California Health Benefits Review Program

Analysis of California Assembly Bill 97 Insulin Affordability

A Report to the 2021–2022 California State Legislature

April 16, 2021



Key FindingsAnalysis of California Assembly Bill 97 Insulin Affordability

Summary to the 2021–2022 California State Legislature, April 16, 2021



SUMMARY

The version of California Assembly Bill (AB) 97 analyzed by CHBRP would prohibit a deductible from being applied to insulin prescriptions. Other cost sharing (copayments, coinsurance) would still be permitted.

In 2022, of the 21.9 million Californians enrolled in state-regulated health insurance, 13.9 million of them would have insurance subject to, and potentially impacted by, AB 97.

Benefit Coverage: At baseline there are 118,014 enrollees who use insulin. 81,265 of enrollees using insulin *do not* have a deductible (69%), while 36,750 enrollees using insulin *have* a deductible (31%). Postmandate, 100% of enrollees would not need to meet their deductible before paying the normal copayment or coinsurance for their insulin prescription. AB 97 appears not to exceed the definition of essential health benefits (EHBs) in California.

Medical Effectiveness: CHBRP found a preponderance of evidence that higher cost sharing reduces adherence to insulin and lower cost sharing increases adherence to insulin. There is insufficient evidence on the associated effect of cost sharing for insulin on diabetes-related health outcomes, including HbA1c levels, outpatient visits, emergency department visits, hospitalizations, long-term complications, and disability/absenteeism rates.

Cost and Health Impacts¹: In 2022, AB 97 would increase total net annual expenditures by \$10,162,000 or 0.008% for enrollees with DMHC-regulated plans and CDI-regulated policies. This is due to an increase in \$23,853,000 in total health insurance premiums paid by employers and enrollees due to the cost-sharing cap, adjusted by a \$13,691,000 decrease in enrollee expenses.

The 31% of enrollees with a deductible at baseline would experience a 3% reduction in cost sharing, which results in a 0.26% increase in utilization of insulin postmandate for those enrollees. Average

cost sharing for these enrollees decreases from \$89 per prescription to \$87 per prescription. Almost 10% of enrollees who use insulin and have a deductible would experience a decrease in costsharing of more than \$20.

Enrollees using insulin at baseline who have a deductible tend to be users of other high-cost medications and other medical services. For example, among enrollees in health savings account (HSA)-eligible high deductible health plans (HDHPs) (and therefore with a combined medical and pharmacy deductible), almost three quarters (70%) of enrollees have expenditures for medical care and non-insulin brand name prescription medications that exceeds \$2,500 annually. As a result, almost all enrollees would reach their deductible or out of pocket maximum within a plan year, regardless of whether insulin is subject to the deductible.

Due to the small decrease in cost sharing and small increase in utilization, CHBRP projects no measurable public health impact. However, at the person-level, for enrollees who would not otherwise meet their deductible or out of pocket maximum and would therefore experience a higher change in cost sharing, AB 97 may result in improved glycemic control, a reduction in healthcare utilization, a reduction in long-term complications attributable to diabetes, and improved quality of life for enrollees that experience a decrease in cost sharing and improved insulin adherence, or begin using insulin due to reduced costs.

CONTEXT

Diabetes mellitus (DM), frequently referred to as diabetes, is one of the most common chronic conditions in California and the United States. According to the 2019 data from the Behavioral Risk Factor Surveillance System, about 10% of the adult population in California has been diagnosed with diabetes. The incidence of diabetes is highest among adults aged 65 and older.

and other aspects of health make stability of impacts less certain as time goes by.

¹ Similar cost and health impacts could be expected for the following year, though possible changes in medical science

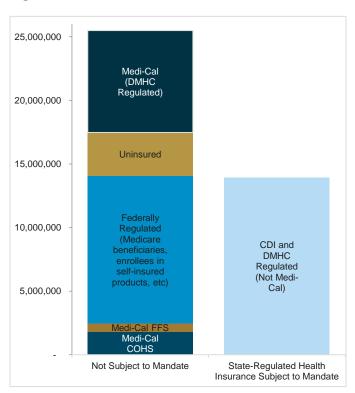


Diabetes is a chronic disease with short- and long-term health effects that prevent the proper production of and/or response to insulin, a hormone that facilitates the transfer of glucose into cells to provide energy.² Insulin can be used to treat all three types of diabetes: Type 1 diabetes mellitus (T1DM); Type 2 diabetes mellitus (T2DM); and gestational diabetes (GDM). The American Diabetes Association recommends different insulin regimens based on the type of diabetes a person has. Insulin is necessary for the treatment of T1DM and sometimes necessary for the treatment of T2DM and GDM.

In general, insulin has become expensive for individuals living with diabetes; therefore, cost may be a barrier to insulin use for some individuals. Other identified barriers to insulin use that are independent of cost include regimen complexity and treatment tolerability, as well as injection-related factors.

BILL SUMMARY

Figure A. Health Insurance in CA and AB 97



Source: California Health Benefits Review Program, 2021.

Assembly Bill (AB) 97 would prohibit a deductible from being applied to insulin prescriptions. Other cost sharing (copayments, coinsurance) would still be permitted. Figure A notes how many Californians have health

insurance that would be subject to AB 97 (approximately 35% of Californians).

IMPACTS

Benefit Coverage, Utilization, and Cost

Benefit Coverage

CHBRP estimates that, at baseline, there are 118,014 enrollees who use insulin in DMHC-regulated plans and CDI-regulated policies, where 81,265 enrollees (69%) using insulin do not have a deductible. CHBRP estimates 36,750 enrollees (31%) using insulin have a deductible (see estimates in Table 1). Postmandate, 100% of enrollees would not need to meet their deductible before paying the normal copayment or coinsurance for their insulin prescription.

Utilization

Postmandate, the group of enrollees with a deductible at baseline would experience an increase in utilization, because this group would experience a decrease in cost sharing due to the bill.

To estimate changes in utilization postmandate, CHBRP applied an estimate of price elasticity of demand to enrollees exceeding the cap at baseline. CHBRP bases the estimate of price elasticity on a Goldman et al. (2004) article that found use of insulin specifically decreased by 8% when copayments doubled. Based on this assumption, CHBRP estimates a 3% reduction in cost sharing for those enrollees who have a deductible at baseline, and therefore estimates a 0.26% increase in utilization of insulin postmandate for those enrollees.

Enrollees using insulin at baseline who have a deductible tend to be users of other high-cost medications and other medical services. A majority of these enrollees also have other prescription drug and medical costs that would cause them to meet their deductible or out-of-pocket maximum in a given year. Among enrollees with a pharmacy deductible, 64% have expenditures for other non-insulin brand name prescription medications that exceed \$500 annually, and therefore would cause them to meet their pharmacy deductible. Among enrollees enrolled in health savings account (HSA)-eligible high deductible health plans (HDHPs) (and therefore with a combined medical and pharmacy deductible), almost three quarters (70%) of enrollees have expenditures for medical care and non-

² Refer to CHBRP's full report for full citations and references.



insulin brand name prescription medications that exceed \$2,500 annually.

Expenditures

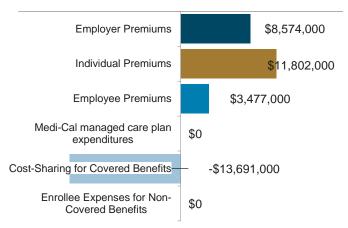
Based on Milliman's 2019 Consolidated Health Cost Guidelines Sources Database (CHSD) claims data, the average cost of insulin per prescription per month is \$491. For enrollees who do not have a deductible at baseline, the average monthly cost sharing for insulin is \$49, and for those enrollees for whom insulin is subject to the deductible at baseline, the average monthly cost sharing for insulin is \$89.

AB 97 would increase total net annual expenditures by \$10,162,000 or 0.008% for enrollees with DMHC-regulated plans and CDI-regulated policies. This is due to an increase in \$23,853,000 in total health insurance premiums paid by employers and enrollees, adjusted by a \$13,691,000 decrease in enrollee expenses.

Total premiums for private employers purchasing group health insurance would increase by \$8,574,000, or 0.02%. Total premiums for purchasers of individual market health insurance would increase by \$11,802,000, or 0.07%. The greatest change in premiums as a result of AB 97 is for the small-group policies (0.12% increase) and individual policies (0.11% increase) in the CDI-regulated market.

Due to the low change in utilization, no offsets are projected as a result of AB 97.

Figure B. Expenditure Impacts of AB 97



Source: California Health Benefits Review Program, 2021.

Enrollee Out-of-Pocket Expenses

For baseline insulin users, AB 97's prohibition of insulin being subject to the deductible only impacts those enrollees who have deductibles for pharmacy benefits.

Overall, 31% of enrollees who use insulin at baseline have deductibles. However, a majority of these enrollees also have other prescription drug and medical costs that would cause them to meet their deductible or out of pocket maximum in a given year. As a result, there is only a 3% decrease in average cost sharing.

It is possible that some enrollees who had deferred insulin treatment due to cost could begin using insulin postmandate; thus, this group of enrollees would incur cost sharing postmandate, whereas they did not have cost sharing at baseline. However, this group is estimated to be relatively small. Literature suggests approximately 2.5% of people who were prescribed insulin never started their prescription in the past year due to cost. Thus, for some enrollees, cost sharing may be the sole barrier to filling their insulin prescription.

The enrollees most likely to experience the greatest cost sharing reductions postmandate are those who are enrolled in plans that require significant deductibles to be met before coinsurance is applied to the insulin purchase, e.g., HDHPs, Bronze, and Silver plans. CHBRP's cost model estimates indicate that for enrollees subject to AB 97, approximately 19% of large-group, 37% of small-group, and 61% of individual market enrollees are in plans or policies with deductibles that apply to prescription drugs, where deductibles may have a material impact on insulin cost sharing. However, as mentioned previously, most enrollees who use insulin are likely to meet their deductible and out of pocket maximum regardless.

Medi-Cal

CHBRP assumes Medi-Cal's pharmacy benefit carve out transition will be complete by 2022. Because AB 97 only impacts DMCH-regulated pharmacy benefits, Medi-Cal managed care plans are not subject to the provisions of AB 97.

CalPERS

For CalPERS HMO enrollees, the impact on premiums is \$0 because there are no enrollees who have deductibles for pharmacy benefits.

Number of Uninsured in California

Because the change in average premiums does not exceed 1% for any market segment, CHBRP would expect no measurable change in the number of uninsured persons due to the enactment of AB 97.

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Medical Effectiveness

CHBRP found a *preponderance of evidence*³ from seven cross-sectional and retrospective studies on cost-related insulin use/adherence that cost sharing affects insulin use and adherence in patients with diabetes. These studies provided a *preponderance of evidence* that higher cost sharing reduces adherence to insulin, and lower cost sharing increases adherence to insulin.

CHBRP found *insufficient evidence*⁴ on the associated effect of cost sharing for insulin on diabetes-related health outcomes, including HbA1c levels, outpatient visits, emergency department visits, hospitalizations, long-term complications, and disability/absenteeism rates. Though the studies presented did report on these health and utilization outcomes, the findings were not specific to the effect of insulin alone, but combined with use of other oral antidiabetic medications and testing supplies.

There were several limitations that contributed to the gradings provided in this review, most notably the inherent differences between the types of diabetes conditions and the multifaceted nature of diabetes treatment. This resulted in a literature base that is not as rigorous and thereby limiting the certainty of conclusions drawn from the evidence.

Public Health

In the first year postmandate, 36,750 enrollees who use insulin and have a deductible at baseline would experience a small reduction in cost sharing. Almost 10% of enrollees who use insulin and have a deductible would experience a decrease in cost sharing of more than \$20. CHBRP projects that as a result, there would be a 0.26% increase in utilization of insulin. CHBRP found a preponderance of evidence that cost sharing for insulin is effective in improving adherence to insulin in patients with diabetes, and insufficient evidence on the effect of cost sharing for diabetes-related health outcomes. Due to the small decrease in cost sharing and small increase in utilization, CHBRP projects no measurable public health impact. However, at the person-level, for enrollees who would not otherwise meet their deductible or out of pocket maximum and would therefore experience a higher change in cost sharing, AB 97 may result in improved glycemic control, a reduction in healthcare utilization, a reduction in longterm complications attributable to diabetes, and improved quality of life for enrollees that experience a decrease in cost sharing and improved insulin adherence, or begin using insulin due to reduced costs.

Long-Term Impacts

CHBRP estimates annual insulin utilization after the initial 12 months from the enactment of AB 97 would likely stay similar to utilization estimates during the first 12 months postmandate. Utilization changes may occur if new diabetes products or medications change the landscape of insulin use for enrollees with diabetes, however CHBRP is unable to predict these types of changes. Similarly, health care utilization due to improved diabetes management may change in the long term. Reductions in significant complications or comorbidities may take years to develop, but are not trivial.

Similarly, reductions in significant complications or comorbidities may take years to develop, as would significant differences in disability and absenteeism. AB 97 is unlikely to impact these public health outcomes statewide, but at a person-level it could make a substantial difference in long-term health care spending, morbidity, and mortality for the enrollees who do experience substantial changes in cost sharing.

