CONCISE REPORT

Hemin-Induced Conversion of Pyruvate Kinase Isozymes in K562 Cells

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The effects of hemin on the conversion of pyruvate kinase (PK) isozymes from the M₂-type to the L-type in K562 cells were investigated. Immunofluorescence, ion exchange chromatography, and electrophoretic studies showed that the untreated K562 cells contained only the M₂-type PK, while eight to 20 days after induction with hemin, concomitant with hemoglobin F synthesis, L-type PK levels increased while M₂-type PK levels decreased. Electrophoretic study revealed three hybrid isozymes of the L-type and M₂-type PK. We conclude that the conversion of PK isozymes from the M₂-type to the L-type in erythroid precursor cells occurs in the early stage of maturation.

We have recently reported the presence of M₂-type pyruvate kinase (PK) in the proerythroblast stage and conversion of the M₂-type PK to L-type PK with erythroid maturation¹ and demonstrated a compensatory production of M₂-type PK in the orthochromatic erythroblasts of PK-deficient patients.² Lozzio and Lozzio³ found that the K562 cell line obtained from the pleural effusion of a patient with Philadelphia chromosome-positive chronic myelogenous leukemia (CML) in blastic crisis can be differentiated into erythroid precursors after the addition of hemin.⁴ Hence, this cell line seems to be suitable for examining changes in PK isozymes in erythroid precursor cells. In this report, we describe conversion of PK isozymes from the M₂-type to the L-type in K562 cells after exposure to an inducing agent, hemin.
PK ISOZYME CONVERSION IN K562 CELLS

MATERIALS AND METHODS

The K562 cells were a kind gift of Dr. Yoshikura at the Department of Bacteriology, The University of Tokyo. Chromosome analysis was performed with the trypsin-Giemsa banding technique. The cultures contained $10^5$ cells per milliliter in RPMI 1640 medium (GIBCO, Grand Island, NY) supplemented with 10% fetal calf serum with or without 50 μmol/L hemin (Sigma Chemical Co, St Louis). After 20 days the cells were harvested by centrifugation at 4 °C. Lysis was performed by addition of four volumes of 10 mmol/L sodium phosphate buffer, pH 6.0, containing 2 mmol/L MgSO₄, 0.2 mmol/L fructose-1,6-diphosphate (FDP) and 0.1% 2-mercaptoethanol (buffer A) and by sonication for two minutes. The cellular debris was removed by centrifugation at 4 °C for 30 minutes at 20,000 g.

Pyruvate kinase activities were determined at 37 °C according to the International Committee for Standardization in Haematology (ICSH) recommended methods using a Gilford Model 250 spectrophotometer. The column was loaded with 0.08 to 0.1 U of the enzyme preparation. For linear gradient elution, a CM-Sephadex C-50 (Pharmacia Fine Chemicals, Piscataway, NJ) column (1 x 18 cm) was equilibrated with buffer A. A 70-mL linear gradient of sodium phosphate buffer from 10 to 100 mmol/L was used for elution; 60 fractions were collected. All the operations were carried out at 4 °C. Pyruvate kinase electrophoresis was performed using polyacrylamide thin layer gel.

Detection of L-type and M₂-type PK by the immunofluorescent method was described previously. Hemoglobin F was determined by the immunofluorescent technique described by Wood et al.

Fig 2. Immunofluorescent stain of K-562 cells. K-562 cells reacted with FITC-labeled anti-M₂-type PK (M₂), anti-L-type PK (L), and anti-hemoglobin F (Hb F). HEMIN(-), untreated K-562 cells; HEMIN(+) 8d, cells on day 8 of induction; HEMIN(+) 20d, cells on 20th day of induction.

Fig 3. Change of PK in K-562 cell. CM-Sephadex column chromatography. Abbreviations used are the same as those in Fig 2.
RESULTS

Cytogenetic Studies

The K562 cells were aneuploid with a prominent, near-triploid mode (66 to 72). The karyotypic analysis revealed that all the cells contained at least seven abnormal chromosomes and three to five markers (Fig 1). The Philadelphia chromosome was identified, but the 9q+ chromosome was not observed. The banding pattern showed that the 17q+ chromosome was probably a translocation between one chromosome 15 and one chromosome 17.

Immunofluorescent Studies

Figure 2 shows the results of the immunofluorescent studies. The fluorescence of M2-type PK was decreased after eight days of hemin induction, while that of L-type PK and hemoglobin F was increased with induction duration.

Ion Exchange Chromatography

The chromatographic patterns are shown in Fig 3. Purified L-type PK was eluted at an earlier position than the M2-type PK (Fig 3A). In the sample from uninduced K562 cells, the elution profiles showed a single peak at the same position as M2-type PK. On the other hand, in hemin-induced cells, a broad fraction was seen with a peak in a position between L- and M2-type PK (Fig 3D).

Pyruvate Kinase Electrophoresis

Electrophoretic studies revealed three bands in addition to the L-type PK band, the mobilities of which were between L-type and M2-type PK (Fig 4).

DISCUSSION

We have reported the conversion of PK isozymes from the M2-type to the L-type during early maturation of normal and PK-deficient erythroblasts.1,2 K562 cells established by Lozzio and Lozzio3 are known to differentiate into erythroid cells after the addition of hemin.4 Many studies concerning embryonic hemoglobin synthesis and membrane proteins have been reported, but little is known about the glycolytic enzymes of this cell line.

REFERENCES

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