

International Association for Vegetation Science (IAVS)

∂ FORUM PAPER

The concept of vegetation class and order in phytosociological syntaxonomy

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Academic editor: Wolfgang Willner + Received 23 October 2020 + Accepted 27 November 2020 + Published 21 December 2020

Abstract

In order to stabilize the defining concepts of the higher rank syntaxonomic units such as class or order, the criteria of floristic content and unity of origin are enunciated. This is done with the aim of preventing the fragmentation of the large classes and the subsequent typological inflation. For orders, the criterion of specific floristic content is discussed, with orders that have been described to encompass seral secondary forests or the separation of forest vegetation from that which is dominated by shrubs rejected, due to their weak floristic characterization. These criteria have been applied to two forest vegetation classes: the European temperate (*Querco-Fagetea*) and the Mediterranean (*Quercetea ilicis*). For the first, it is argued in favor of maintaining a single class for all temperate deciduous forests in Europe instead of dividing them into four. Within this single class five orders are distinguished: *Fagetalia*, *Quercetalia roboris*, *Quercetalia pubescenti-petraeae*, *Alno-Fraxinetalia* and *Populetalia albae*, rejecting the orders that have been proposed for secondary forests because they have few characteristic taxa. For the sclerophyllous and macchia forests of Mediterranean Europe, the *Quercetea ilicis* class can be split into two or three geographical orders, rejecting the *Pistacio-Rhamnetalia alaterni* as a shrubby physiognomic unit.

Taxonomic reference: Castroviejo S (coord. gen.) (1986–2012) Flora iberica 1–8, 10–15, 17–18, 21. Real Jardín Botánico, CSIC, Madrid, ES.

Syntaxonomic reference: Mucina et al. (2016).

Keywords

Floristic content, Quercetea ilicis, Querco-Fagetea, syntaxonomy, unity of origin, vegetation class, vegetation order

Introduction

The concept of vegetation class needs to be formalized in such a way that instability is prevented. The higher rank syntaxonomic units, such as classes, as far as they represent the fundamental parts of the ecological and floristic variability in a given territory, should have a well-defined ecology and a sufficient biological content (Pignatti et al. 1995). In terms of phytosociological syntaxonomy, each class should have a certain number of characteristic taxa; a minimal "floristic content". As syntaxonomy reflects the floristic and ecologic relationships between the subordinate units, the higher rank units should include a significant content of a specific flora. This is particularly important in the case of classes, but it is also valid for orders. The number of taxa required to fulfill these minimal requirements are variable and must be estimated in the context of the vegetation types involved. Species rich vegetation, such as the *Querco-Fagetea* or *Molinio-Arrhenatheretea*, would need a longer list of characteristic taxa to justify the classes and orders, while species poor vegetation, such as the *Calluno-Ulicetea* or *Cytisetea scopario-striati*, have fewer taxa available for this purpose and it is usually difficult to distinguish more than one or two orders within a class in such cases.



Copyright Javier Loidi. This is an open access article distributed under the terms of the Creative Commons Attribution License (CC-BY 4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited. The syntaxonomic scheme structuring a class is the expression of the ecological and dynamic diversity occurring within it, but, to a certain extent, it can also express a certain biogeographical-evolutionary background. The floristic set characterizing a class has potentially originated in a particular evolutionary episode that occurred in a particular geographical area under specific environmental conditions. We could hypothesize a type of "unity of origin" in the floristic element characteristic of the class. This eco-evolutionary background should prevent us from making arbitrary decisions which are not based on real patterns in nature.

I will illustrate these ideas with a short discussion on the classes *Querco-Fagetea* and *Quercetea ilicis*.