CHBRP estimates that AB 97 could improve disparities related to income for some enrollees who have cost-related barriers to insulin use. CHBRP is unable to estimate reductions in existing disparities. However, because the prevalence of diabetes is higher for Black persons than for Whites, and there is evidence that cost-related medication nonadherence is also more associated with Black persons, it is possible that this disparity may be reduced for the population AB 97 impacts.

The impact of AB 97 on premature mortality is unknown due to the lack of evidence that reduced cost sharing for insulin reduces mortality. However, well-controlled blood glucose results in fewer diabetes-related comorbidities (blindness, amputations, kidney disease, etc.). Therefore, for those patients who attain good glycemic control through increased adherence to insulin, these diabetes-related comorbidities that are known to lead to premature death could be prevented, delayed, or ameliorated.

effective, either because there are too few studies of the treatment or because the available studies are not of high quality. It does not indicate that a treatment is not effective.

³ Preponderance of evidence indicates that the majority of the studies reviewed are consistent in their findings that treatment is either effective or not effective.

⁴ Insufficient evidence indicates that there is not enough evidence available to know whether or not a treatment is



Essential Health Benefits and the Affordable Care Act

AB 97 would not require coverage for a new state benefit mandate and instead modifies cost sharing terms and conditions of an already covered medication. Therefore,

AB 97 appears not to exceed the definition of EHBs in California.

A Report to the California State Legislature

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April 16, 2021

California Health Benefits Review Program MC 3116; Berkeley, CA 94720-3116 www.chbrp.org

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The California Health Benefits Review Program (CHBRP) was established in 2002. As per its authorizing statute, CHBRP provides the California Legislature with independent analysis of the medical, financial, and public health impacts of proposed health insurance benefit-related legislation. The state funds CHBRP through an annual assessment on health plans and insurers in California.

An analytic staff based at the University of California, Berkeley, supports a task force of faculty and research staff from multiple University of California campuses to complete each CHBRP analysis. A strict conflict-of-interest policy ensures that the analyses are undertaken without bias. A certified, independent actuary helps to estimate the financial impact. Content experts with comprehensive subject-matter expertise are consulted to provide essential background and input on the analytic approach for each report.

More detailed information on CHBRP's analysis methodology, authorizing statute, as well as all CHBRP reports and other publications, are available at www.chbrp.org.

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Table 1. AB 97 Impacts on Benefit Coverage, Utilization, and Cost, 2022

	Baseline (2022)	Postmandate Year 1 (2022)	Increase/ Decrease	Change Postmandate
Benefit Coverage				
Total enrollees with health insurance				
subject to state-level benefit mandates (a)	21,945,000	21,945,000	0	0.00%
Total enrollees with health insurance	· · ·	, ,		
subject to AB 97	13,940,470	13,940,470	0	0.00%
Total percentage of enrollees with coverage				
subject to AB 97	64%	64%	0%	0.00%
Utilization and Cost				
Number of enrollees using insulin	118,014	118,014	-	0.00%
Enrollees whose claims are not	-,-	- , -		
subject to the deductible	81,265	118,014	36,750	45.22%
Enrollees whose claims are	·			
subject to the deductible	36,750	0	-36,750	-100.00%
Utilization per insulin user (# of 30-day				
supply insulin prescriptions per month)	0.85	0.85	0.00	0.18%
Utilization for enrollees whose claims <i>are not</i>			_	
subject to the deductible at baseline	0.85	0.85	0	0.00%
Utilization for enrollees whose claims are	0.05	0.05	0.00	0.000/
subject to the deductible at baseline	0.85	0.85	0.00	0.26%
Average monthly cost sharing for insulin per insulin user	\$76.95	\$75.17	-\$1.78	-2.32%
Average monthly cost sharing for enrollees	φ10.95	φ/3.1/	-\$1.70	-2.32/0
whose claims <i>are not</i> subject to the				
deductible at baseline	\$49.31	\$49.31	0	0.00%
Average monthly cost sharing for enrollees	ψ10.01	Ψ10.01	<u> </u>	0.0070
whose claims are subject to the deductible				
at baseline	\$89.45	\$86.86	-\$2.59	-2.89%
Average cost of insulin per prescription per	A 404	A 404		0.000/
month Expanditures	\$491	\$491	0	0.00%
Expenditures Premium (expenditures) by Payer				
	#FF 000 000 000	# 55.044.077.000	#0.574.000	2.000/
Private Employers for group insurance	\$55,032,803,000	\$55,041,377,000	\$8,574,000	0.02%
CalPERS HMO employer expenditures (b) (c)	\$5,765,017,000	\$5,765,017,000	\$0	0.00%
Medi-Cal Managed Care Plan expenditures	\$24,150,529,000	\$24,150,529,000	\$0	0.00%
Enrollee Premiums (expenditures)				
Enrollees for individually purchased insurance	\$15,847,507,000	\$15,859,309,000	\$11,802,000	0.07%
Individually Purchased – Outside Exchange	\$4,890,852,000	\$4,896,539,000	\$5,687,000	0.12%
Individually Purchased – Covered California	\$10,956,655,000	\$10,962,770,000	\$6,115,000	0.06%
Enrollees with group insurance, CalPERS HMOs, Covered California, and Medi-Cal				
Managed Care (c)	\$20,753,446,000	\$20,756,923,000	\$3,477,000	0.02%
Enrollee cost-sharing expenses	Ψ20,733,440,000	Ψ20,130,323,000	ψ5,+77,000	0.0270
Cost sharing for covered benefits	¢42 460 022 000	¢12 154 244 000	¢12 604 000	0.400/
(deductibles, copayments, etc.)	\$13,168,032,000	\$13,154,341,000	-\$13,691,000	-0.10%
Expenses for noncovered benefits (d) (e)	\$0	\$0	\$0	0.00%
Total Expenditures	\$134,717,334,000	\$134,727,496,000	\$10,162,000	0.008%

Source: California Health Benefits Review Program, 2021.

Notes: (a) Enrollees in plans and policies regulated by DMHC or CDI aged 0 to 64 years as well as enrollees 65 years or older in employer-sponsored health insurance. This group includes commercial enrollees (including those associated with Covered California or CalPERS) and Medi-Cal beneficiaries enrolled in DMHC-regulated plans.

- (b) About one in five (20.5%) of these enrollees has a pharmacy benefit not subject to DMHC. CHBRP has projected no impact for those enrollees. However, CalPERS could, postmandate, require equivalent coverage for all its members (which could increase the total impact on CalPERS).
- (c) Enrollee premium expenditures include contributions by employees to employer-sponsored health insurance, health insurance purchased through Covered California, and contributions to Medi-Cal Managed Care
- (d) Includes only expenses paid directly by enrollees (or other sources) to providers for services related to the mandated benefit that are not covered by insurance at baseline. This only includes those expenses that will be newly covered postmandate. Other components of expenditures in this table include all health care services covered by insurance.
- (e) Although enrollees with newly compliant benefit coverage may have paid for some tests before AB 97, CHBRP cannot estimate the frequency with which such situations may have occurred and therefore cannot estimate the related expense. Postmandate, such expenses would be eliminated, though enrollees with newly compliant benefit coverage might, postmandate, pay for some tests for which coverage is denied (through utilization management review), as some enrollees who always had compliant benefit coverage may have done and may continue to do, postmandate.

Key: CalPERS HMOs = California Public Employees' Retirement System Health Maintenance Organizations; CDI = California Department of Insurance; DMHC = Department of Managed Health; COHS = County Operated Health Systems, OPD = Outpatient Prescription Drug

POLICY CONTEXT

The California Assembly Committee on Health has requested that the California Health Benefits Review Program (CHBRP)⁵ conduct an evidence-based assessment of the medical, financial, and public health impacts of Assembly Bill (AB) 97, Insulin Affordability.

Bill-Specific Analysis of AB 97, Insulin Affordability

Bill Language

AB 97 would prohibit a deductible from being applied to insulin prescriptions. Other cost sharing (copayments, coinsurance) would still be permitted.

The full text of AB 97 can be found in Appendix A.

CHBRP previously analyzed a similar bill, AB 2203 Insulin Cost Sharing Caps, in 2020. Where applicable, this report builds on that analysis.

Relevant Populations

If enacted, AB 97 would apply to the health insurance of approximately 13.9 million enrollees (35% of all Californians). This represents 64% of the 21.9 million Californians who will have health insurance regulated by the state that may be subject to any state health benefit mandate law — health insurance regulated by the California Department of Managed Health Care (DMHC) or the California Department of Insurance (CDI). If enacted, the law would affect the health insurance of enrollees in DMHC-regulated plans and CDI-regulated policies, excluding enrollees with coverage through Medi-Cal managed care plans.

Although Medi-Cal managed care plans are subject to the Health and Safety Code, cost sharing for all Medi-Cal services is determined through the Welfare and Institutions Code (Section 14134).⁶ Therefore, because AB 97 only impacts cost sharing, Medi-Cal managed care plans are not subject to the provisions of AB 97.

Additionally, as of a to-be-determined date⁷, outpatient prescription drugs for Medi-Cal managed care beneficiaries are paid for on a fee-for-service basis and are "carved out" of care provided by Medi-Cal managed care plans. Although Medi-Cal managed care plans are subject to the Health & Safety Code, which AB 97 amends, the "carve out" of outpatient prescription drugs results in no impact to the coverage provided to Medi-Cal managed care plan beneficiaries. More information about outpatient prescription drug coverage among Californians with state-regulated health insurance is available in CHBRP's resource *Estimates of Pharmacy Benefit Coverage in California for 2022*.

Table 2 below indicates the presence of enrollees in the various market segments regulated by DMHC or CDI with a pharmacy benefit deductible. Enrollees in HSA qualified HDHPs have a combined medical and pharmacy benefit deductible.

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⁵ CHBRP's authorizing statute is available at www.chbrp.org/faqs.php.

⁶ Communication with the Department of Managed Health Care, March 2020; Communication with the Department of Health Care Services, April 2020.

⁷ The implementation originally scheduled for April 1, 2021 was delayed due to contracting issues, not due to a change in approach. CHBRP assumes implementation will occur by January 1, 2022.

Table 2. Enrollment in Plans/Policies with a Pharmacy Deductible Among Enrollees with a Pharmacy Benefit. 2022

Market Segment	Total Enrollment	No Deductible (\$0)	Low (<\$500)	High (a) (>=\$500)	HSA Qualified HDHP
CDI/DMHC Large					
Group (e)	8,097,000	81%	13%	0%	6%
CDI/DMHC Small					
Group (b)(c)(d)(f)	2,129,000	64%	23%	5%	9%
CDI/DMHC Individual	2,093,000	39%	27%	24%	10%
DMHC CalPERS	672,000	100%	0%	0%	0%
Total	12,991,000	72%	16%	5%	7%

Source: California Health Benefits Review Program, 2021.

Notes: CHBRP assumes only enrollees in HDHPs have combined medical and pharmacy deductibles.

- (a) does not include enrollees in HSA Qualified HDHP plans.
- (b) For this analysis, Small Group Gold plans are assumed to have no pharmacy deductible. In 2021, two of thirteen plans offered in this metal tier have a \$250 pharmacy deductible and CHBRP is unable to estimate the number of enrollees in those two plans.
- (c) For this analysis, Small Group Silver plans are assumed to be in the "Low Deductible" category. In 2021, nine of thirteen plans offered in this tier have a \$300-\$350 pharmacy deductible, three have \$0 deductible, and one has a deductible of \$500 and CHBRP is unable to estimate the varied enrollment among these plans.
- (d) For this analysis, Small Group Bronze plans are assumed to have a \$500 pharmacy deductible. In 2021, four of five plans offered in this tier have a \$500 pharmacy deductible and one plan has no pharmacy deductible. CHBRP is unable to estimate enrollment in that plan.
- (e) Large group distribution between no deductible and a deductible is from the Kaiser Family Foundation's 2019 Employer Health Benefits Survey (KFF, 2019), which reports the average pharmacy deductible of large groups with a pharmacy deductible is \$190. For this analysis, all large group enrollees with a deductible are in the "Low Deductible" category.
- (f) According to the Kaiser Family Foundation's 2015 Employer Health Benefits Survey (KFF, 2015), the average pharmacy deductible of small groups with a pharmacy deductible is \$160. For this analysis, all grandfathered small group enrollees with a deductible are assumed to be in the "Low Deductible" category.

Key: CalPERS HMOs = California Public Employees' Retirement System Health Maintenance Organizations; CDI = California Department of Insurance; DMHC = Department of Managed Health Care; HDHP = high deductible health; HSA = health savings account

Interaction with Existing Requirements

Health benefit mandates may interact and align with the following state and federal mandates or provisions.

California Policy Landscape

California law and regulations

Pharmacy benefits regulated by DMHC or CDI are required to provide coverage for insulin.⁸ As mentioned above, some enrollees with DMHC or CDI regulated health insurance do not have a pharmacy benefit or have a pharmacy benefit not regulated by DMHC or CDI.

