POLYPEPTIDE COMPOSITION OF THYLAKOIDS FROM VIRIDIS AND XANTHA MUTANTS IN BARLEY

by

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The chloroplast membrane proteins from four recessive nuclear gene mutants of barley were studied with sodium dodecyl sulfate gel electrophoresis. The protein patterns of two mutants, *xantha-b*¹² and *viridis-zd*⁶⁹, consist of a summation of the polypeptides present in the chloroplast and etioplast membranes of the wild type. The pattern of *viridis-c*¹², a mutant deficient in photosystem II activity was lacking two bands, one of which is considered to be a component of the reaction center of photosystem II. The pattern of *viridis-k*²³ lacked three chloroplast specific bands, two of which are components of chlorophyll-protein complex II. The results are discussed in relation to previously established functional and structural characteristics of the mutant plastids.

1. INTRODUCTION

A survey of chlorophyll deficient seedling mutants in barley for alterations in the electrophoretic patterns of the chloroplast thylakoid proteins was carried out by N. C. NIELSEN (6, 13, 14). Among the recessive nuclear gene mutants

(20) studied by solubilization of their plastid membranes with phenol-acetic acid-urea two groups could be distinguished. Mutants with leaky blocks in chlorophyll synthesis and therefore limited capacities to form grana had reduced amounts of a membrane protein band

designated as J. Mutants with a tendency for stacking of thylakoids into giant grana and deficient in stroma thylakoids were deficient in the high molecular weight chloroplast membrane protein B.

In continuation of studies on the characterization of thylakoid proteins from barley mutants using membrane solubilization with sodium dodecyl sulfate (SDS) (12) and various conditions for electrophoresis (8) we report here on the thylakoid membrane pattern of four lethal nuclear gene mutants. Two of these are representative for the above mentioned two groups, mutant *xantha-b*¹² producing giant grana and mutant *viridis-k*²³ (like *xantha-l*³⁵) being defective in the conversion of Mg-protoporphyrin to protochlorophyllide (7). The other two mutants, *viridis-c*¹² and *viridis-zd*⁶⁹, are devoid of light induced CO₂ fixation and oxygen evolution, although the mutant seedlings contain as much as 20 per cent of the amount of chlorophyll present in wild-type seedlings (7, 9).

Chloroplast preparations of mutant *viridis-c*¹² lack significant photosystem II activity but could display photosystem I activity reaching one-third of wild-type rates (14, 17). HENNINGSEN (personal communication) found chloroplast preparations of *viridis-zd*⁶⁹ lacking in photosystem II activity but active in a photosystem I dependent reaction with rates surpassing those of *viridis-c*¹².

2. MATERIAL AND METHODS

Wild-type (*Hordeum vulgare* c.v. Svalöf's Bonus) and mutant barley seedlings were grown on tap water moistened vermiculite for 7 days at 20°C at constant illumination (1500 lux). The leaves were harvested just prior to homogenization and treated according to the flow sheet on the next page.

Electrophoresis was performed as described by HØYER-HANSEN, MACHOLD and KAHN (8) for the uniform gel system.

3. RESULTS

Chlorophyll contained in SDS micelles moved

as a band near the front during electrophoresis of the pigment-containing, SDS-solubilized plastid membranes from all four mutants. The green band of chlorophyll-protein complex II (CP II) was visible for the membrane preparations of *xan-b*¹² and *vir-zd*⁶⁹, whereas a band corresponding to chlorophyll-protein complex I (CP I) could not be discerned. The preparation of *vir-k*²³ yielded no CP II band but a weak CP I band was visible at the beginning of the electrophoretic run and disappeared as electrophoresis proceeded. Mutant *vir-c*¹² preparations gave neither CP I nor CP II bands.

The polypeptide pattern obtained after separation of acetone extracted and chemically reduced membrane proteins from the thylakoids of the four mutants is presented in Figure 1 together with the patterns from wild-type chloroplasts and wild-type etioplasts on either side. Aggregated polypeptides were found at the top of the gel with both wild-type and mutant membrane preparations. Such aggregates form upon removal of pigments from CP I (4, 11) and it is therefore possible that all four mutant membranes as well as those of the etioplasts (cf.8) contain the apoprotein of CP I. In Figure 2 the top portion of electrophoretograms from SDS-solubilized but not pigment extracted membrane preparations after staining with Coomassie blue are given. Only the wild-type membranes yielded a distinct CP I band corroborating the above mentioned observations of chlorophyll containing bands. The etioplasts and mutant plastids yielded aggregates not visible in the wild-type. Band I with an apparent molecular weight of 69,400 dalton representing the apoprotein of CP I (4, 8, 11) is present in chloroplast membranes of the wild-type but absent in those of etioplasts. It is present but less intense in mutants *xan-b*¹², *vir-k*²³ and *vir-zd*⁶⁹, and only represented by a very weak band in *vir-c*¹² (Figs. 1,2).