The class Querco-Fagetea

The recent publication of the EuroVeg checklist (Mucina et al. 2016) has revived the question of the syntaxonomic structuring of temperate deciduous forests in Europe. Contrary to the concept of a broad class Querco-Fagetea sylvaticae, maintained by several European phytosociologists (Mucina et al. 1993, Stortelder et al. 1999), Mucina et al. (2016) decided to accept the division of this syntaxon into four European forest classes: Quercetea robori-petraeae, Carpino-Fagetea sylvaticae, Quercetea pubescentis and Alno glutinosae-Populetea albae. A different approach was presented by Rivas-Martínez et al. (2011), who proposed one class for non-riparian forests and another one for riparian forests. Other approaches have included all the fen alder and riparian ash forests in a specific class, the Alnetea glutinosae (Berg et al. 2004). A solution to this controversy requires the application of clear and objective criteria. Here, I advocate the recognition of a single class that includes most of the deciduous forests of Europe. This is supported by three broad arguments that are discussed:

1. Biogeographical-evolutionary criterion: common origin and evolution. The current European deciduous forests descend from the deciduous forests that occupied a wide area in the extratropical latitudes of the Northern Hemisphere in the first half of the Tertiary. They encompassed the territories of ancient North America and those of Eurasia. In that time they were connected by the proximity of the land masses that formed Laurasia (Axelrod 1983). The separation of North America and Eurasia and the subsequent climatic changes that occurred in the late Tertiary, gave rise to the fragmentation of that continuous zone into the three current ones: (i) North America (basically the eastern half of the USA and the southeast of Canada), (ii) Europe (to which are added areas of western Asia bordering the Black and Caspian seas), and (iii) East Asia (Sino-Japanese region comprising northeast China, the Korean peninsula, the south of the Russian Far East and the Japanese archipelago). This general context shows that the European temperate deciduous forests have a common origin followed by a common evolutionary history up to the present. Since the phytosociological class is the highest unit of the system, this unity of origin should

be reflected in a single class *Querco-Fagetea sylvaticae*. The diversity existing within it can be represented by lower syntaxonomic ranks, such as the order and the alliance.

2. Floristic criterion: common set of characteristic species. After being isolated from the American and East Asian deciduous forests, the European ones shared the climatic oscillations of the Pleistocene. It is generally assumed, since Gray (1878) and Reid (1935), that the European forest flora was impoverished as a result of the extinctions that occurred during the Pleistocene glaciations, whose effects were intensified by the east-west disposition of the mountain ranges. Later it was postulated that these extinctions must have been more severe in the siliceous flora than in the basiphilous one, because the siliceous territories, located mainly in the north and in the center of Europe, suffered more from the climatic rigors of the glaciations than those in the south of the continent where base-rich rocks are more abundant, allowing the basiphilous flora to find more refuge areas and suffer fewer extinctions (Chytrý et al. 2003). In spite of these differences, there is a set of specific forest plants occurring throughout the European temperate deciduous forests, regardless of the substrates on which they are found, from northwestern Iberia to the Hyrcanian region south of the Caspian sea. No less than 25 forest species, from herbs such as Athyrium filix-femina, Stellaria holostea, Poa nemoralis or Brachypodium sylvaticum, to shrubs such as Rubus caesius, Daphne laureola or Frangula alnus and trees such as Acer campestre, Fraxinus excelsior, Quercus robur, Taxus baccata or Ulmus glabra, are common to most regions where these forests exist.

3. Application criterion: coherence in the presentation. Syntaxonomy has to be presented to non-phytosociologists in a useful and coherent way. It is very difficult to argue, in a teaching context, that forests dominated by the same species belong to different classes. This is also valid in other fields such as conservation, vegetation mapping or land management and planning. If we recognize four separate classes, there would be forests of *Fagus sylvatica* and *Quercus petraea* or *Q. robur* that would be classified into different classes. This would be unusual compared to other phytosociological syntaxonomy and further explanation would be required for students, land managers and other stakeholders, adding inconsistency and discrediting the system.

Riparian and fen forests

Riparian forests are different in several aspects from most other deciduous forests. There are three main eco-topographical positions within them: the lower bed, usually occupied by willow scrub, the upper bed, normally populated by alder and poplar forests, and the floodplain, usually dominated by ashes and elms. The forests belonging to the second and third level can be included in the class *Querco-Fagetea sylvaticae* because they share a high number of species with the non-hygrophilous deciduous forests. However, due to their floristic and ecological specificity, they have been grouped in various ways by different authors, e.g. as a class of their own (*Alno glutinosae-Populetea albae*; Mucina et al. 2016) or joined together with the willow scrub formations of the lower river bed (*Salici purpureae-Populetea nigrae*; Rivas-Martínez et al. 2011). Concerning this second option, the nemoral *Querco-Fagetea* species are very rare in the willow scrub of the lower river bed. This lower level is subject to very frequent flooding episodes which cause strong erosion by the river current, preventing the edaphic development of the soil horizons and making the establishment of these herbaceous nemoral species impossible. Therefore, the lower-bed willow scrub vegetation would better be grouped in the independent class *Salicetea purpureae*, leaving the rest to be included in the *Querco-Fagetea*.