DMHC-regulated plans and CDI-regulated policies are required to cover equipment and supplies for the management and treatment of insulin-using diabetes, non-insulin using diabetes, and gestational diabetes as medically necessary, even if the items are available without a prescription. This provision is not specific to enrollees with a DMHC or CDI regulated pharmacy benefit.

Senate Bill (SB) 852, signed into law in 2020, requires the California Health and Human Services Agency to enter into partnerships to increase patient access to affordable drugs, including producing or

⁸ H&SC 1367.51; IC 10176.61.

⁹ H&SC 1367.51; IC 10176.61.

distributing generic prescription drugs and at least one form of insulin. The Agency is required to submit a report to the Legislature by July 2023 that assesses the feasibility of these directives.

Existing California law limits cost sharing for prescription drugs to up to \$250 for a 30-day supply. Departed pharmacy deductibles are limited to \$500 for nongrandfathered individual and small group plans and policies. Departed in the policies of the prescription drugs to up to \$250 for a 30-day supply. Departed in the policies are limited to \$500 for nongrandfathered individual and small group plans and policies. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to \$250 for a 30-day supply. Departed in the prescription drugs to up to u

Similar requirements in other states

At least ten states have passed laws that limit cost sharing (copayment, coinsurance, and deductibles) for insulin, as of March 2021. Colorado, ¹² Illinois, ¹³ New York, ¹⁴ Washington, ¹⁵ Vermont, ¹⁶ and West Virginia ¹⁷ currently limit cost sharing for an insulin prescription to \$100 per 30-day supply, regardless of the amount or type of insulin. Connecticut limits total cost sharing to \$25 per 30-day supply, regardless of the amount or type of insulin. ¹⁸ Maine limits cost sharing for insulin to \$35 for a 30-day supply, regardless of the amount. ¹⁹ New Mexico limits cost sharing for a 30-day supply of preferred formulary insulin or the medically necessary equivalent to \$25. ²⁰ Utah limits cost sharing for a 30-day supply of at least one insulin in each "therapy category" to \$30 and prohibits insulin from being subject to the deductible. ²¹

Similar legislation has been recently introduced in at least 18 other states and DC.²² State bills would limit cost sharing for insulin prescriptions anywhere between \$25 for a 30-day to \$100 for a 30-day supply. A few states have legislation that would limit cost sharing for a 90-day supply. Additionally, several states have multiple proposals moving through the legislature with varying levels of cost sharing limits.

No other states have passed or have introduced legislation in 2021 that only prohibits insulin from being subject to the deductible.

Federal Policy Landscape

On March 11, 2020, the Centers for Medicare & Medicaid Services (CMS) announced the Part D Senior Savings Model, a voluntary model that enables participating Part D enhanced plans²³ to lower Medicare beneficiaries' cost sharing for insulin to a maximum \$35 copay per 30-day supply throughout the benefit year.²⁴ The program began January 1, 2021.

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¹⁰ H&SC 1342.73; IC 10123.1932.

¹¹ Ibid.

¹² Colorado House Bill 19-1216.

¹³ Illinois Senate Bill 667.

¹⁴ New York State Senate Bill 7506-B.

¹⁵ Washington Senate Bill 6087; although the bill language states cost sharing is limited regardless of amount of insulin prescribed, the bill does not state whether this applies regardless of type of insulin.

¹⁶ Vermont House Bill 969

¹⁷ West Virginia House Bill 4543.

¹⁸ Connecticut House Bill 6003

¹⁹ Maine Legislative Document 2096.

²⁰ New Mexico House Bill 292.

²¹ Utah House Bill 207 Insulin Access Amendments, 2020.

²² Legislative search through PoliticoPro, conducted on March 6, 2021.

²³ Approximately 60% of Medicare Part D prescription drug plans, nationally, are "enhanced" in 2020. KFF, Medicare Part D: A first look at prescription drug plans in 2020. 2019. Accessed on March 31, 2020 at https://www.kff.org/report-section/medicare-part-d-a-first-look-at-prescription-drug-plans-in-2020-issue-brief/.

²⁴ Centers for Medicare and Medicaid Services. 2020. CMS launches groundbreaking model to lower out of pocket expenses for insulin. Accessed on March 13, 2020. Available at: https://www.cms.gov/newsroom/press-releases/cms-launches-groundbreaking-model-lower-out-pocket-expenses-insulin.

Affordable Care Act

A number of Affordable Care Act (ACA) provisions have the potential to or do interact with state benefit mandates. Below is an analysis of how AB 97 may interact with requirements of the ACA as presently exists in federal law, including the requirement for certain health insurance to cover essential health benefits (EHBs).^{25,26}

For the 2021 plan year for nongrandfathered group plans, the annual out-of-pocket maximums for an individual are \$8,550 and \$17,100 for a family.²⁷ This means once an enrollee or a family reach these outof-pocket maximums, they are no longer responsible for additional cost sharing responsibilities for the remainder of the plan year.

Any changes at the federal level may impact the analysis or implementation of this bill, were it to pass into law. However, CHBRP analyzes bills in the current environment given current law and regulations.

Essential Health Benefits

Nongrandfathered plans and policies sold in the individual and small-group markets are required to meet a minimum standard of benefits as defined by the ACA as essential health benefits (EHBs). In California, EHBs are related to the benefit coverage available in the Kaiser Foundation Health Plan Small Group Health Maintenance Organization (HMO) 30 plan, the state's benchmark plan for federal EHBs.^{28,29} CHBRP estimates that approximately 4.2 million Californians (11%) have insurance coverage subject to EHBs in 2022.30

States may require plans and policies to offer benefits that exceed EHBs.31 However, a state that chooses to do so must make payments to defray the cost of those additionally mandated benefits, either by paying the purchaser directly or by paying the qualified health plan. 32,33 Health plans and policies sold outside of the health insurance marketplaces are not subject to this requirement to defray the costs. State rules related to provider types, cost sharing, or reimbursement methods would not meet the definition of state benefit mandates that could exceed EHBs.34

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²⁵ The ACA requires nongrandfathered small-group and individual market health insurance — including but not limited to QHPs sold in Covered California — to cover 10 specified categories of EHBs. Policy and issue briefs on EHBs and other ACA impacts are available on the CHBRP website: www.chbrp.org/other_publications/index.php.

²⁶ Although many provisions of the ACA have been codified in California law, the ACA was established by the federal government, and therefore, CHBRP generally discusses the ACA as a federal law. ²⁷ HealthCare.gov. Out-of-pocket maximum/limit. Accessed on March 6, 2021. Available at

https://www.healthcare.gov/glossary/out-of-pocket-maximum-limit/.

²⁸ CCIIO, Information on Essential Health Benefits (EHB) Benchmark Plans. Available at: https://www.cms.gov/cciio/resources/data-resources/ehb.html.

²⁹ H&SC Section 1367.005; IC Section 10112.27.

³⁰ CHBRP, Estimates of Sources of Health Insurance in California in 2022. Available at: www.chbrp.org/other_publications/index.php.

³¹ ACA Section 1311(d)(3).

³² State benefit mandates enacted on or before December 31, 2011, may be included in a state's EHBs, according to the U.S. Department of Health and Human Services (HHS). Patient Protection and Affordable Care Act: Standards Related to Essential Health Benefits, Actuarial Value, and Accreditation. Final Rule. Federal Register, Vol. 78, No. 37. February 25, 2013. Available at: https://www.gpo.gov/fdsys/pkg/FR-2013-02-25/pdf/2013-04084.pdf.

³³ However, as laid out in the Final Rule on EHBs which HHS released in February 2013, state benefit mandates enacted on or before December 31, 2011, would be included in the state's EHBs, and there would be no requirement that the state defray the costs of those state-mandated benefits. For state benefit mandates enacted after December 31, 2011, that are identified as exceeding EHBs, the state would be required to defray the cost.

³⁴ Essential Health Benefits. Final Rule. A state's health insurance marketplace would be responsible for determining when a state benefit mandate exceeds EHBs, and QHP issuers would be responsible for calculating the cost that must be defrayed.

AB 97 would not require coverage for a new state benefit mandate and instead modifies cost sharing terms and conditions of an already covered medication. Therefore, AB 97 appears not to exceed the definition of EHBs in California.

Trends in Cost of Insulin Prescriptions

The average list price of brand-name insulin nearly tripled between 2007 and 2018, increasing by 262% (Table 2) (Hernandez et al., 2020). While the average net price also increased, the increase was smaller (51%) and was offset by discounts such as those paid by manufacturers. The price increases were higher between 2012 and 2015, but began to level out in 2016. More recently, insulin has been increasing at a rate closer to 1.5% per year. The reasons for insulin prices increasing are not entirely clear, but they are due in part to the complexity of drug pricing in general and of insulin pricing in particular (Cefalu et al., 2018).

Table 3. Summary of 2007-2018 Changes in List and Net Prices

	All Drug Classes	Insulins
List Price		
Change from 2007-2018	159%	262%
Annual mean change per year	9.1%	12.6%
Net Price		
Change from 2007-2018	60%	51%
Annual mean change per year	4.5%	4.2%
List price increase offset by discounts	62%	81%

Source: California Health Benefits Review Program, 2021, as adapted from Hernandez et al., 2020.

Notes: List price is defined as the price of a product as listed by the manufacturer. Net price is defined as the ratio between company-reported sales for each product and the number of units sold in the US.

As the price of insulin has increased, so too have patient out-of-pocket costs. Between 2006 and 2013, average cost sharing per insulin user among Medicare Part D enrollees increased by 10% per year for all insulin types (Cefalu et al., 2018).

The increases in list price, net price, and out-of-pocket costs are substantially higher than increases due to inflation. Overall inflation between 2006 and 2013 was 2.2%, medical care service costs increased by 3.8%, and spending for all prescription drugs increased by an average of 2.8% (Cefalu et al., 2018).

Cost Sharing and Outpatient Prescription Drug Benefits

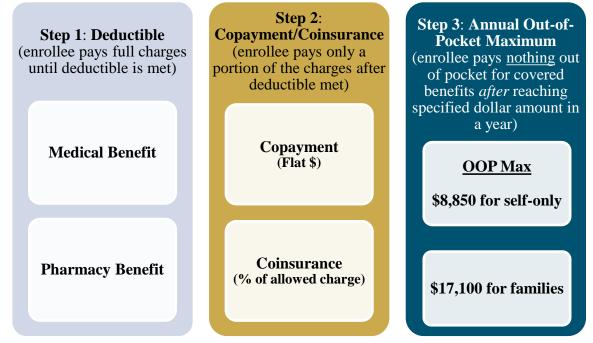
This section provides an overview of the cost sharing structures used for health insurance benefits, including prescription drugs. Payment for covered health insurance benefits is shared between the payer (e.g., health plan/insurer or employer) and the enrollee. Common cost-sharing mechanisms include copayments, coinsurance, and/or deductibles (but do not include premium payments). There are a variety of cost-sharing mechanisms that can be applicable to covered benefits (Figure 3). Some health insurance benefit designs incorporate higher enrollee cost sharing in order to lower premiums. Reductions in

allowed copayments, coinsurance, and/or deductibles can shift the cost to premium expenses or to higher cost sharing for other covered benefits.³⁵ Annual out-of-pocket maximums for covered benefits limit annual enrollee cost sharing (medical and pharmacy benefits). After an enrollee has reached this limit through payment of coinsurance, copayments, and/or deductibles, insurance pays 100% of the covered services. The enrollee remains responsible for the full cost of any tests, treatments, or services that are not covered benefits.

An enrollee using insulin may experience multiple forms of cost sharing. If an enrollee has a plan with a deductible and the enrollee has not yet met the deductible, the enrollee would be responsible for the full cost of care and prescriptions until that deductible is met. Once an enrollee has met their deductible, the enrollee would be responsible for the copayment or coinsurance associated with the insulin prescriptions. Should an enrollee's out-of-pocket expenses meet the annual out-of-pocket maximum, the enrollee would no longer be responsible for cost-sharing responsibilities.

AB 97 would instead require that an enrollee only pay the cost sharing for insulin prescriptions, regardless of whether they have met their deductible.

Figure 3. Overview of the Intersection of Cost-Sharing Methods Used in Health Insurance



Source: California Health Benefits Review Program, 2021.

Note: Steps 1 and 2 are not mutually exclusive. Under certain circumstances (e.g., preventive screenings or therapies), enrollees may pay coinsurance or copayments prior to their deductible being met; also, copayments and coinsurance may be applied against the deductible in some circumstances. The figure assumes that the enrollee is in a plan with a deductible. If there is no deductible, then enrollee pays a coinsurance and/or a copayment beginning with the first dollar spent (Step 2).

Key: OOP Max = annual out-of-pocket maximum.

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³⁵ Plans and policies sold within Covered California are required by federal law to meet specified actuarial values. The actuarial value is required to fall within specified ranges and dictates the average percent of health care costs a plan or policy covers. If a required reduction in cost sharing impacts the actuarial value, some number of these plans or policies might have to alter other cost-sharing components of the plan and/or premiums in order to keep the overall benefit design within the required actuarial value limits.

