ULTRASTRUCTURAL CHANGES IN CHLOROPLASTS OF MESOPHYLL CELLS OF CHLOROTIC AND PREMATURELY YELLOWED LEAVES OF BETULA PENDULA ROTHR.

KRYSTYNA PRZYBYŁ1, KRYSTYNA IDZIKOWSKA2

1 Department of Tree Diseases, Institute of Dendrology, Polish Academy of Sciences
ul. Parkowa 5, 62-035 Kórnik, Poland
e-mail: kmprz@man.poznan.pl
2 Laboratory of Electron Microscopy, Faculty of Biology, Adam Mickiewicz University
ul. Grunwaldzka 6, 60-780 Poznań, Poland
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ABSTRACT

The ultrastructure of chloroplasts was studied in mesophyll cells of the leaves of silver birch (Betula pendula) showing interveinal chlorosis or premature yellowing, in comparison with leaves without symptoms or exhibiting symptoms of natural senescence. The leaves were collected between May 26 to June 7 and additionally in the September 10-12 from the upper part of the crown, from increments of the past four years. No major difference in ultrastructure of chloroplasts was found between spongy and palisade mesophyll cells. The following senescence-related changes were observed in chloroplasts of prematurely yellowed leaves and showing interveinal chlorosis: reduced chloroplast size, degeneration of the membrane systems of thylakoids and increased electron density of plastoglobuli. The most electron dark globules (liquid droplets) were found together with starch grains in cells of spongy mesophyll of leaves showing interveinal chlorosis. Abnormal, spherical and rounded chloroplasts with electron-dark inside of thylakoids or the electron-dark stroma between thylakoids were found only in yellowed and chlorotic leaves in spring.

KEY WORDS: Betula pendula, leaves, interveinal chlorosis, premature yellowing, chloroplasts, TEM.

INTRODUCTION

For ten years, decline of birch stands (Betula pendula) has been recorded in Poland, but its intensity varied from one birch stand to the next. The most noticeable macroscopic symptoms occurring in above-ground organs are as follows: crown thinning due to dieback of fine twigs and branches in the tree-top, wounds and slime flux on the trunk, brown bark and sapwood inside branches and trunk, showing sometimes a water-saturated appearance (Przybył et al. 1998; Przybył and Złobińska-Podejna 2000; Przybył 2001). Leaves of the declining trees usually turn yellow prematurely. Irrespective of this symptoms, chlorosis between veins of leaves is observed in majority of trees. It is suspected that the symptoms in birch leaves could be related to the prolonged periods of drought recorded in Poland in the past few years. Moreover, it seems plausible that the K, Mg, and N deficiency is involved in accelerated yellowing (Przybył and Mańka 2000).

The ultrastructure of birch leaves has been studied during the autumnal senescence and in relation to injuries induced by ozone together with drought by Dodge (1970), Günthardt et al. (1993), Pääkkönen et al. (1995), Pääkkönen et al. (1997) and Pääkkönen et al. (1998). However to our knowledge there is a little information available from the literature on ultrastructural changes in chloroplasts with regard to macroscopic disease symptoms occurring on the leaves of declining trees.

The purpose of this research was to compare chloroplast ultrastructure in mesophyll cells of birch leaves showing interveinal chlorosis or yellowing of whole leaf blades in spring and of leaves without symptoms or exhibiting symptoms of natural senescence.

MATERIALS AND METHODS

Plant material and sampling methods

The study was conducted on silver birch trees aged 40-46. The trees grew on previously established experimental plots situated in following forest districts: Bierzwnik, Lipinki Lużyckie and Jarocin. Leaves showing some visual
TABLE 1. Collection sites and periods of Betula pendula leaves showing disease changes.