The two bands E and F (60,600 and 56,500 daltons respectively) ascribed to two subunits of the coupling factor (11, 21) are present in the membranes of the four mutants. Their intensity

relative to the other bands varies being highest in the pattern of *vir-k*²³ (Fig. 1). A comparison of the band pattern for polypep-

tides with an electrophoretic mobility higher than the coupling factor subunits E and F reveals drastic differences between the chloro-

Figure 1. Electrophoretic separation of membrane proteins from wild-type chloroplasts, wild-type etioplasts and plastids of the mutants *xantha-b*¹², *viridis-c*¹², *viridis-k*²³ and *viridis-zd*⁶⁹. Membrane were extracted with acetone prior to SDS-solubilization and reduction with mercaptoethanol. Electrophoresis was carried out with the uniform gel system.

plast and etioplast internal membranes. These have been described in (8) and the major band differences are diagramed in Figure 3, which is

directly comparable to Figure 1. A band by band comparison reveals the pattern of *xan-b*¹² and *vir-zd*⁶⁹ to be a summation of the band pat-

Figure 3. Diagram of electrophoretogram shown in Figure 1.

■ band present in chloroplast pattern only or in chloroplast and etioplast patterns
 ▨ band present in etioplast pattern only
 ▩ aggregates

terns of wild type thylakoid polypeptides and etioplast internal membrane polypeptides.

These two mutants thus appear to contain in their plastid membranes the polypeptides specific to the chloroplast, those specific to the etioplast and those common to the internal membranes of both chloroplasts and etioplasts. The membrane patterns of mutants *vir-c*¹² and *vir-k*²³ also contain chloroplast specific, etioplast specific and common polypeptides. The membrane pattern of mutant *vir-c*¹² contains all chlo-

Figure 2. Electrophoretic separation of the high molecular weight polypeptides of thylakoids from wild-type chloroplasts and plastids of four mutants. Membranes disintegrated in their native state with SDS.

roplast specific bands with the exception of two bands. A band with an apparent molecular weight of about 46,000 dalton is not detectable (arrow in Fig. 2). A polypeptide of this size has previously been implicated as a photosystem II reaction center protein (3, 10). Another chloroplast specific polypeptide with a somewhat higher electrophoretic mobility (double headed arrow in Fig. 2) is likewise missing or at least strongly reduced in this mutant. A reduction in the intensity of this particular band is observed for *vir-zd*⁶⁹.

The pattern of *vir-k*²³ is missing three bands. Two of these, IIa and IIb are the two polypeptides contained in chlorophyll-protein complex II (12). The third is a chloroplast specific polypeptide with a somewhat higher electrophoretic mobility.

4. DISCUSSION

The polypeptide patterns of the SDS-solubilized plastid membranes of the four mutants contained no unique bands, *i. e.*, apparently no polypeptides with an altered electrophoretic mobility. All bands present in the mutants had

counterparts either in wild-type chloroplast thylakoids or etioplast membranes or both. The membrane patterns of the mutants *vir-c*¹² and *vir-k*²³ were lacking two and three bands respectively. The simultaneous presence of chloroplast specific and etioplast specific polypeptides in the membrane patterns of the four mutants implies that the internal membranes of the mutant plastids can be viewed as a mixture of etioplast and chloroplast membranes. A general consequence of the four mutations thus is the continued synthesis of etioplast specific polypeptides in the light. This is associated with retarded plastid development (7) and suggests that the characteristic morphology of the membranes in etioplasts and mutant plastids, such as their arrangement in prolamellar bodies, is an expression of the presence of these etioplast specific membrane polypeptides. At this point it cannot be ruled out that the etioplast specific polypeptides represent contaminations from other cellular membranes.