High Deductible Health Plans (HDHPs)

Both DMHC-regulated plans and CDI-regulated policies may be designated HDHPs. HDHPs are a type of health plan and requirements are set by federal regulation.³⁶ As the name implies, these plans include a deductible – but they are not allowed to have separate medical and pharmacy deductibles. For the 2020 plan year, the Internal Revenue Service (IRS) defines an HDHP as any plan with a deductible of at least \$1,400 for an individual and \$2,800 for a family. Annual out-of-pocket expenses for coverage of innetwork tests, treatments, and services, which would result from cost sharing³⁷ applicable after the deductible is met, are not allowed to be more than \$6,900 for an individual and \$13,800 for a family.³⁸

Health Savings Account (HSA) Qualified HDHPs

To be eligible to establish a Health Savings Accounts (HSA) for taxable years beginning after December 31, 2003,³⁹ (and so to be eligible to make tax-favored contributions to an HSA), a person must be enrolled in an HSA qualified HDHP.

In order for a HDHP to be HSA qualified, it must follow specified rules regarding cost sharing and deductibles, as set by the IRS. Generally, an HDHP may not provide benefits for any year until the deductible for that year is satisfied - but federal law provides a safe harbor for the absence of a deductible applicable to preventive care.⁴⁰ Therefore an HDHP may cover preventive care benefits without any deductible or with a deductible below the minimum annual deductible – but is not required to do so for a specified list of preventive services. The list of preventive services for which application of a deductible is not required includes treatments for chronic conditions.⁴¹ Insulin is listed a treatment for chronic conditions and therefore the requirements of AB 97 would not interfere with an HDHP's qualification for an HSA.

Allowed Cost Amounts for Medical Services

Insurers usually negotiate how much they will pay for the costs of covered health care services with health care providers and suppliers (Center on Budget and Policy Priorities, 2018). These negotiated amounts are known as the "allowed cost amount." Health care providers, including hospitals and physicians, participating in a plan's network agree to accept these payment amounts when an enrollee covered by the plan uses covered services. The cost-sharing charges the enrollee owes (for example, a 20% coinsurance rate) are based on this allowed cost amount. If an enrollee uses a service that is not covered or sees a provider that is not within the insurer's network, the overall charge, including an enrollee's cost sharing, could be higher than the allowed amount.

³⁶ HealthCare.gov, Glossary: High Deductible Health Plans. Available at https://www.healthcare.gov/glossary/high-deductible-health-

plan/#:~:text=For%202019%2C%20the%20IRS%20defines,or%20%2413%2C500%20for%20a%20family.Accessed on March 5, 2021.

³⁷ Such as copays and coinsurance applicable to the covered test, treatment, or service.

³⁸ There is no annual out-of-pocket expenses limit for coverage of out-of-network tests, treatments, and services.

³⁹ Section 1201 of the Medicare Prescription Drug, Improvement, and Modernization Act of 2003, Pub. L. No. 108-173, added section 223 to the Internal Revenue Code

⁴⁰ For more information on screening services, see Notice 2004-23, 2004-15 I.R.B. 725, available at IRS.gov/irb/2004-15 IRB#NOT-2004-23.

For additional guidance on preventive care, see Notice 2004-50, 2004-2 C.B. 196, Q&A 26 and 27, available at IRS.gov/irb/2004-33_IRB#NOT-2004-50; and Notice 2013-57, 2013-40 I.R.B. 293, available at IRS.gov/pub/irs-drop/n-13-57.pdf.

⁴¹ For information on preventive care for chronic conditions, see Notice 2019-45, 2019-32 I.R.B. 593, available at IRS.gov/pub/irs-drop/n-19-45.pdf.

BACKGROUND ON DIABETES MELLITUS AND INSULIN FOR GLYCEMIC CONTROL

Maintaining a proper blood sugar (glucose) level is critical to maintaining good health and preventing complications for people with diabetes mellitus (DM). This section defines DM, the prevalence of DM, and describes the subject of AB 97, insulin for management of diabetes.

What Is Diabetes Mellitus?

DM, commonly referred to as diabetes, is a chronic disease with short- and long-term health effects (discussed below) that prevent the proper production of and/or response to insulin, a hormone that facilitates the transfer of glucose into cells to provide energy (NIDDKD, 2017a). There are three primary types of diabetes, and insulin can be used to treat all three types:

- Type 1 diabetes mellitus (T1DM) is an autoimmune disease, most commonly diagnosed during childhood/adolescence that attacks and destroys the insulin-producing cells in the pancreas. In addition to dietary modifications, treatment requires lifetime use of daily insulin injections and/or an insulin pump used to replace the patient's impaired ability to produce insulin, and attention to diet.
- Type 2 diabetes mellitus (T2DM) is most commonly diagnosed in middle-aged or older adults, although it has been increasingly diagnosed in children and adolescents at a rate of 5% between 2002 and 2015 (CDC, 2015; CDC 2020). Type 2 diabetes prevents the body from properly responding to insulin (known as insulin resistance). In some cases, people with T2DM also do not make enough insulin. It is associated with obesity, genetics, and lifestyle patterns. Treatments for T2DM include diet modifications, exercise, weight loss, oral medications, non-insulin injected medications, and/or insulin depending on the severity of the disease, which progresses over time especially with inadequate treatment.
- Gestational diabetes (GDM) develops only in women who are pregnant and is generally
 diagnosed in the second trimester (Blumer et al., 2013). For most, this is a transient condition that
 resolves following delivery; however, these women remain at higher risk for T2DM later in life.
 Treatments include diet modifications, exercise, oral medication, and insulin.

Diabetes Mellitus: Short- and Long-Term Effects

Short-term effects

Achieving stable, healthy blood glucose levels is challenging for individuals with diabetes. On a daily basis, people with diabetes can experience swings between very high blood glucose levels (*hyperglycemia*) and extremely low blood glucose levels (*hypoglycemia*). Changes in stress, sleep, physical activity, diet, acute illnesses, and changes in non-diabetes medications can contribute to hyperand hypoglycemic events. *Hyperglycemia* is exhibited through increased thirst or hunger, frequent urination, headache, and fatigue. Left untreated, particularly in T1DM, it may develop into ketoacidosis where the body develops a toxic amount of ketones (toxic acids) for energy, which can lead to coma or death.

Symptoms of *hypoglycemia* can begin as mild (e.g., anxiety, sleepiness, and tremors) and, if left untreated, escalate to serious health events such as cognitive dysfunction, seizures, coma, and death (Unger, 2012). Some patients (between 20% and 40% of T1DM patients and 10% of T2DM patients) are diagnosed with *hypoglycemia unawareness*, a condition in which individuals are unable to sense dangerously low blood sugar early enough to reverse it, which puts them at high risk for severe hypoglycemic events requiring hospitalization (Martin-Timon and Canizo-Gomez, 2015). People with this condition are required to perform more frequent blood glucose testing than those who can feel their blood glucose levels dropping. Vigersky et al. (2015) estimated that among people with hypoglycemic

unawareness, 2.4 to 8.1 hospitalizations occur annually among T1DM patients, and 2.1 to 5.9 hospitalizations per year among T2DM patients. Hypoglycemia unawareness occurs more frequently among those with a longer duration of diabetes, who are insulin dependent, and/or have a history of hypoglycemic events (Martin-Timon and Canizo-Gomez, 2015).

For pregnant women, uncontrolled GDM may lead to complications during pregnancy including abnormal fetal growth, need for extra testing during pregnancy, preeclampsia, and possible early and/or more invasive delivery methods including cesarean. Infants of women with GDM can suffer complications during and directly after birth, including hypoglycemia and hyperbilirubinemia (jaundice), but most are transient with some infants requiring NICU care (NIDDKD, 2017b).

Long-term effects

Time spent in hyperglycemia and frequency and severity of hyper- and hypoglycemia over a lifetime are associated with serious morbidity and mortality outcomes. In the United States, diabetes is the leading cause of blindness, amputations, and kidney failure, and a key contributor to stroke, heart disease, dental disease, nerve damage, and premature death (NIDDKD, 2017a) due to suboptimal blood sugar control. In the long term, uncontrolled GDM puts pregnant women and their infants at higher risk of developing T2DM later in life (NIDDKD, 2017b). Although people with diabetes may not avoid all associated comorbidities, tightly controlled blood glucose over time may prevent, delay, or ameliorate some comorbidities.

COVID-19 effects

According to the CDC, individuals with T2DM are at an increased risk of severe illness from COVID-19 and individuals with T1DM or GDM *might* be at an increased risk of severe illness from COVID-19 (CDC, 2021). The CDC and American Diabetes Association (ADA) indicate that individuals with well-managed diabetes are likely at lower risk from severe complications and recommend that individuals with diabetes take their insulin and other diabetes oral medications as prescribed, regularly test and keep track of blood glucose levels, and to ensure that individuals have a 30-day supply of all diabetes medications, including insulin (CDC, 2021; ADA, 2021). Viral infections, like COVID-19, as well as elevated blood glucose levels, can increase inflammation in individuals with diabetes and that dual effect on inflammation could contribute to more severe symptoms and complications. Tightly controlled blood glucose by adhering to prescribed insulin and diabetes medication regimens may help mitigate these effects (ADA, 2021).

Prevalence of Diabetes Mellitus in California

Diabetes is one of the most common chronic conditions in California and the United States. According to the 2019 Behavioral Risk Factor Surveillance System, about 10% of the adult population in California has been diagnosed with diabetes (America's Health Rankings, 2020). Approximately 2.6% of adults age 18 to 44, 13.9% of adults ages 45 to 64, and 22.8% of adults over age 65 have diabetes (America's Health Rankings, 2020).

The following are the most recent prevalence estimates for the privately insured population⁴² by type of diabetes for adults, pregnant women, and youth:

- Adults: Of the estimated 6% (906,000) privately insured adult (aged 18–64 years) enrollees with diabetes, about 9.4% have T1DM and about 87.3% have T2DM (Table 3) (CHIS, 2019b).
- Pregnant women: The 2018 CHIS estimates that GDM occurs in 5.1% of pregnancies among non-diabetic enrollees (CHIS, 2018), which is similar to national estimates that range between 2% and 10% of pregnancies are affected by gestational (CDC, 2019a). According to the CDC, approximately 50% of women with GDM develop T2DM (CDC, 2019a).

⁴² As discussed in the Policy Context section, Medi-Cal managed care plans are not impacted by AB 97.

• Youth: CHIS does not report diabetes in those under age 18 years after 2007; however, national data published by the CDC estimates that in 2018 0.25% of youth under age 20 years are diagnosed with T1DM (~89%) and T2DM (~11%) (CDC, 2020).

Table 4. Prevalence of Type 1 and Type 2 Diabetes among Privately Insured Californians Diagnosed with Diabetes. 2019

Diabetes Type	Percent (n) Diagnosed with Diabetes			
California Adults Aged 18–64 Years with Diabetes (n=906,000)				
Type 1	9.4% (85,000)			
Type 2	87.3% (791,000)			
Unknown/another type*	3.3% (29,000)			

Source: California Health Benefits Review Program, 2021. Based on 2019 data from the California Health Interview Survey (CHIS).

Diabetes Management Using Insulin

In individuals without diabetes, beta cells within the pancreas release the hormone insulin when food is ingested to help the body use or store blood sugar. As described earlier in this section, the hallmark difference between T1DM and T2DM is the body's ability to create or utilize insulin to regulate blood sugar levels. Clinical practice recommendations for prescribing insulin for glycemic control are presented in the *Medical Effectiveness* section. This section summarizes the types of insulin products available and mechanisms of delivery.

Types of insulin

Insulin is classified by the rate at which it acts in the body. The differences for each type depend on onset, peak, duration, concentration, and delivery method. Table 4 summarizes types of insulin products. Short or rapid-acting insulin (bolus or prandial) is used to control blood sugar during meals as fat tissue absorbs it quickly from the bloodstream. Intermediate or long-acting insulin (basal insulin) is absorbed at a slower stabilizing rate, which is used to control blood sugar during one's sleep or fasting periods (Shah et al., 2016). Premixed insulin products may be useful for those with poor eyesight or dexterity, or who have trouble measuring the correct dosages for injection (ADA, 2020b).

Insulin products may also be identified as animal, human, or analog. The first insulin products were isolated from animals, and later, the technology to create a synthetic insulin allowed for greater production volume. These synthetic versions were called *human* to distinguish them from the insulin derived from animals (Tibaldi, 2014). Later advances included the development of rapid-acting insulin analogs and long-acting basal analogs (Tibaldi, 2014). The long-acting basal analogs are one of the most widely prescribed, and have been used to help patients with T2DM achieve glycemic control with lower risk of hypoglycemia. However, the cost of insulin analogs is much greater than the original human and animal-derived insulins (Cefalu et al., 2018).