<table>
<thead>
<tr>
<th>Location</th>
<th>Disease changes on leaves</th>
<th>Collection</th>
</tr>
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<tbody>
<tr>
<td>1. Forest Division</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2. Department</td>
<td></td>
<td></td>
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<tr>
<td>3. Type of forest site and age of trees</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1. Bierzwnik</td>
<td>chlorosis between veins of blade leaf</td>
<td>late May</td>
</tr>
<tr>
<td>2. Kosobudka (protective zone)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3. Ca. 40</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1. Lipinki Lużyckie</td>
<td>yellowing of blade leaf</td>
<td>early June</td>
</tr>
<tr>
<td>2. Lipinki 781</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3. Wet coniferous mixed forest, 46</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1. Jarocin</td>
<td>spring – without disease symptoms (control 1)</td>
<td>early June</td>
</tr>
<tr>
<td>2. Gorzyce, 89</td>
<td>autumn – yellowing of blade leaf (control 2)</td>
<td>mid September</td>
</tr>
<tr>
<td>3. Wet coniferous mixed forest, 40</td>
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symptoms (Table 1, Fig. 1) and without macroscopic changes were harvested in the morning between May 26 to June 7 and additionally in the September 10-12 from the upper part of the crown of uprooted or removed trees, from increments of the past four years. The material was transported directly to the laboratory in portable iceboxes (at 5-10°C).

Sample preparation for transmission electron microscopy (TEM)

After cool transportation to the laboratory within one day, small sections (ca 2 mm x 2 mm) were cut from middle part of the leaf (near the main vein) of five leaves for each stand. The samples were immediately fixed in chilled 4% glutaraldehyde in 0.1 M cacodylate buffer at pH 7.2 for 20 h at 4°C and then rinsed in the same buffer. Postfixation was carried out in 1% osmium tetroxide (OsO₄) in the cacodylate buffer for 2 h at 4°C and then treated with 2% uranyl acetate for 2 h. Subsequently the material was dehydrated as follows: a) in a graded series of ethanol (30-90%), b) in ethanol-aceton mixtures (90% ethanol and 90% acetone (1:1) and 90% ethanol and 96% acetone (1:1) c) in absolute acetone and d) in 99% propylene oxide. The samples were embedded in Epon 812. Ultrathin sections were obtained with using ultramicrotome Leica-UltraTrent S. Ultrathin (ca. 70-90 nm thick) sections were poststained with uranyl acetate and lead citrate (Reynolds 1963) and viewed using Jeol type 1200 EX transmission electron microscope at an accelerating voltage of 80 keV.

The length of chloroplasts were measured from the photographs (Pääkkönen et al. 1995).

RESULTS

Leaves without symptoms in spring (control 1)

No apparent cytological differences were noted between the palisade and spongy mesophylls. Two types of the cells were observed in both kinds of mesophylls with regard to the degree of vacuolisation. One type had a large central vacuole with the other organelles situated in the periphery of cells. Mesophyll cells of the other type contained several vacuoles, smaller than in the first type, scattered in the cytoplasm. The cytoplasm of all observed cells was dense and contained minute granules. Besides, other organelles were evident inside both types of cells: nucleus, mitochondria, chloroplasts and endoplasmic reticulum.

Lens-shaped and elongated chloroplasts in cells of both palisade and spongy mesophylls were 4.0-5.5(-6) μm long. They exhibited a well-developed internal membrane system, which was electron-light (more transparent than stroma). Some chloroplasts contained usually electron-light globules of various size and number (Fig. 2). Small and single starch grains were sporadically observed.