The non-willow scrub perifluvial forests separate clearly into two large biogeographic-climatic units which are represented by two orders: the eurosiberian Alno-Fraxinetalia excelsioris, and the mediterranean Populetalia albae (Biurrun et al. 2016). The proposal to join riparian forests with alder fen forests into one class Alnetea glutinosae (Berg et al. 2004) can be supported using floristic arguments, the dilemma is whether to recognize the class Alnetea glutinosae including only the Alnus fen forests, or to add the riparian humid ash forest to that class. The decision on how to establish the limit between two units, which have a certain number of plants in common, should also take account of the biogeographic context, as far as there are floristic arguments for either keeping them separated or joining them. The Alno-Fraxinetalia occur in many areas of southern Europe where the Alnetalia are lacking. The areas where both units coexist add weight to the argument to consider both as a single class, separated from the rest of Querco-Fagetea. However, in areas where humid forests all belong to the Alno-Fraxinetalia or to the Populetalia albae and the Alnetalia fen forests are absent, the recognition of a single class Querco-Fagetea could be a better solution. As the *Alnetalia* forests have peaty soils with stagnant anoxic water, there are a number of plants specifically adapted to that situation. Overall, I suggest maintaining a single class for all European deciduous forests: basiphilous, acidophilous, sub-mediterranean and riparian, but excluding the lower-bed willow scrub that are grouped in the Salicetea purpureae and the fen alder forests grouped in the Alnetea glutinosae.

Secondary forests

Several proposals to classify secondary forests have been published. Willner et al. (2016) described the order *Lonicero periclymeni-Betuletalia pubescentis*, which partially corresponds to the previously described *Betulo pendulae-Populetalia tremulae* (Rivas-Martínez et al. 2002). Both descriptions encompass secondary forests which have become established after a certain level of disturbance. Both are also very poorly defined floristically, with the *Betulo pendulae-Populetalia tremulae* including a few tree species of the genus *Betula, Populus tremula* or *Corylus avellana*. The rest of the flora is unspecific to secondary forests and occurs in many other units of the European deciduous forest, or even in forest mantles or shrublands. The floristic definition of these orders is significantly poorer than that of the other orders such as the Fagetalia sylvaticae, Quercetalia pubescenti-petraeae, Quercetalia roboris or Alno-Fraxinetalia excelsioris. These orders have a clear ecological character and they have been long recognized. Secondary forests are short-lived, filling a short time window in the succession and therefore can hardly develop a specific understory flora. Before humans started transforming the landscape at the beginning of the Neolithic age, the forests of temperate Europe were overwhelmingly primary; the secondary forests only occurred after natural disturbance episodes, which were much less frequent and less extensive than after the spread of agriculture and domesticated cattle herds. Thus, they temporarily occupied naturally disturbed patches cleared by wildfire, animal herbivory, windfalls, tree disease, etc. As their floristic differentiation is weak, it would be more consistent to frame these secondary forests in associations and alliances, rather than in one or two orders.

For the reasons stated, I believe that maintaining the unity of deciduous forests in the rank of a class would be a reflection of their unity of origin and their physiognomic and ecological similarity. This argument is also supported by a significant set of species that inhabit all of them. Additionally, secondary forests do not deserve the order status due to their weak floristic characterization. I propose that within the *Querco-Fagetea* class there are five European orders: *Fagetalia sylvaticae*, *Quercetalia roboris*, *Quercetalia pubescenti-petraeae*, *Alno-Fraxinetalia excelsioris* and *Populetalia albae*.