Patients with T1DM require insulin for their diabetes management, and will use both bolus and basal insulin. Therefore, the prevalence of insulin use among individuals with T1DM is 100%. Insulin may or may not be used for someone with T2DM, and for those who are prescribed insulin, they may use one or both types (ADA, 2019). The prevalence of insulin use among T2DM varies; however, the CDC reports that for adults aged 20 and older with diagnosed T1DM or T2DM, 10.9% started using insulin within a year of their diagnosis (CDC, 2020). Women with GDM may also be prescribed one or both types of insulin (ADA, 2019).

^{*}CHIS reports these data as statistically unstable. CHIS permits respondents to select "Unknown or Another type" in response to its "type of diabetes" question. Examples of other types of diabetes may include maturity-onset diabetes of youth; from surgery, medications, infections, pancreatic disease, or other illnesses including cystic fibrosis.

Table 5. Types of Insulin Products

Categories of Insulin	Delivery	Onset	Peaktime	Duration
Rapid-acting insulin	Vial, pen, cartridge, or inhaler	15 minutes	1-2 hours	3-4 hours
Regular or short-acting insulin	Vial	30 minutes	2-3 hours	3-6 hours
Intermediate-acting	Vial or pen	2-4 hours	4-12 hours	12-18 hours
Long-acting	Vial or pen	Several hours	Does not peak	Up to 24 hours
Ultra long-acting	Vial or pen	6 hours	Does not peak	36+ hours
Premixed insulin products	Vial or pen	Varies	Varies	Varies

Source: California Health Benefits Review Program, 2021, based on Cefalu et al., 2018; ADA, 2019; and ADA, 2020b.

Delivery mechanisms

There are various delivery methods of insulin, but subcutaneous injections with a vial and syringe or prefilled pen are the most common forms (Shah et al., 2016; Zhang et al., 2019). Insulin pumps are devices that are worn by the individual and mimic the function of the pancreas to deliver small steady doses of insulin (HHN, 2018). Insulin pumps can deliver both basal and bolus insulin, and the decision to use one depends on the patient's needs and preferences (HHN, 2018). Conventional delivery mechanisms of syringes, pens, and pumps may be uncomfortable or inconvenient for some with diabetes. A non-injection insulin product available since 2015 is an inhaled insulin (ADA, 2020b). This delivery method is used as a rapid-acting insulin before meals and must be used in conjunction with injectable long-acting insulins (ADA, 2020b). Insulin is not available as a pill; because it is a peptide hormone, the body would digest it and it would not reach the blood stream (ADA, 2020a; Shah et al., 2016). Developments to oral routes of administration are currently under investigation, as are buccal, peritoneal, and transdermal (Shah et al., 2016).

Barriers to Diabetes Control

Insulin-Associated Barriers

In general, insulin has become expensive for individuals living with diabetes. See the *Policy Context* section for information on the rising cost of insulin and common cost-sharing mechanisms. As mentioned, for those with insurance, the patient is responsible for applicable cost sharing for insulin. See more details about the cost of insulin in the *Benefit Coverage*, *Utilization*, *and Cost Impacts* section. Additionally, the *Medical Effectiveness* section describes how the effects of cost sharing impact insulin use and adherence. Patients with T1DM have less flexibility in altering use due to cost as insulin is required for their glycemic control.

Other identified barriers to insulin use that are independent of cost include regimen complexity and treatment tolerability (Brod, 2012; Peyrot et al., 2010), as well as injection-related factors (Peyrot et al., 2010; Rubin et al., 2009). Patients reported that injections interfered with daily activities, caused pain at the injection site, and caused embarrassment in social situations (Pawaskar et al., 2007; Peyrot et al., 2010). A systematic review by Davies et al. (2013) also cited difficulty with insulin use while travelling,

challenging social situations, and forgetting as barriers. Additionally, fear of weight gain and hypoglycemia were cited as barriers to starting insulin therapy, though were less of a concern once insulin treatment had started (Davies et al., 2013). Following a set dosing schedule is also cited as challenging and inconvenient for patients (Pawaskar et al., 2007). The most common reasons for dosing irregularities range from inconsistent eating patterns to running low on insulin (Brod et al., 2012).

Additional Barriers to Diabetes Control

Barriers to insulin use present challenges in glycemic control for individuals with diabetes that are prescribed insulin therapy. However, additional barriers to glycemic control exist for patients that may or may not be taking insulin. In order to effectively manage diabetes, additional non-insulin prescription medications or medical supplies such as blood glucose testing strips and devices may be required. For example, affordability of blood glucose testing devices is one barrier. In a retrospective database analysis, Yeaw and colleagues identified that testing strips and supplies accounted for 27% of the cost of insulin prescription and supplies required for self-management of blood glucose levels (Yeaw et al., 2012). Similarly, it was reported that for patients with lower incomes, nearly two-thirds experienced challenges with affording diabetes equipment (Herkert et al., 2019). While the economic implications of insulin costs seem to be well-understood, there is a need for additional studies to provide greater understanding of costs associated with monitoring supplies. If a patient encounters barriers in accessing or using devices to monitor blood glucose levels regularly, they have reduced ability to administer insulin correctly and safely. Another important component to diabetes management is a change in behaviors and lifestyle factors, which each present a wide variety of barriers on their own. Lifestyle changes required for diabetes management include self-management education, weight control through diet and exercise, and regular medical care to monitor for comorbid conditions or complications from diabetes (ADA, 2018).

Disparities⁴³ and Social Determinants of Health⁴⁴ in Diabetes

Per statute, CHBRP includes discussion of disparities and social determinants of health (SDOH) as it relates to diabetes. Disparities are differences between groups that are modifiable. CHBRP found literature identifying disparities in diabetes by race/ethnicity, gender, and age.

Disparities

Race or ethnicity

In California, Hispanics (10.5%), Blacks (8.8%), American Indian/Alaska Natives (7.5%), and Asian/Pacific Islanders (6.2%) have higher prevalence of T2DM than non-Hispanic Whites (4.9%), and Hispanics and Blacks have two times higher prevalence: 1 in 20 non-Hispanic Whites have T2DM, compared with 1 in 10 Hispanics and 1 in 11 Blacks (Conroy et al., 2014). This is consistent with racial/ethnic differences found nationally: prevalence of diagnosed diabetes was highest among American Indians/Alaska Natives (14.7%), people of Hispanic origin (12.5%), and non-Hispanic blacks (11.7%), followed by non-Hispanic Asians (9.2%) and non-Hispanic Whites (7.5%) (CDC, 2020). However, Whites are more likely to develop T1DM than Blacks and Hispanic/Latino Americans (CDC, 2019b).

These differences may be attributed to biological factors, health system factors, and social factors (Spanakis et al., 2013). Multiple studies have shown that compared to non-Hispanic Whites, non-Hispanic

⁴³ Several competing definitions of "health disparities" exist. CHBRP relies on the following definition: Health disparity is defined as the differences, whether unjust or not, in health status or outcomes within a population (Wyatt et al., 2016).

⁴⁴ CHBRP defines social determinants of health as conditions in which people are born, grow, live, work, learn, and age. These social determinants of health (economic factors, social factors, education, physical environment) are shaped by the distribution of money, power, and resources and impacted by policy (adapted from: CDC, 2014; Healthy People 2020, 2019). See CHBRP's SDOH white paper for further information: http://chbrp.com/analysis_methodology/public_health_impact_analysis.php.

blacks and Mexican Americans have increased insulin resistance and differences in insulin secretion (Golden et al., 2012; Spanakis et al., 2013). Evidence is mixed regarding significant racial or ethnic differences in adherence to diabetes medication, including insulin (Brod et al., 2012; Golden, 2012). However, Kang et al. (2018) report significant racial/ethnic disparities for cost-related medication nonadherence for non-Hispanic blacks compared to non-Hispanic Whites. Obesity is correlated with diabetes risk in racial or ethnic minority populations (Golden et al., 2012). This is due in part to racial disparities observed in obesity, particularly among non-Hispanic blacks and Hispanics (Golden et al., 2012). Additional research is needed to establish the underlying risk factors that contribute to disparities in obesity rates, but it is hypothesized that cultural norms, obesity definition cut-points, and immigration status may be factors (Golden et al., 2012).

Gender⁴⁵

The prevalence of T2DM is higher in men (7.2%) than women (6.4%) in California (Conroy et al., 2014). This trend is consistent with national prevalence rates: approximately 11% of men in the United States have diabetes, while 9.5% of women do (CDC, 2020). Gender was also found as a correlate of nonadherence to insulin therapy in a large systematic review (Davies et al., 2013). Female gender was associated with lower adherence. Among younger females in particular, intentional insulin omission may be related to weight control and eating disorders (Peyrot et al., 2010).

Age

Across all age groups, the prevalence of T1DM is low in California (<2%) (Conroy et al., 2014). However, differences exist across age groups in the state: the prevalence of T2DM is less than 2% for adults aged 44 years and under, but rises sharply to 10% for those aged 45 to 64 years, and to 17% for those aged 65 years and older (Conroy et al., 2014). Similarly, in the United States, the rate of adults with diagnosed diabetes (T1DM or T2DM) increases with age, though national rates report reaching 26.8% among those aged 65 years and older (CDC, 2020). Davies et al. (2013) noted that for studies within the review (one study of T1DM, two studies of T2DM, one study of both T1DM and T2DM, and one with type of diabetes not reported), age was a predictor for adherence to insulin therapy; however, two studies indicated older patients were more adherent, while one showed that younger patients were more adherent. Peyrot et al. found no association between age and intentional insulin omission among patients with T1DM, and it was proposed that perhaps patients "age-out" of the behavior as they get older (Peyrot et al., 2010). Conversely, when including cost as a factor, younger age (<55) was at significantly greater risk for cost-related medication nonadherence for diabetes when compared to older adults age 75 and over (Kang et al., 2018).

Social Determinants of Health (SDOH)

SDOH include factors outside of the traditional medical care system that influence health status and health outcomes (e.g., income, education, geography). CHBRP found literature indicating that level of education, income, and health literacy have a role in diabetes.

Education

The prevalence of diagnosed T2DM is twice as high in California adults without a high-school diploma (9.9%) compared to those with a college degree (4.8%) (Conroy et al., 2014). Studies show that there is a relationship between low educational attainment and high prevalence of T2DM (Borrell et al., 2006). Those with more education are thought to utilize resources and knowledge to prevent or better control their diabetes (Borrell et al., 2006). Higher levels of education are associated with better health outcomes,

⁴⁵ CHBRP uses the NIH distinction between "sex" and "gender:" "'Sex' refers to biological differences between females and males, including chromosomes, sex organs, and endogenous hormonal profiles. 'Gender' refers to socially constructed and enacted roles and behaviors which occur in a historical and cultural context and vary across societies and over time." (NIH, 2019).

as research shows that individuals will take part in more preventive measures (Clark and Utz, 2014). Additionally, a higher level of education is associated with higher socioeconomic stability, which in turn promotes healthy behaviors (Borrell et al., 2006).

Income

The percentage of adults in California with diagnosed T1DM or T2DM is almost double for those with family incomes below 200% of the federal poverty level (FPL) (100–199% below FPL have 9.1% diagnosed diabetes, 0–99% below FPL have 8.7% diagnosed diabetes) compared to those whose income is 300% or more above the FPL (5.3%) (Conroy et al., 2014). In a systematic review and meta-analysis of 23 studies, socioeconomic status was strongly associated with an increased risk of T2DM (Agardh et al., 2011).

Peyrot and colleagues (2010) also found that respondents with higher household income were less likely to skip insulin injections as prescribed. This may be due in part to easier access to medications and supplies among individuals with higher income, but it is also likely that higher socioeconomic status is associated with more access to diabetes education, higher health literacy, greater control over one's daily routines, and better problem-solving skills (Peyrot et al., 2010). In addition, individuals with better socioeconomic status have lower cost-related medication nonadherence for diabetes (Herkert et al., 2019; Kang et al., 2018). The rate of cost-related nonadherence decreased as annual household income level increased. The rate is tripled for those without insurance compared to those with insurance, and is higher for individuals on insulin therapy compared to those who are not on insulin therapy (Kang et al., 2018).

Health literacy

Health literacy refers to an individual's capacity to read, understand, and make use of healthcare-related information for decision making and self-care. CHBRP did not find that lower health literacy was a risk factor for diabetes diagnosis. However, there was strong evidence in the literature that for individuals diagnosed with diabetes, health literacy was significantly correlated with management of diabetes and health outcomes.

In diabetes, health literacy is particularly important for disease management elements such as understanding treatment regimens, reading and interpreting food labels, carbohydrate counting, and appropriate insulin administration (Ahola and Groop, 2013). While low levels of health literacy are not necessarily shown to prevent blood glucose monitoring, interpreting the results and acting accordingly in response may be compromised among individuals with low health literacy. Poor health literacy is also related to reduced ability to recall oral medical instructions (Ahola and Groop, 2013). In a study examining the relationship between racial disparities and poor glycemic control in diabetes, the authors concluded that health literacy was associated with diabetes medication adherence (Osborn et al., 2011).

