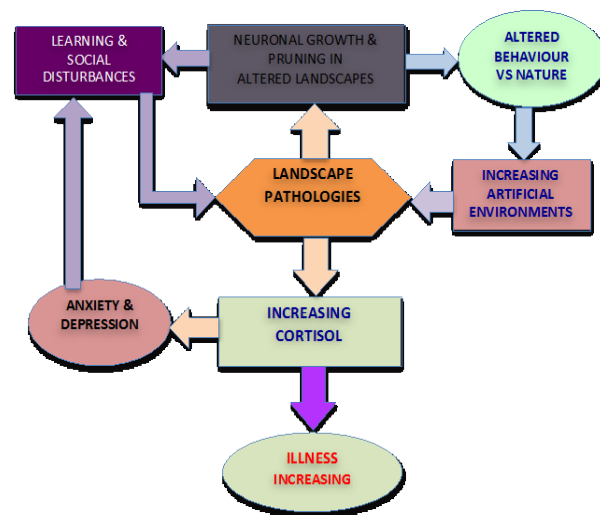
## 4.2 Main consequences of Landscape Pathologies on Human Health

As expressed in the introduction, the environment influences our health through exposures to physical, chemical and biological risk factors, and through related changes in our behaviour in response to those factors. So, we must note that in case of landscape pathologies not only insidious stressors and consequent illness compare, but also an altered behaviour which in turn worsens the conditions of the alteration of the landscape, initiating a dangerous exalting feedback.

Today we know that, during adolescence, different parts of the brain mature at different rates. Neural connections increase after the birth to about 5 years. An average decrease in “grey matter” volumes between ages 5 and 20, thanks to the “pruning” of neural connections (for best environmental fitness) has been demonstrated. Areas that mediate “executive functioning” mature later than areas responsible for basic functions [37]. Therefore, artificial environments seems to attract more and more young people and to submit urban green to technology as sign of a New Babylon.

Let us observe that the two processes we

exposed before, the ‘growth & pruning neural synapsis’ and the ‘environmental stress alarm’, have to be obviously integrated in relation to landscape pathologies and human health. In figure 10 a logic flow diagram tries to explain their main linkages and loops.



*Fig. 11. Logic flow diagram on the main consequences of landscape pathologies on human health. Note the formation of three feedback loops, which can aggravate the landscape syndromes, therefore the consequences on illness and MR.*

If neuronal growth and pruning process operates in altered landscapes, all this can results in an altered behaviour Vs. Nature, bringing to increase artificial environments, than intensifying landscape pathologies: a typical positive feedback loop. But another behavioural feedback loop appears when an increase in cortisol brings to increase arousal and aggression; then learning and social disturbances lead to act against natural lows, thus degrading ecological components and producing a new intensification of landscape pathologies. Chronically elevated cortisol levels have been linked to problems including abdominal fat gain, cognitive decline and compromised immune function. Note that learning and social disorders are reinforced by the altered neuronal growth & pruning.

## 4.3 Importance of applications

In presence of pollution, drastic changes of impact assessment derive from the found correlation MR/BF (Fig. 12). When pollution is strong, the BF risk factor enhances the effects on premature death. The case study of Vado Ligure carbon-powered plant of electric energy can be of great significance. Lung cancer mortality (x 100.000 inhabitants)



results 54.6 in Italy, 83.5 in Savona Province, but 112.3 in Vado Ligure. This municipality is not only polluted, it's heavily altered by landscape syndromes (Fig. 12), from industrial and harbour areas to forest and agricultural degradations. Therefore, after the cited theory, the so high MR is due not only by chemical pollutants, but also from BF alterations, when our defence systems have to fight strongly to avoid the effects of toxins and degenerative diseases.