Fig. 1. Disease symptoms of leaves of B. pendula: chlorosis between veins – A, B, C; leaves without symptoms and yellowed in late spring – D.
Fig. 2. Electron-light plastoglobuli (p) in chloroplasts of palisade mesophyll of leaves without symptoms in late spring (bar = 1 μm).
Fig. 3. Plastoglobuli (p) differing in electron density in chloroplasts of palisade mesophyll of leaves yellowed in autumn (disappearance of some part of thylakoids – arrow; bar = 1 μm).
Fig. 4. Spherical and rounded chloroplasts in mesophyll cells of leaves with interveinal chlorosis in late spring (s – starch, 1 – lipids; bar = 1 μm).
Fig. 5. Type of chloroplast in cells of spongy mesophyll of leaves with interveinal chlorosis in late spring (disappearance of envelope, swelling of thylakoids – arrow, strongly electron-dark stroma between thylakoids; bar = 1 μm).
Fig. 6. Type of chloroplasts in cells of mesophylls of leaves yellowed in late spring (disappearance of envelope and thylakoids – arrow and occurrence of plastoglobuli (p) differing in electron density (bar = 1 μm).
Fig. 7. Type of chloroplasts in cells of mesophylls (short sections of thylakoids with strongly osmiophilic inside, numerous plastoglobuli (p) varied in size (bar = 1 μm).
Leaves yellowed in autumn (control 2)

The vacuolization increased visually in cells of both spongy and palisade mesophyll, as compared with the leaves without symptoms and collected in spring. Chloroplasts were generally elongated and lens-shaped, 2.5-4.5 μm in length. Many of them contained plastoglobuli (Fig. 3), which differed in electron density. In this aspect, the following types of plastoglobuli were observed: 1) more electron-light than stroma; 2) similarly electron-dark to stroma; 3) with central part more electron-dark than the border. Some chloroplasts lacked electron-light thylakoids. Starch grains were rare and observed only in some cells. Osmophilic materials in the form of the granularity were observed in the cytoplasm of both kinds of mesophyll cells.

Leaves with interveinal chlorosis in spring

The cytoplasm of cells of both palisade and spongy mesophyll was rich in electron-dark granular deposit. The electron-dark material resembled that found in certain vacuoles. The majority of chloroplasts of both kinds of mesophyll cells were spherical shape or/and rounded (2.5-4.0 μm in length). Elongated and lens-shaped chloroplasts were rarely observed (Fig. 4). Thylakoids were not well-defined. In some cells, especially in spongy mesophyll, only electron-light residues of the thylakoids could be observed. These chloroplasts were filled with electron-dark (osmophilic) globules. Single starch grains were also found in the majority of cells of both kinds mesophyll (Fig. 4). Occasionally, spongy mesophyll cells contained chloroplasts that were elongated or lens-shaped but much smaller (2-2.5 μm in length) than those mentioned above. The smaller chloroplasts were lacking the envelope and some parts of their parallel membranes showed swelling (Fig. 5). The thylakoids were less crowded than in other chloroplasts. The stroma between thylakoids was strongly electron-dark, similarly to the osmophilic substances being in close contact with the chloroplasts (Fig. 5).

Leaves yellowed in spring

The cell cytoplasm of both palisade and spongy mesophyll was filled with electron-dark granules. The delimitation of certain organelles, like mitochondria and RE, appeared less clear than in the leaves without symptoms. Some cells contained a large and unshapely vacuole whereas in the others, several smaller vacuoles occurred. The majority of these vacuoles were filled with osmophilic globules.

Two types of chloroplasts showing some abnormalities in comparison with the ones observed in the leaves of control 1 were found in both palisade and mesophyll cells. One type was characterized by spherical chloroplast (length: 1.5-2.5 μm) whose envelopes were difficult to distinguish. The parallel thylakoids of these chloroplasts were more electron-light than the stroma. The disappearance of some sections of thylakoid membranes was also noted. Plastoglobuli showed differences in electron density. Usually their border was darker than the internal part (Fig. 6). Another type of chloroplasts, showing spherical shape and length range of 2-2.5 μm, was characterized by thylakoids looking as if they were divided into short sections. The inside of thylakoids was strongly osmophilic. In chloroplasts of this type, plastoglobuli were numerous and varied in size. The center of the plastoglobuli was more osmophilic than their border (Fig. 7). In these chloroplasts, starch grains were scarce and sometimes on many sections they could not be found.