Societal Impact of Diabetes in California

The presence of diabetes in California creates a societal impact. In dollar terms, the societal impact can be indirect (e.g. lost wages, etc.), as well as direct (e.g. medical care, etc.). Total economic costs for T1DM and T2DM (direct plus indirect costs) in California were reported to be \$55.5 billion in 2013 (median \$5.9 billion) (Shrestha et al., 2018). For non-Medicare or Medicaid payers (private insurance, other payers, and out of pocket from patients), medical costs were \$11.7 billion in California (Shrestha et al., 2018). According to the American Diabetes Association⁴⁶, total direct medical expenses in California were estimated to be \$27.6 billion in 2012 for diagnosed and undiagnosed diabetes, prediabetes, and GDM. An additional \$9.5 billion was spent on indirect costs due to lost productivity. Indirect costs have

⁴⁶ American Diabetes Association (ADA). The Burden of Diabetes in California. Available at: http://main.diabetes.org/dorg/PDFs/Advocacy/burden-of-diabetes/california.pdf

also been reported as high as \$32.6 billion when including morbidity and premature mortality costs (Shrestha et al., 2018). Please note, the societal impact discussed here is relevant to a broader population than AB 97 impacts, which would affect the health insurance of a subset of Californians (see *Policy Context*). See the *Benefit Coverage, Utilization, and Cost Impacts* section for estimates of cost impacts for the specific population targeted by AB 97.

MEDICAL EFFECTIVENESS

As discussed in the *Policy Context* section, AB 97 would prohibit a deductible from being applied to insulin prescriptions. Other cost sharing (copayments, coinsurance) would still be permitted. Additional information on the management of diabetes and insulin cost sharing is included in the *Error! Reference s ource not found.* section. The medical effectiveness review summarizes findings from evidence⁴⁷ on the effects of cost sharing on insulin use and adherence for patients with diabetes (type 1 diabetes mellitus [T1DM], type 2 diabetes mellitus [T2DM], and gestational diabetes [GDM]) and how insulin treatment adherence affects the management of diabetes.

Clinical Practice Guidelines for Diabetes Mellitus

The American Diabetes Association (ADA, 2020b) recommends different insulin regimens based on the type of diabetes a person has. Insulin is necessary for the treatment of T1DM and sometimes necessary for the treatment of T2DM and GDM usually after diet, lifestyle, and oral anti-diabetic medications are insufficient to lower HbA1c levels to a goal of less than 7% for most adults. According to the guidelines, T1DM patients typically inject insulin subcutaneously in two patterns, one basal (continuous) form of insulin and one bolus (mealtime) form of insulin. This is achieved by four injections per day of insulin (of a long-acting insulin analog, typically dosed one to two times daily, and a rapid-acting insulin analog dosed three times daily before meals) or using an insulin pump (where a rapid-acting insulin is delivered both as the basal and bolus insulin). The most common types of insulin used are rapid-acting insulins (reaches bloodstream 15 minutes after injection, peaks at 1 to 2 hours, continues to work for 3 to 4 hours), and long-acting insulins (takes several hours to reach bloodstream and maintains glucose levels throughout a 24-hour period). Less frequently used insulins are regular human insulin (30 minutes to reach bloodstream, peaks at 2 to 3 hours, works for 3 to 6 hours) and intermediate-acting insulin (2 to 4 hours to reach bloodstream, peaks 4 to 12 hours, works for 12 to 18 hours) (see Table 5 in the Error! Reference s ource not found. section). Insulin regimens (i.e., types, timing, and doses) are typically determined by health care providers' recommendations, but may vary, and be self-adjusted by an individual based on diet, exercise, and other factors. Despite the variety of insulin regimens, the long-term complications of diabetes (e.g., eye, kidney, and nerve damage) can be best prevented by reaching glycemic targets/A1c goals with intensive insulin therapy or continuous subcutaneous administration through an insulin pump (ADA, 2020b).

T2DM is a progressive disease and use of insulin is often required for its management, especially with increased diabetes duration (ADA, 2020b). According to the guidelines, Metformin, an oral glucose-lowering medication, is the preferred initial pharmacologic agent for the treatment of T2DM, in combination with lifestyle modifications. The choice for the next step in therapy depends on patient-specific factors (e.g., presence of atherosclerotic cardiovascular disease, heart failure, chronic kidney disease, obesity). For patients who are on combination therapy, insulin therapy is generally initiated after a patient is on a class of medications called glucagon-like peptide-1 receptor agonists (GLP-1). When insulin is initiated in a patient with T2DM, usually a long-acting insulin is added as one injection daily to the medication regimen. Over time, a patient may require prandial insulin, and a rapid-acting insulin is added at mealtimes. Similar to T2DM, treatment of GDM may require insulin therapy and depends on patient-specific factors (ADA, 2020b).

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⁴⁷ Much of the discussion in this section is focused on reviews of available literature. However, as noted in the section on Implementing the Hierarchy of Evidence on page 11 of the Medical Effectiveness Analysis and Research Approach document (posted at http://chbrp.com/analysis_methodology/medical_effectiveness_analysis.php), in the absence of fully applicable to the analysis peer-reviewed literature on well-designed randomized controlled trials (RCTs), CHBRP's hierarchy of evidence allows for the inclusion of other evidence.

Research Approach and Methods

Studies of cost sharing related to insulin use and adherence for diabetes were identified through searches of PubMed, the Cochrane Library, Web of Science, and the Cumulative Index of Nursing and Allied Health Literature. Websites maintained by the following organizations that produce and/or index meta-analyses and systematic reviews were also searched: the Agency for Healthcare Research and Quality (AHRQ), the International Network of Agencies for Health Technology Assessment (INAHTA), the National Health Service (NHS) Centre for Reviews and Dissemination, the National Institute for Health and Clinical Excellence (NICE), and the Scottish Intercollegiate Guideline Network. The American Diabetic Association (ADA) website and available resources were also searched as pertinent to this bill and reviewed.

The search was limited to abstracts of studies published in English. The search was limited to studies published from 2020 to present, because CHBRP had previously conducted thorough literature searches on these topics in 2020 for AB 2203. Of the 122 articles found in the literature review, 19 were reviewed for potential inclusion in this report on AB 97, and a total of two new studies were included in the medical effectiveness review for this report, as well as five studies that were included in the previous review for AB 2203. The other articles were eliminated because they did not focus on a specific treatment, were from outside the United States, were of poor quality, or did not report findings from clinical research studies. A more thorough description of the methods used to conduct the medical effectiveness review and the process used to grade the evidence for each outcome measure is presented in Appendix B.

The conclusions below are based on the best available evidence from peer-reviewed and grey literature. 48 Unpublished studies are not reviewed because the results of such studies, if they exist, cannot be obtained within the 60-day timeframe for CHBRP reports.

Key Questions

- What are the effects of deductibles on insulin use/adherence for patients with T1DM, T2DM, or GDM?
- 2. What are the associated effects of deductibles for insulin on health outcomes and utilization?

Methodological Considerations

The primary focus of this review and analysis is on insulin use and adherence related to cost sharing, including deductibles, as related to the bill language. Thus, it does not include adherence for overall diabetes management, for which there are multiple components. Additionally, this bill would apply to patients with T1DM, T2DM, or GDM diagnosis, and there are disease differentiations between the types that inherently affect adherence. It should also be noted that there are several barriers to conducting randomized controlled trials (RCTs) of differential cost sharing on insulin use (i.e. ethical considerations, medical necessity of insulin for treatment of type 1 diabetes, multi-faceted treatment regimens required to effectively treat diabetes), resulting in a literature base that is not as rigorous and thereby limiting the certainty of conclusions drawn from the evidence.

CHBRP did not review the evidence on the effectiveness of insulin for the treatment of diabetes in general, as this has been well documented, and is included in the American Diabetic Association (ADA) treatment guidelines as referenced in the "clinical practice guidelines for diabetes mellitus" section above.

⁴⁸ Grey literature consists of material that is not published commercially or indexed systematically in bibliographic databases. For more information on CHBRP's use of grey literature, visit http://chbrp.com/analysis_methodology/medical_effectiveness_analysis.php.

CHBRP found no studies that specifically addressed the effect of deductibles only on insulin adherence, thus the review below includes the effects of all forms of cost sharing on insulin adherence.

Outcomes Assessed

The primary outcome of interest for the effect of cost sharing on insulin use for patients with diabetes is utilization of insulin, defined as fills after prescription and adherence to prescribed insulin regimens. The associated effect of insulin adherence on health was measured by glycemic control (HbA1c levels), healthcare utilization (e.g., emergency department visits, hospitalizations), productivity (disability, absenteeism) and diabetes-related complications or comorbidities (e.g., amputations, ulcers, blindness, heart attack, stroke). No literature included in the medical effectiveness review examined hyperglycemic events or ketoacidosis events specifically, so while these are common health outcomes associated with diabetes, they are not reflected in these studies.

Study Findings

This section summarizes CHBRP's findings regarding the strength of evidence for the effects of cost sharing on insulin use and adherence for patients with diabetes. Each section is accompanied by a corresponding figure. The title of the figure indicates the test, treatment, or service for which evidence is summarized. The statement in the box above the figure presents CHBRP's conclusion regarding the strength of evidence about the effect of a particular test, treatment, or service based on a specific relevant outcome and the number of studies on which CHBRP's conclusion is based. Definitions of CHBRP's grading scale terms is included in the box below, and more information is included in Appendix B.

CHBRP found a *preponderance of evidence* on the effect of cost sharing on insulin use for diabetes treatment.

The following terms are used to characterize the body of evidence regarding an outcome:

Clear and convincing evidence indicates that there are multiple studies of a treatment and that the large majority of studies are of high quality and consistently find that the treatment is either effective or not effective.

Preponderance of evidence indicates that the majority of the studies reviewed are consistent in their findings that treatment is either effective or not effective.

Limited evidence indicates that the studies have limited generalizability to the population of interest and/or the studies have a fatal flaw in research design or implementation.

Inconclusive evidence indicates that although some studies included in the medical effectiveness review find that a treatment is effective, a similar number of studies of equal quality suggest the treatment is not effective.

Insufficient evidence indicates that there is not enough evidence available to know whether or not a treatment is effective, either because there are too few studies of the treatment or because the available studies are not of high quality. It does not indicate that a treatment is not effective.

More information is available in Appendix B.

Cost Sharing for Prescription Drugs

It is well established in the literature that persons who face higher cost sharing use fewer services than persons with lower cost sharing (CHBRP, 2018). In addition, there is a preponderance of evidence across multiple health conditions that, as cost sharing increases, adherence to drug regimens decreases, with a majority of studies indicating that decreased adherence is associated with worse outcomes (CHBRP, 2014). Goldman et al. (2007) found that for every 10% increase in cost sharing, there was a 2% to 6% decrease in utilization. The results are clear for those with chronic conditions that increased cost sharing is associated with decreased adherence and worse health outcomes (Goldman et al., 2007). Similar results were found in a meta-analysis of publicly insured patients (Sinnott et al., 2013). However, there is also evidence that the effect of cost sharing may differ depending on the specific disease and the specific drug (CHBRP, 2018).

Effect of Cost Sharing on Insulin Use and Adherence for Diabetes Mellitus

CHBRP identified seven studies that examined the effects of cost sharing on insulin use for diabetes treatment. In a cross-sectional survey study by Herkert et al. (2019), the authors analyzed the prevalence of cost-related insulin underuse and its association with glycemic control. The survey was administered at the Yale Diabetic Center to patients with T1DM or T2DM for whom insulin was prescribed in the past 6 months. Cost-related insulin underuse was defined by a "yes" response to any of the six questions: "In the last 12 months did you... (1) use less insulin than prescribed, (2) try to stretch out your insulin, (3) take smaller doses of insulin than prescribed, (4) stop using insulin, (5) not fill an insulin prescription, (6) not start insulin... because of cost." Of 354 eligible patients, 199 completed the survey and 51 (25.5%) reported cost-related underuse. Cost-related insulin underuse did not significantly differ between patients with T1DM and T2DM.

A systematic review by Davies et al. (2013), identified studies reporting factors associated adherence to insulin therapy in adults with T1DM or T2DM. Seventeen studies were identified and two of these studies examined the effects of financial burden on adherence.

The first of these studies was a retrospective pre-post comparison study of a cohort of patients with T1DM and T2DM who switched from a traditional formulary to a value-based insurance design, which reduces or eliminates copayments for highly effective preventive medications (Nair et al., 2009). This involved placing all diabetic drugs and testing supplies on the lowest copay tier for one employer group (n=225) of which 53 patients were receiving insulin. Differences in insulin adherence (proportion of days covered [PDC] ratio) were found to be significant at both year 1 (7.7% increase; p=.0068) and year 2 (7.48%; p=.0251) compared to the pre-period. However, the proportion of adherent patients (defined as \geq 80% PDC ratio) did not significantly change between the three time points and remained at about 20% (20.8% pre-period, 22.6% Y1, 20.8% Y2). It should be noted that these rates of adherent patients include both T1DM and T2DM patients, for which adherence to insulin is known to inherently differ (see "Insulin-Associated Barriers" in the *Background* section).

