GREEN SHEET REDIGEST

HB 883 **2024 Regular Session** **Phelps**

HEALTH CARE/FACILITIES: Allows the provision of data relative to sickle cell disease.

DIGEST

Present law establishes a registry to record data related to individuals diagnosed with sickle cell disease in this state.

Present law requires a healthcare provider to provide the La. Dept. of Health with patient date when an individual is diagnosed with sickle cell disease.

Present law directs the La. Dept. of Health to promulgate rules to govern the collection of data.

Proposed law directs the La. Dept. of Health to facilitate an agreement between sickle cell clinics and sickle cell associations to communicate patient information.

Proposed law directs the La. Dept. of Health to utilize existing software to expedite the communication between the department and the sickle cell associations.

(Amends R.S. 40:1125.13)

Summary of Amendments Adopted by House

The Committee Amendments Proposed by House Committee on Health and Welfare to the original bill:

1. Directs the La. Dept. of Health to facilitate an agreement between healthcare providers and sickle cell associations to communicate patient information.

Summary of Amendments Adopted by Senate

Committee Amendments Proposed by Senate Committee on Health and Welfare to the engrossed bill

1. Directs the La. Dept. of Health to facilitate an agreement between sickle cell clinics and sickle cell associations to communicate patient information.
2. Directs the La. Dept. of Health to utilize existing software to expedite the communication between the department and the sickle cell associations.

Prepared by Amanda Trapp.