Evidence is accumulating that CP I contains at least part of the reaction center of photosystem I (2,19). Measurements of photochemical activities during greening of etiolated barley have shown that photosystem I is functioning within the first hour of illumination (5,15). Thus a very rapid induction of synthesis of photosystem I

protein components after onset of illumination is required, if the components are not present already in the etioplast. If the polypeptide associated with the reaction center of photosystem I indeed corresponds to the band I polypeptide, and if this polypeptide is preformed in etioplast membranes, it could be present and hidden in the aggregated protein regularly found at the top of the gels after electrophoresis of etioplast membranes. The mutants *xan-b*¹², *vir-c*¹², *vir-zd*⁶⁹ yielded no CP I and *vir-k*²³ an unstable CP I upon membrane solubilization. The mutant membranes contained the 69,400 dalton band, which has been identified as the CPI apoprotein (4,11) and have an active photosystem I (7, 9, 14, 17 and HENNINGSEN, personal communication). Photosystem I activity of *vir-c*¹² is low on a chlorophyll basis. Interestingly, the 69,400 dalton band is weak in intensity in the membrane polypeptide pattern of this mutant. We interpret the absence or transitory presence of a CP I band in the patterns of the solubilized mutant membranes to be due to a decreased stability of the chlorophyll-protein complex I towards SDS rather than a deficiency of its protein component.

It is generally accepted that CP II does not contain the reaction center protein of photosystem II (1, 3, 18, 19). In previous studies it was shown that the four allelic chlorophyll b-less mutants, *chlorina-f* 2, -2800, -2808 and -3613 had good photosystem II activity (16, 18) in spite of the absence of CP II. In urea-containing gels CP II is resolved into the two protein bands, IIa and IIb, and the membranes of chlorophyll b-less mutants were found to lack IIa but to contain IIb (12). In the present study it is shown that mutant *vir-k*²³ is deficient in both polypeptides IIa and IIb, but nevertheless, it exhibits significant Hill activity (14,17). On the other hand, *vir-c*¹² and *vir-zd*⁶⁹ membranes, which are devoid of significant photosystem II activities have both the polypeptides IIa and IIb. From these three observations it is concluded that neither IIa nor IIb can be a reaction center protein for photosystem II.

The location of the reaction center protein of photosystem II in electrophoretic patterns of solubilized thylakoids has been investigated by CHUA & BENNOUN (3) with mutants of

Chlamydomonas. They found the absence of a protein with an apparent molecular weight of 47,000 to be correlated with lack of photosystem II activity. According to KLEIN & VERNON (10) a band corresponding to a polypeptide with a molecular weight of 44,000 is prominent in intensity in electrophoretograms of Triton X-100 membrane particles of spinach enriched in photosystem II activity. If the apparent molecular weight of the reaction center protein of barley is similar to that of spinach and *Chlamydomonas*, the 46,000 dalton band, marked with an arrow in Figure 1 is a candidate for this protein. This band is absent from the pattern of *vir-c*¹², a mutant which is deficient in photosystem II activity. Mutant *vir-zd*⁶⁹, which also is defective in photosystem II activity does contain the 46,000 dalton band. As mentioned above the pattern of *vir-c*¹² is missing a second chloroplast specific band (double-headed arrow in Figure 1), which is reduced in intensity in *vir-zd*⁶⁹. This band thus might also be important for photosystem II activity. Alternatively the *vir-zd*⁶⁹ mutation might cause an amino acid substitution in the 46,000 dalton polypeptide and thereby render it inactive. It has been suggested that CP II is necessary for thylakoid stacking (1,18). We have pointed out (12) that polypeptide IIa is not needed for stacking, as the chlorophyll b-less mutants, which lack this polypeptide, form considerable amounts of grana. The grana deficient mutant *vir-k*²³ investigated here lacks both the polypeptides IIa and IIb. It is therefore not excluded that polypeptide IIb functions in stacking. Since this mutant lacks an additional chloroplast specific polypeptide with a somewhat higher electrophoretic mobility (cf. Figs. 1,3), it is worth exploring with additional grana deficient mutants if this latter polypeptide is needed for stacking. Further studies are necessary to determine which of the three bands missing in the pattern of SDS-solubilized membranes from mutant *vir-k*²³ corresponds to the J band found affected when the membranes are solubilized in phenol-acetic acid-urea. Mutants *xan-b*¹², *vir-c*¹² and *vir-zd*⁶⁹ are rich in grana and relatively deficient in stroma thylakoids. NIELSEN (13) found the membranes of such mutants deficient in protein B. The relative low intensity of the high molecular weight protein I

band in the three mutants investigated here suggests that the B band corresponds to the protein I of the chlorophyll-protein complex I.

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