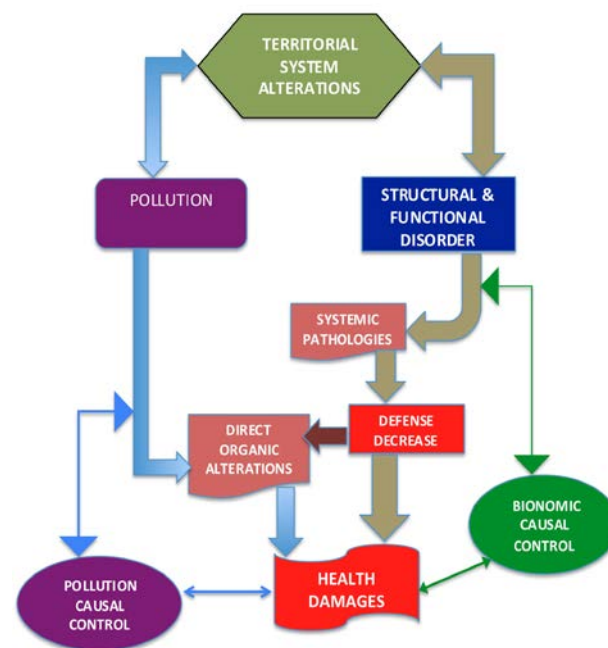
**Fig. 11.** The municipality of Vado Ligure (Liguria, Italy), with the carbon-powered electric plant.

This fact must be added to the effects of endocrine disruption chemicals (EDC) in humans. It is well known [38] that, for a clinician taking care of an individual patient, numerous challenges in ascertaining EDC involvement in a particular disorder are possible:

- each person has unique exposure to a variety of both known and unknown EDCs. Considerable variability in the half-life and persistence of EDCs, as well as their degradation in body fluids and tissues, are created by individual differences in metabolism and body composition;
- Susceptibility to EDCs may vary according to genetic polymorphisms. In addition, human disorders are more likely the result of chronic exposure to low amounts of mixtures of EDCs. The latency between exposure to EDCs and occurrence of clinical disorders creates further challenges when one attempts to establish a relationship at the level of a given individual.

Anyway, even if the International Association for Impact Assessment (IAIA) correctly indicates 'cumulative impacts' as the most important health/environment studies [39], today the procedure to evaluate a case study like Vado Ligure is the scheme limited to the left of Fig. 12. The difference with the right part of this figure is sharp. The effects of landscape pathologies on human health bring not only to an integration among

different types of EDC and geo-physics parameters, but even among the biologic-structural parameters of the landscape (see the flow chart differences!). Thus, the bionomic indexes and models become very important also in bio-monitoring and we need to refer to a new professional figure, the ecologist-physician or "ecoiatra", able to face the diagnostic evaluation of the landscape.



**Fig. 12.** Logic diagram of the screening for a case study like Vado Ligure. Note that the health damages go beyond the direct organic alterations.

## 5. Conclusion

To understand better the relation between the environmental dysfunctions and human health, the new discipline of Landscape Bionomics confirms to be indispensable. It allows the formation of an "ecoiatra" as the physician of the environment. Also a deeper exchange between Medicine and Ecology-Bionomics should help these researches, as proved after the Seminar on this argument held by Ingegnoli (2016, May) at the Sapienza University of Rome, Department of Experimental Medicine.

The Author is in touch with IAIA (International Association for Impact Assessment) for the same argument. In facts, the importance of environmental protection and rehabilitation may be reinforced by the linkage health-environment, while in too many cases people follow banal urban-gardening solutions. It must be known that we cannot arrive to protect even the human health if we do not follow a

true scientific approach on environmental rehabilitation [40].

Evident limits are related to the dominant reductionism in scientific researches. For instance, in the evolution processes, an overvaluation of the genetic determinism comes from the underestimation of the environment. Therefore, a new approach to the environment is needed. The discipline of Ecology or “*Speech on our House*” is necessary, but not sufficient: Bionomics or the “*Doctrine of the Laws of Life organisation on the Earth*”, even outside the organism, is needed too; the landscape has to be recognised and investigated as a peculiar biological level. This new perspective inevitably leads to significant changes in how to assess and manage the environment.

We hope that a study like the one exposed in this paper should be repeated in other countries both in Europe and in USA, to arrive to a better definition of risk factor on the estimation of premature death (PD) probability due to BF (PD<sub>BF</sub>).