The class Quercetea ilicis

This class encompasses the sclerophyllous arboreal and shrubby vegetation in the Mediterranean region of Europe, the Middle East, and north Africa. It is a relatively species-poor unit with few herbaceous elements, due to summer drought stress and persistent shading of the understory. Distinguishing the different orders within this class has always been controversial, particularly in the western Mediterranean portion. Many of the genuine species of this class have a wide distribution across the whole Mediterranean region. In the central and eastern portion of the Mediterranean two orders have been recognized: Quercetalia calliprini and Pinetalia halepensis and for the central-western area the Quercetalia ilicis has been described. Rivas-Martínez (1975) described the order Pistacio lentisci-Rhamnetalia alaterni with representation throughout the Mediterranean. This order includes the sclerophylle macchia vegetation which can act as a mantle for the arboreal communities, i.e. their first seral stage, as well as potential vegetation in the areas where climatic or edapho-topographic conditions impede the establishment of a true forest. In Mucina et al. (2016) the Quercus-dominated forest communities remained in the order Querceta*lia ilicis* or the *Quercetalia calliprini*. However, it should be noted that some specialists in Mediterranean vegetation have published opposing views (Bolòs and Vigo 1984). The characteristic species listed by Rivas-Martínez (1975) frequently occur in forests: Asaparagus horridus, Chamaerops humilis, Clematis cirrhosa, C. flammula, Daphne gnidium, Euphorbia characias, Jasminum fruticans, Myrtus communis, Osyris alba, O. lanceolata, Pistacia lentiscus, P. terebinthus, Rhamnus alaternus, R. lycioides, but the order Pistacio-Rhamnetalia was described based on physiognomical criterion more than floristic citerion. In our opinion, it is more sustainable and consistent to separate the Quercetea ilicis into geographic orders, with the orders supported by the different endemic species which provide a firm floristic basis and confer stability to the syntaxonomy.

The relationship between the Pistacio-Rhamnetalia and Quercetalia ilicis seems to parallel that of the order Prunetalia spinosae and the rest of the Querco-Fagetea. When the order Pistacio-Rhamnetalia was described (Rivas-Martínez 1975), the Prunetalia spinosae was still accepted as an order of the Querco-Fagetea by most European phytosociologists, following the original concept of Tüxen (1952). The argument for establishing the Pistacio-Rhamnetalia was a physiognomic one (shrublands versus forests), following the criterion accepted at that time for the Prunetalia spinosae. However, more recently the general opinion regarding temperate forests and forest mantles has changed, and most authors have started to accept a separate class for the spiny hedges: Rhamno-Prunetea, restricting the Querco-Fagetea to forests. At the same time, with increased knowledge of the Quercetea ilicis, particularly in northern Africa (Quézel et al. 1988, 1992), a significant number of associations with a tree layer (Pinus halepensis, Juniperus turbinata, Tetraclinis articulata, etc.) have been described and attributed to the Pistacio-Rhamnetalia.

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Thus, the initial separation of forests and shrubland was broken and the separation of *Pistacio-Rhamnetalia* from *Quercetalia ilicis* became inconsistent.

The *Rhamno-Prunetea* is a relatively well defined class because it has few plants in the herb layer, in strong contrast to the *Querco-Fagetea*. Concerning the shrub species, it is true that many of them are shared between both classes, particularly with the order *Quercetalia pubescenti-petraeae*, but in the other orders, such as *Fagetalia*, most of these shrub species grow with diminished vitality. They hardly produce flowers and fruits and their populations are maintained by the constant seed rain provided by birds, as they are mostly endozoochorous. The spiny shrubs occurring inside the forests are therefore maintained by a metapopulational dynamic as sink populations. These shrub species need high light levels to produce a good set of fruit and fertile seeds.

The *Pistacio lentisci-Rhamnetalia alaterni* is quite different from the *Prunetalia spinosae* in this respect as the shrub species exhibit good vitality within most forests. It is impossible to recognize truly characteristic shrub or tree taxa within the *Pistacio lentisci-Rhamnetalia alaterni* and the herb species are limited and generally unspecific.

Acknowledgements

I wish to thank Jim Martin for the linguistic editing and suggestions in the text and the comments of an anonymous reviewer that have helped me to arrive at a better version of the manuscript, but I am especially grateful to Wolfgang Willner for his in-depth review and participation in the production of a more complete and scientifically sound final version.

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