DISCUSSION

Some of the senescence-related changes were observed in the chloroplasts of leaves of B. pendula showing premature yellowing and interveinal chlorosis. The changes include: reduced chloroplast size, degeneration of the membrane systems of thylakoids (swelling or disappearance) and increased electron density of plastoglobuli. Plastoglobuli differing in electron density occurred simultaneously in the chloroplast and the process of the density change started from their center or border. Generally, the features of senescence observed in chloroplasts in this study confirmed the results of Dodge (1970) and Pääkkönen et al. (1995), but in the present study, most of the strongly osmophilic globules (lipids) and most starch grains were found in the chloroplasts of the spongy mesophyll in leaves showing interveinal chlorosis. The ultrastructural symptoms seem to be rather non-specific. They were also detected in conifer needles with some stresses, such as ozone, sulphur dioxide and K and Mg deficiencies (Fink 1989, 1991; Sutinen 1987).

The abnormal rounded and spherical shape of the chloroplasts with electron-dark inside of thylakoids and electron-dark stroma between thylakoids were found only in the leaves exhibiting yellowing and chlorosis in spring. Moreover, in these leaves thylakoids were less crowded. Similar ultrastructural changes in chloroplasts of mesophyll cells of birch leaves were found after ozone treatment combined with drought (Pääkkönen et al. (1998), whereas protrusions of chloroplasts and disappearance of the envelope were observed after ozone exposure in spruce (Sutinen et al. 1992). Generally, no major differences in ultrastructure of chloroplasts were found between spongy and palisade mesophyll cells. The changes in chloroplast ultrastructure in mesophyll cells of leaves showing premature yellowing and interveinal chlorosis resembled those observed during natural senescence. However, certain traits of chloroplast ultrastructure can suggest the influence of drought and ozone which injured these organelles. Ozone sensitivity of birch (Betula pendula) was indicated by some authors (Mattyssek et al. 1991; Pääkkönen et al. 1993). In nature plants are usually exposed to ozone and drought stress simultaneously during warm periods of summer (McLaughlin and Downing 1996).

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LITERATURE CITED


ZMIANY W ULTRASTRUKTURZE CHLOROPLASTÓW MIĘKSIUU LIŚCII BETULA PENDULA WYKAZUJĄCYCH CHLOROZĘ I PRZEDWZESNE ŻÓŁKNIĘCIE

STRESZCZENIE

Zmiany w ultrastrukturze chloroplastów miękkisu palisadowego i assimilacyjnego badano u liści Betula pendula wykazujących objawy chlorozy i przedwczesne żółknienie. Liście do badań pobierano z górnej części korony (z przyrostów z czterech lat) w okresie od 26 maja do 7 czerwca. Kontrolę stanowiły liście nie wykazujące zmian chorobowych oraz liście pobierane w pierwszej połowie września.

Nie stwierdzono zasadniczych różnic w ultrastrukturze chloroplastów pomiędzy palisadowym a gębczastym miękkiem. Niektóre zmiany w chloroplastach u liści wykazujących przedwczesne żółknienie oraz chlorozy podobne były do zmian zachodzących w procesie naturalnego żółknienia (wielkość chloroplastów, degeneracja tylakoïdów gran i stromy oraz wzrastająca gęstość plastoglobuli). Najbardziej elektronowo cienkie ciała (lipidy) łącznie z ziarnami skrob i obserwowano w miękku gębczastym liści wykazujących chlorozę. Sferyczne i okrągłe chloroplasty z elektronowo cienkim wnętrzem tylakoïdów oraz chloroplasty z elektronowo cienmą stromą stwierdzono u liści chlorotycznych i przedwczesne żółtych. W dyskusji podkreślono czynniki, które mogły wpłynąć na obserwowane zmiany w chloroplastach.

SŁOWA KLUCZOWE: Betula pendula, liście, chloroza, przedwczesne żółknienie, chloroplasty, mikroskop elektronowy-transmisyjny.