## 6. References

- [1] W.H.O. (2011) *Public Health and Environment, Global Strategy Overview*. World Health Organisation, Geneva
- [2] W.H.O. (2014) *Seven million premature deaths annually linked to air pollution*. World Health Organisation, Geneva
- [3] Meffe GK, Carroll CR (1997) *Principles of conservation biology*. Sinauer Associates Inc Publ, Sunderland, Massachusetts
- [4] Ingegnoli V. (1991) *Human influences in landscape change: thresholds of metastability*. in: O Ravera (ed.) *Terrestrial and aquatic ecosystems: perturbation and recovery*. Ellis Horwood Ltd., Chichester, England pp. 303-309
- [5] Ingegnoli V (2001) *Landscape Ecology*. in: D Baltimore, R Dulbecco, F Jacob, R Levi-Montalcini (eds.) *Frontiers of Life*. vol IV. Academic Press, Boston, pp. 489-508.
- [6] Ingegnoli V (2015) *Landscape Bionomics. Biological-Integrated Landscape Ecology*. Springer-Verlag, Heidelberg, Milan, New York, pp. XXIV+431
- [7] Ingegnoli V (2002) *Landscape Ecology: A Widening Foundation*. Springer-Verlag, Berlin, Heidelberg, New York, pp. XXIII+357
- [8] Ingegnoli V, Giglio E (2005) *Ecologia del paesaggio: manuale per conservare, gestire e pianificare l'ambiente*. Simone Edizioni-Esse Libri, Napoli, pp. 685+XVI
- [9] Maas J, Verhij RA, Groenewegen PP, de Vries S, Spreeuwenberg P (2006) *Evidence based public health policy and practice. Green space, urbanity and health: how strong is the relation?* J Epidemiol Community Health. Vol. 60, pp. 587-592. doi:10.1136/jech.2005.043125
- [10] Mitchell R, Popham F (2008) *Effect of exposure to natural environment on health inequalities: an observational population study*. Lancet, Vol. 372, pp1655-1660 (Issue 9650)
- [11] Nieuwenhuijsen MJ, Kruize H, Gidlow C et al. (2014) *Positive health effects of the natural outdoor environment in typical populations in different regions in Europe (PHENOTYPE): a study programme protocol*. BMJ Open. Vol. 4, e004951 doi:10.1136/bmjopen-2014-004951
- [12] Lorenzini G, Nali C (2005) *Le piante e l'inquinamento dell'aria*. Springer-Verlag, Milano
- [13] Bottaccioli F (2014) *Epigenetica e Psiconeuroendocrino-immunologia; le due facce della Rivoluzione in corso nelle scienze della vita*. Edra Spa, Milano, pp.192
- [14] Selye H (1956) *The Stress of Life*. Mc Graw-Hill, New York
- [15] Altman J. (1963) *Autoradiographic investigation of the cell proliferation in the brains of rats and cats*. Anathomy Research 145:573-591
- [16] Holliday R (2006) *Epigenetics: a historical overview*. Epigenetics, Vol. 1, pp. 76-80
- [17] Crick F (1970) *Central Dogma of Molecular Biology*. Nature, Vol. 227, pp. 561-563
- [18] Ingegnoli V (2011) *Bionomia del paesaggio. L'ecologia del paesaggio biologico-integrata per la formazione di un "medico" dei sistemi ecologici*. Springer-Verlag. Milano, pp. XX+347
- [19] Naveh Z, Lieberman A (1984) *Landscape Ecology: theory and application*. Springer-Verlag New York, Inc, pp.
- [20] Naveh Z, Lieberman A (1994) *Landscape Ecology: theory and application*. Springer-Verlag New York, Inc, pp.
- [21] Pignatti S, Box EO, Fujiwara K (2002) *A new paradigm for the XXIth Century*. Annali di Botanica, Vol. II, pp. 31-58
- [22] Ingegnoli V (2005) *An innovative contribution of landscape ecology to vegetation science*. Israel Journal of Plant Sciences. Vol. 53, pp. 155-166
- [23] Ingegnoli V, Pignatti S (2007) *The impact of the widened Landscape Ecology on Vegetation Science: towards the new paradigm*. Springer Link: Rendiconti Lincei Scienze Fisiche e Naturali. s.IX, vol.XVIII, pp. 89-122
- [24] Ingegnoli V (1999) *Definition and Evaluation of the BTC (Biological Territorial Capacity) as an Indicator for Landscape Ecological Studies on Vegetation*. in: W Windhorst, PH Enckell (Eds) *Sustainable Landuse Management: The Challenge of Ecosystem Protection*. EcoSys: Beitrage zur Oekosystemforschung, Suppl Bd 28, pp. 109-118