In the second of these studies, a large (n=20,176) retrospective database study of patients with primarily type 2 diabetes (approximately 90%) who switched to a value-based insurance design was compared to a random control sample of n=190,889 who remained on a traditional tiered formulary design (Chang et al., 2010). They found that adherence to insulin improved over the first year in those in the value-based insurance group, compared to a decline over the same period in the traditional formulary group. They also found the initiation rate for T2DM patients starting on insulin was significantly higher at year one in the value-based insurance group than in the control group.

A 2016 systematic review by Capoccia et al. synthesized the evidence on general medication adherence with prescribed glucose-lowering agents (including insulin and oral anti-diabetics). They identified a total of 98 studies and found cost and copays to significantly affect adherence, among several other factors. Of these, CHBRP identified two retrospective studies that specifically related to cost sharing and insulin adherence as relevant to AB 97.

One was a second retrospective pre-post comparison study by Nair et al. (2010), in which they examined the effects of a value-based insurance design for diabetics within a different employer group. The sample consisted of n=589 patients with T1DM and T2DM, with n=132 of these patients receiving insulin. Differences in mean insulin adherence rates were again found to be significant at both year 1, with a 9.4% increase of baseline mean adherence, and year 2, with a 11.3% increase of baseline mean adherence. Contrary to their first study, they did find significant differences in the percentage of insulin adherent individuals (defined as \geq 80% PDC ratio) from the pre-period to year 1 (22% vs. 30.3%; OR=1.57) and the pre-period to year 2 (22% vs. 33.3%; OR=1.80).

In the second, a retrospective, cross-sectional study by Gibson et al. (2010) assessed the relationship between cost sharing and adherence to medications in patients with T2DM. This study combined insulin and oral antidiabetic medications (OAD) in their examination of adherence rates to prescribed regimens and did not analyze results by insulin alone. The analysis included 96,734 patients on a combination of OADs and insulin with employer-sponsored insurance in the 2003–2006 MarketScan Database. They reported that an increase from \$10 to \$20 in the cost-sharing index resulted in an average 4.8% reduction in adherence (defined as \geq 80% PDC ratio).

A 2021 retrospective analysis by Chandra et al., examined the effects of cost sharing on prescription drug utilization in a 20% random sample of all Medicare Part D enrollees from 2007-2012. After applying their inclusion criteria, their total analytic sample was 358,706 beneficiaries across a variety of health conditions. Approximately 16% (n=57,392) of the sample was observed to have a diagnosis of T1DM or T2DM. Using predictive modeling strategies based on changes in percent coverage from Medicare's drug benefit structure, the authors estimated changes in prescriptions filled for specific drug classes. They found that for every one percent increase in coinsurance costs for diabetes drugs that lower blood sugar, including insulin and other OADs, patients made .00288 fewer diabetes prescription fills. This translates to approximately a 3% decrease in adherence to insulin and other OADs for every 10% increase in cost-sharing expense.

In another retrospective analysis of Medicare Part D beneficiaries, Trish et al. (2021) examined the association between out-of-pocket spending and insulin adherence. Specifically, the authors sought to assess the potential outcomes associates with the newly announced Medicare program to limit out-ofpocket spending on insulin to \$35 per month (refer to the *Policy Context* section for more information on the Senior Savings Model). To analyze the potential outcomes of this cost sharing limit, changes in insulin use (defined as change in PDC ratio) were compared between beneficiaries in individual plans and those in an employer group-waiver plan across three coverage phases. 49 The study used a 100% sample of Medicare Part D claims from 2018, which included a total sample of 474,929 individuals with a prescription for insulin with n=303,616 in individual plans and n=171,313 in employer plans. The mean cost sharing for insulin per 30-day supply among individual plan enrollees in the initial coverage phase was \$50.57, \$117 in the coverage gap, and \$36.86 in catastrophic coverage. The mean cost sharing among employer plan enrollees across the same phases was \$32.73 (initial), \$31.99 (coverage gap), and \$19.73 (catastrophic coverage). They found that beneficiaries in individual plans who were in the coverage gap reduced their insulin use by 5.4% (PDC=67.5% during initial coverage vs. PDC=62.1% in the coverage gap). Comparatively, beneficiaries in employer plans increased their insulin use by 2.8% in the coverage gap (PDC=70.1% during initial coverage vs. PDC=72.9% in the coverage gap). Compared to the initial coverage phase, there was no change in insulin use for individual plan enrollees who ended the year in catastrophic coverage and a 2.4% increase for employer plan enrollees.

Summary of findings regarding cost sharing on insulin use and adherence: There is a *preponderance of evidence* from one cross-sectional self-report study, one retrospective cross-sectional study, three retrospective pre-post studies, and 2 retrospective cohort studies on cost-related insulin use/adherence that cost sharing affects insulin use and adherence in patients with diabetes; higher cost sharing reduces adherence and lower cost sharing increases adherence. Some of the limiting factors that

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⁴⁹ These included (1) The initial coverage phase (2) The coverage gap phase and (3) The catastrophic coverage phase

contributed to this evidence grading are the quality of the studies, the inherent differences between the types of diabetes conditions, and confounding adherence issues (i.e., insulin side effects, fear of injection, social factors, health literacy).

Figure 4. Effect of Cost Sharing for Insulin Use & Adherence



Effect of Cost Sharing for Insulin on Health Outcomes and Utilization

CHBRP identified no studies that examined the effects of cost sharing for insulin alone on diabetes-related health outcomes. Four of the studies discussed above reported health outcome and utilization results, though these findings are not specific to insulin alone, and include the effect of cost for insulin, other OADs, and diabetic testing supplies. These findings are discussed in this section to provide the available evidence on cost sharing for insulin and the related health outcomes.

The Herkert et al. (2019) cross-sectional survey study found that patients who reported cost-related insulin underuse, compared to those who did not, were significantly more likely to have poor glycemic control (p=.03). Poor glycemic control was defined has HbA1c \geq 9% collected at time of the visit or within 3 months.

The Nair et al. (2009) retrospective pre-post comparison study of switching to a value-based insurance design examined changes in medical utilization at each of the three time points (pre-period, year 1, year 2). The authors reported a 25% decrease in diabetes-specific emergency department visits and a 20% decrease in hospitalizations in year 2 compared to year 1, though these comparisons were not found to be statistically significant. It should be noted that these outcomes included the entire sample of patients with diabetes, not only those patients using insulin.

The 2010 study by Nair et al. also examined the effects of switching to a value-based insurance design, within a different employer group than the 2009 study, and reported on diabetes-related medical utilization effects for the entire sample at each of the three time points (pre-period, year 1, year 2). The authors reported a 12% decrease in diabetes-specific office visits, a 31% decrease in emergency room visits, and a 53% decrease in hospitalizations in year 1 compared to the pre-period. However, only the comparisons for office visits and emergency department visits from the pre-period to year 1 were found to be statistically significant. These effects include the entire sample of patients, not only those on insulin, and the associated effects of the lowered cost of diabetic testing supplies and other diabetic drugs should also be considered.

The Gibson et al. (2010) retrospective cross-sectional study also assessed the relationship between cost sharing for diabetes medications and the associated health outcomes in patients with T2DM that resulted from improved adherence. They examined the relationship between improved adherence to the prescribed diabetes treatment regimen (OADs with and without insulin) and health outcomes and found significant reductions in long term complications, emergency department visits and hospitalizations. However, number of physician visits (non-ED visits) were higher among adherent patients. For measures of productivity and quality of life, they also reported that the number of short-term disability days was significantly lower for adherent patients, but found no significant difference in absenteeism.

Summary of findings regarding cost sharing for insulin on health outcomes and utilization: There is *insufficient evidence* on the effect of cost sharing for insulin on diabetes-related health outcomes and

utilization. Though the studies presented in the above section provide some evidence on health and utilization outcomes, these findings were not specific to insulin alone, but to patients on insulin and other OADs. Additional limiting factors that contributed to this evidence grading are the quality of studies, the inability to separate outcomes based on type of diabetes, confounding variables (i.e., lowered cost of testing supplies), and the multifaceted nature of diabetes treatment. A grading of insufficient evidence does not indicate that there is no effect, but rather means that the effect is unknown.

Figure 3. Effect of Cost Sharing for Insulin on Health Outcomes & Utilization



Summary of Findings

CHBRP found a *preponderance of evidence* from seven cross-sectional and retrospective studies on cost-related insulin use/adherence that cost sharing affects insulin use and adherence in patients with diabetes. Though no studies specifically examined the effect of deductibles only, these studies provided a *preponderance of evidence* that higher cost sharing reduces adherence to insulin and lower cost sharing increases adherence to insulin. CHBRP found *insufficient evidence* on the associated effect of cost sharing for insulin on diabetes-related health outcomes, including HbA1c levels, outpatient visits, emergency department visits, hospitalizations, long-term complications, and disability/absenteeism rates. Though the studies presented did report on these health and utilization outcomes, the findings were not specific to the effect of insulin alone, but combined with use of other OADs and testing supplies. There were several limitations that contributed to the gradings provided in this review, most notably the inherent differences between the types of diabetes conditions and the multifaceted nature of diabetes treatment, resulting in a literature base that is not as rigorous and thereby limiting the certainty of conclusions drawn from the evidence.

BENEFIT COVERAGE, UTILIZATION, AND COST IMPACTS

As discussed in the *Policy Context* section, AB 97 would prohibit commercial and CalPERS DMHC-regulated plans and CDI-regulated policies from applying a deductible to insulin prescriptions. Other cost sharing (copayments, coinsurance) would still be permitted.

This section reports the potential incremental impacts of AB 97 on estimated baseline benefit coverage, utilization, and overall cost. This analysis makes the following assumptions:

- The population subject to AB 97 includes individuals covered by DMHC-regulated commercial insurance plans, CDI-regulated policies, and publicly funded plans (including CalPERS) subject to the requirements of the Knox-Keene Health Care Service Plan Act. Based on DMHC and DHCS guidance, Medi-Cal managed care enrollees are not subject to AB 97 since pharmacy benefit is carved out from DMHC-regulated plans.
- CHBRP assumes the insulin products available in Milliman's 2019 Consolidated Health Cost
 Guidelines Sources Database (CHSD) that was used for this analysis will continue to be available
 in 2022. These claims data do not incorporate insulins newly available between 2019 and 2021.
 CHBRP is unable to predict the number, type, or price of new insulin products that may come to
 the market in 2022, nor how new products might affect the price and cost sharing for existing
 products.
- The estimated changes in cost sharing reported here include deductible amounts incurred by enrollees in plans where deductible amounts must be reached (e.g., high deductible health plans [HDHPs], Bronze and Silver plans offered through Covered California). More information about enrollees with deductibles is presented in Table 2 in the *Policy Context* section.

For further details on the underlying data sources and methods used in this analysis, please see Appendix C.

Baseline and Postmandate Benefit Coverage

At baseline, 100% of the 13,945,000 enrollees in commercial and CalPERS DMHC-regulated plans and CDI-regulated policies would be subject to AB 97. The 13,945,000 enrollees in DMHC-regulated plans and CDI-regulated policies make up 64% of all enrollees subject to state-level benefit mandates and excludes enrollees in DMHC-regulated Medi-Cal managed care plans.

CHBRP estimates at baseline there are 118,014 enrollees who use insulin in DMHC-regulated plans and CDI-regulated policies, where 81,265 enrollees using insulin do not have a deductible (69%). CHBRP estimates 36,750 enrollees using insulin have a deductible (31%) (see estimates in Table 1). Postmandate, 100% of enrollees would not need to meet their deductible before paying the normal copayment or coinsurance for their insulin prescription.

A majority— over 93% - of enrollees in commercial and CalPERS plans and policies regulated by DMHC or CDI have a pharmacy benefit regulated by DMHC or CDI that covers both generic and brand-name outpatient prescription medications. Approximately 3.1% do not have a pharmacy benefit and 3.6% have a pharmacy benefit that is not regulated by DMHC or CDI. Because AB 97 does not require creation of a pharmacy benefit — only compliant benefit coverage when a pharmacy benefit is present — baseline benefit coverage for enrollees without a pharmacy benefit or whose pharmacy benefit is not regulated by DMHC or CDI is compliant.

Baseline and Postmandate Utilization

Using relevant codes from the International Classification of Diseases, 10th Revision, Clinical Modification (ICD-10-CM), and National Drug Codes (NDCs), CHBRP used data from Milliman's 2019 Consolidated

Health Cost Guidelines Sources Database (CHSD) to develop baseline estimates of utilization of insulin. CHBRP calculated utilization rates for enrollees whose claims for insulin exceed the cost-sharing cap at baseline and for those who did not exceed the cap. See estimates in Table 1. Utilization (measured as number of 30-day supply insulin prescriptions per month per user) is the same (0.85) for all enrollees at baseline. Postmandate, the group of enrollees with a deductible at baseline would experience an increase in utilization because this group would experience a decrease in cost sharing due to the bill.

To estimate changes in utilization postmandate, CHBRP applied an estimate of price elasticity of demand to enrollees with a deductible at baseline. CHBRP assumes reduced cost sharing for insulin increases the utilization of outpatient prescription insulin based on literature that establishes evidence of price elasticity of demand for prescription drugs (Goldman et al., 2004).

