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# FISCAL IMPACT REPORT

SPONSOR	PONSOR Komadina		ORIGINAL DATE LAST UPDATED	02/09/07	нв			
SHORT TITLE		Birth Defects Regi	stry		SB	667		
				ANAI	AYST	Geisler		
<b>APPROPRIATION (dollars in thousands)</b>								

Appropr	iation	Recurring or Non-Rec	Fund Affected
FY07	FY08		
	\$250.0	Recurring	General

(Parenthesis () Indicate Expenditure Decreases)

# SOURCES OF INFORMATION

LFC Files

<u>Responses Received From</u> Department of Health (DOH)

## SUMMARY

#### Synopsis of Bill

Senate Bill 667, for the Legislative Health and Human Services Committee, would appropriate \$250,000 from the general fund to the Public Health division of the Department of Health (DOH) for the purposes of supporting a birth defects registry. Any unexpended funds remaining at the end of FY08 shall revert to the general fund.

## FISCAL IMPLICATIONS

DOH notes that the funding in SB 667 is not part of the Executive Budget Request.

DOH notes that in order to implement an effective birth defects registry, \$150,000 of the funding would be used for two FTEs to support data management and referral of identified children to appropriate programs for follow up. The remaining \$100,000 would be used for other costs associated with developing the registry. As this is a long term project, recurring funding may be needed to sustain the registry.

# SIGNIFICANT ISSUES

Each year in New Mexico, approximately 600 infants (about 1 out of every 45 births) are born with a major birth defect. Birth defects were the second leading cause of infant mortality in New Mexico (New Mexico Selected Health Statistics Annual Report for 2004, NMDOH). Since 2000, birth defects have been a reportable condition in New Mexico. Babies born with birth defects have a greater chance of illness and long term disability than babies without birth defects. By identifying these children early in their life, we have the opportunity to assure that they receive the medical care and early intervention services to allow them to be productive adults and fulfill their potential.

NM had a Birth Defects Registry between 1995 and 2005 that was federally funded by the Center for Disease Control (CDC). The state's proposal for continued support from CDC was approved but not funded in 2005. There is currently no state or other funding to support this activity. The goal of the birth defects registry is to 1) monitor the prevalence of birth defects in the state, 2) provide data for the planning for services needed by these children and their families, 3) assess the impact of screening and prevention activities, and 4) to link children to necessary services such as case management for those that have complex medical needs and early intervention services to support the child's development.

GG/mt