- [25] Ingegnoli V, Giglio E (1999) *Proposal of a synthetic indicator to control ecological dynamics at an ecological mosaic scale*. *Annali di Botanica*. Vol. LVII, pp. 181-190
- [26] DUSAF (2010) *Usa del suolo in Regione Lombardia*. Regione Lombardia, Milano
- [27] European Space Agency (2004) *Annual Report*. Paris
- [28] Ingegnoli V (2013) *Concise evaluation of the bionomic state of natural and human vegetation elements in a landscape*. *Rend. Fis. Acc. Lincei*; doi 10.1007/s12210-013-0252-2
- [29] Lorenz K. *Vergleichende Verhaltensforschung: Grundlagen der Ethologie*. Springer-Verlag, Berlin, Wien; 1978
- [30] Hiramoto RN, Solvason HB, Hsueh CM (1999) *Psychoneuro-endocrine immunology: perception of stress can alter body temperature and natural killer cell activity*. *Int. J. Neurosci*. Vol. 98 (1-2): 95-129
- [31] Slavich GM, Way BM, Eisenberger NI, Taylor SE (2010) *Neural sensitivity to social rejection is associated with inflammatory responses to social stress*. *PNA*, Vol. 107 (33), pp. 14817-14822 doi: 10.1073/pnas.1009164107
- [32] Berne RM, Levy MN (1990) *Principles of Physiology*. The CV Mosby Company. U.S.A
- [33] Genuth SM (1990) *The Adrenal Medulla*. in: RM Berne, MN Levy. *Principles of Physiology*. The CV Mosby Company, USA, pp. 572- 578
- [34] Hansel A, Hong S, Càmara RJ, von Kanel R (2010) *Inflammation as a psychophysiological biomarker in chronic psychological stress*. *Neuroscience and Biobehavioral Reviews*. Vol. 35, pp. 115-121
- [35] Bottaccioli F (2012) *Stress e Vita. La scienza dello stress e la scienza della salute alla luce della Psiconeuroendocrinoimmunologia*. Tecniche Nuove, Milano
- [36] Melse JM, de Hollander AEM (2001) *Human Health and the Environment. Background document for the OECD*. Environmental Outlook, OECD, Paris
- [37] Gogtay N, Giedd JN, Lusk L et al (2009) *Dynamic mapping of human cortical development during childhood through early adulthood*. *Proceedings of the National Academy of Sciences of the United States of America*. Vol. 101 (21), pp. 8174-8179
- [38] Diamanti-Kandarakis E, Bourguignon JP, Giudice LC et al.(2009) *Endocrine-disrupting chemicals: an Endocrine Society Scientific Statement*. *Endocr Rev*. 2009; Vol. 30 (4), pp. 293-342 doi: 10.1210/en.2009-0002.
- [39] Magro G. et al. (2007) *Quantifying Cumulative Impact Assessment with Dynamic Computational G.I.S. System in a multi stressor area: a case study on a waste treatment plant in Italy*. Annual Conference “Growth, Conservation and Responsibility”, IAIA Seoul, Corea
- [40] Ingegnoli V, Giglio E (2016) *Landscape project can limit bionomics dysfunction risk factor Vs. premature death increase*. *Modern Environmental Science and Engineering*, vol. 2 ( 7): 435-444. Academic Star Publishing Company, USA; doi 10.15341/mese(2333-2581)/07.02.2016/001
- [41] Szif M, McGowan P, Meaney M (2008) *The social environment and epigenome*. *Environmental and Mol Mutagenesis*, vol. 49, pp. 46-60.