There is limited literature on the price elasticity of demand for insulin specifically; recent studies examining the effect of value-based insurance design (VBID) on insulin use also include oral antidiabetic (OAD) medications in the impacts of cost sharing. Because these OADs are in a different medication class than insulin, they may impact the elasticity measure due to different cost sharing levels for that drug class. Because of this, CHBRP bases the estimate of price elasticity on a Goldman et al. (2004) article that found use of insulin specifically decreased by 8% when copayments doubled. Thus, CHBRP applied this elasticity estimate to calculate increase in insulin utilization postmandate for enrollees who would experience a decrease in cost sharing postmandate.

As shown in Table 1, CHBRP estimates a 3% reduction in cost sharing for those enrollees who have a deductible at baseline, and therefore estimates a 0.26% increase in utilization of insulin postmandate for those enrollees. Enrollees using insulin at baseline who have a deductible tend to be users of other high-cost medications and other medical services. As a result, almost all enrollees would hit their deductible or out of pocket maximum within a plan year, regardless of whether insulin is subject to the deductible.

Baseline and Postmandate Per-Unit Cost

The average cost of insulin per prescription per month is \$491. Using 2019 CHSD data, per-unit cost is calculated based on the allowed costs and is trended to 2022; the per-unit cost is not reduced by potential rebates that may be received by the health plans. AB 97 would not change the unit or per-prescription cost for insulin.

Baseline and Postmandate Expenditures

Table 77 and Table 88 present baseline and postmandate expenditures by market segment for DMHC-regulated plans and CDI-regulated policies. The tables present per member per month (PMPM) premiums, enrollee expenses for both covered and noncovered benefits, and total expenditures (premiums as well as enrollee expenses).

AB 97 would increase total net annual expenditures by \$10,162,000 or 0.008% for enrollees with DMHC-regulated plans and CDI-regulated policies. This is due to an increase in \$23,853,000 in total health insurance premiums paid by employers and enrollees, adjusted by a \$13,691,000 decrease in enrollee expenses.

Premiums

Total premiums for private employers purchasing group health insurance would increase by \$8,574,000, or 0.02%. Total premiums for purchasers of individual market health insurance would increase by \$11,802,000, or 0.07%. Changes in premiums as a result of AB 97 would vary by market segment. Note that such changes are related to the number of enrollees (see Table 1, Table 7, and Table 8), with health insurance that would be subject to AB 97. The greatest change in premiums as a result of AB 97 is for

small-group policies (0.12% increase) and individual policies (0.11% increase) in the CDI-regulated market.

Among publicly funded plans, DMHC-regulated Medi-Cal managed care is not subject to AB 97. For CalPERS HMO enrollees, the impact on premiums is \$0 because these plans do not have pharmacy deductibles.

Enrollee Out-of-Pocket Expenses

AB 97–related changes in enrollee expenses for covered benefits (deductibles, copays, coinsurance, etc.) and enrollee expenses for noncovered benefits would vary by market segment. Note that such changes are related to the number of enrollees (see Table 1, Table 7, and Table 8) with health insurance that would be subject to AB 97 that are expected to use insulin during the year after enactment.

The largest reduction in enrollee cost sharing due to AB 97 would be for small-group policies in the CDI-regulated market, with reductions of approximately \$0.43 per member per month.

Average enrollee cost-sharing expenses per user

For baseline insulin users, AB 97's prohibition of insulin being subject to the deductible only impacts those enrollees who have deductibles. Overall, 31% of enrollees who use insulin at baseline have deductibles. For enrollees who do not have a deductible at baseline, the average cost sharing for insulin is \$49 per month, and for those enrollees for whom insulin is subject to the deductible at baseline, the average cost sharing for insulin is \$89 per month (Table 1). On average, cost sharing would decrease by \$2 per month for enrollees who have deductibles, postmandate. A majority of these enrollees also have other prescription drug and medical costs that would cause them to meet their deductible or out-of-pocket maximum in a given year. Among enrollees with a pharmacy deductible, 64% have expenditures for other non-insulin brand name prescription medications that exceed \$500 annually, and therefore would cause them to meet their pharmacy deductible (Table 6). Among enrollees in health savings account (HSA)-eligible high deductible health plans (HDHPs) (and therefore with a combined medical and pharmacy deductible), almost three quarters (70%) of enrollees have expenditures for medical care and non-insulin brand name prescription medications that exceed \$2,500 annually. Due to this, there is a 3% decrease in average cost sharing as a result of AB 97.

Table 6. Annual non-insulin brand name prescription medication and medical care spending for enrollees using insulin

	Percent of Enrollees using Insulin with spending by category
Annual Non-Insulin Brand Name Prescription Medication Spending (a)	
\$0 - \$250	28%
\$250 - \$500	8%
\$500+	64%
Annual Medical Care and Non-Insulin Brand Name Prescription Medication Spending (b)	
\$0 - \$1,500	23%
\$1,500 - \$2,500	7%
\$2,500+	70%

Source: California Health Benefits Review Program, 2021.

Note: The table above was calculated using total medical and prescription drug allowed amounts for insulin-using enrollees in the

2019 CHSD database. Insulin claims and generic drug claims were excluded from the totals. The total annual spend was calculated for each enrollee individually, so that each enrollee was assigned to a range band of non-insulin brand drug or medical and non-insulin brand drug spend.

- (a) Among enrollees with a pharmacy deductible.
- (b) Among enrollees enrolled in HSA-qualified HDHPs, and therefore with a combined medical and pharmacy deductible.

Among enrollees with a deductible who experience a decrease in cost sharing of greater than the average of \$2:

- 9.5% of insulin users would experience a reduction of more than \$20:
- 7.7% of insulin users would experience a reduction of more than \$100:
- 3.3% of insulin users would experience a reduction of more than \$250; and
- 1.5% of insulin users would experience a reduction of more than \$500.

It is possible that some enrollees who had deferred insulin treatment due to cost could begin using insulin postmandate; thus, this group of enrollees would incur cost sharing postmandate where they did not have cost sharing at baseline. However, this group is estimated to be relatively small. Per CHBRP's content expert, forgoing insulin completely after a physician has prescribed it is something that will occur among only those with type 2 diabetes mellitus (T2DM) where symptoms or the clinical consequences of not having the insulin are not felt by the patient. Literature suggests approximately 2.5% of people who were prescribed insulin never started their prescription in the past year due to cost. ⁵⁰ Thus, for some enrollees, cost sharing may be the sole barrier to filling their insulin prescription; however, it is not known what the baseline cost sharing is for this group if they did fill their prescription (i.e., what proportion of non-users have deductibles and what the plan benefit design is regarding copayments or coinsurance), nor is it known what cost-sharing threshold would stimulate utilization among these enrollees. While CHBRP expects some demand response from this group when cost sharing is lowered postmandate, CHBRP expects it would be a relatively low utilization increase that would not substantially change the results of this analysis.

The enrollees most likely to experience the greatest cost sharing reductions postmandate are those who are enrolled in plans that require significant deductibles to be met before coinsurance is applied to the insulin purchase, e.g., HDHPs, Bronze, and Silver plans. CHBRP's cost model estimates indicate that for enrollees subject to AB 97, approximately 19% of large-group, 37% of small-group, and 61% of individual market enrollees are in plans or policies with deductibles that apply to prescription drugs, where deductibles may have a material impact on insulin cost sharing. However, as mentioned previously, most enrollees who use insulin are likely to meet their deductible and out of pocket maximum regardless.

It is possible that at baseline some enrollees incurred insulin-related expenses when coverage was denied, delivered through another vendor or purchased outside of the health insurance plan, but CHBRP cannot estimate the frequency with which such situations occur and so cannot offer a calculation of impact.

Potential Cost Offsets or Savings in the First 12 Months After Enactment

Due to the low change in utilization of insulin, no offsets are projected as a result of AB 97.

Postmandate Administrative Expenses and Other Expenses

CHBRP estimates that the increase in administrative costs of DMHC-regulated plans and/or CDI-regulated policies would remain proportional to the increase in premiums. CHBRP assumes that if health care costs increase as a result of increased utilization or changes in unit costs, there is a corresponding

⁵⁰ Personal communication with corresponding author of Herkert et al., 2019, on March 10, 2020

proportional increase in administrative costs. CHBRP assumes that the administrative cost portion of premiums is unchanged. All health plans and insurers include a component for administration and profit in their premiums.

Other Considerations for Policymakers

In addition to the impacts a bill may have on benefit coverage, utilization, and cost, related considerations for policymakers are discussed below.

Postmandate Changes in the Number of Uninsured Persons

Because the change in average premiums does not exceed 1% for any market segment (see Table 1, Table 7, and Table 8), CHBRP would expect no measurable change in the number of uninsured persons due to the enactment of AB 97.

Changes in Public Program Enrollment

CHBRP estimates that the mandate would produce no measurable impact on enrollment in publicly funded insurance programs due to the enactment of AB 97.

How Lack of Benefit Coverage Results in Cost Shifts to Other Payers

Enrollees may take part in cost sharing assistance programs to help offset high copayments or coinsurance. CHBRP is unable to provide a quantifiable estimate of the number of enrollees who take part in patient assistance programs and the potential impact AB 97 would have on the number of enrollees who use these programs.

Table 7. Baseline Per Member Per Month Premiums and Total Expenditures by Market Segment, California, 2022

	DMHC-Regulated							CDI-Regulated			
	Privately Funded Plans (by Market) (a)			Publicly Funded Plans			Privately Funded Plans (by Market) (a)				
	Large Group	Small Group	Individual	CalPERS HMOs (b)	MCMC (Under 65) (c)	MCMC (65+) (c)	Large Group	Small Group	Individual	Total	
Enrollee counts											
Total enrollees in plans/policies subject to state mandates (d)	8,405,000	2,086,000	1,989,000	889,000	7,218,000	787,000	384,000	43,000	144,000	21,945,000	
Total enrollees in plans/policies subject to AB 97	8,405,000	2,086,000	1,989,000	889,000	0	0	384,000	43,470	144,000	21,945,470	
Premiums											
Average portion of premium paid by employer	\$426.28	\$374.49	\$0.00	\$540.40	\$226.61	\$478.87	\$530.80	\$421.81	\$0.00	\$84,948,349,000	
Average portion of premium paid by employee	\$141.02	\$180.89	\$624.47	\$96.86	\$0.00	\$0.00	\$186.55	\$212.07	\$545.57	\$36,600,954,000	
Total premium	\$567.30	\$555.38	\$624.47	\$637.27	\$226.61	\$478.87	\$717.35	\$633.88	\$545.57	\$121,549,303,000	
Enrollee expenses											
For covered benefits (deductibles, copays, etc.)	\$43.61	\$121.70	\$173.51	\$50.75	\$0.00	\$0.00	\$134.75	\$197.13	\$184.11	\$13,168,032,000	
For noncovered benefits (e)	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0.00	\$0	
Total expenditures	\$610.91	\$677.07	\$797.97	\$688.02	\$226.61	\$478.87	\$852.10	\$831.01	\$729.68	\$134,717,335,000	

Source: California Health Benefits Review Program, 2021.

Notes: (a) Includes enrollees with grandfathered and nongrandfathered health insurance acquired outside or through Covered California (the state's health insurance marketplace).

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⁽b) Approximately 54.1% of CalPERS enrollees in DMHC-regulated plans are state retirees, state employees, or their dependents. About one in five (20.5%) of these enrollees has a pharmacy benefit not subject to DMHC. CHBRP has projected no impact for those enrollees. However, CalPERS could, postmandate, require equivalent coverage for all its members (which could increase the total impact on CalPERS).

⁽c) Medi-Cal Managed Care Plan expenditures for members over 65 include those who are also Medicare beneficiaries. This population does not include enrollees in COHS.

- (d) Enrollees in plans and policies regulated by DMHC or CDI aged 0 to 64 years as well as enrollees 65 years or older in employer-sponsored health insurance. This group includes commercial enrollees (including those associated with Covered California or CalPERS) and Medi-Cal beneficiaries enrolled in DMHC-regulated plans.
- (e) Includes only those expenses that are paid directly by enrollees or other sources to providers for services related to the mandated benefit that are not currently covered by insurance. This only includes those expenses that will be newly covered, postmandate. Other components of expenditures in this table include all health care services covered by insurance.

Key: CalPERS HMOs = California Public Employees' Retirement System Health Maintenance Organizations; CDI = California Department of Insurance; COHS = County Organized Health Systems; DMHC = Department of Managed Health Care; MCMC = Medi-Cal Managed Care.

Table 8. Postmandate Per Member Per Month Premiums and Total Expenditures by Market Segment, California, 2022

	DMHC-Regulated							CDI-Regulated			
	Privately Funded Plans (by Market) (a)			Publicly Funded Plans			Privately Funded Plans (by Market) (a)				
	Large Group	Small Group	Individual	CalPERS HMOs (b)	MCMC (Under 65) (c)	MCMC (65+) (c)	Large Group	Small Group	Individual	Total	
Enrollee counts								·			
Total enrollees in plans/policies subject to state mandates (d)	8,405,000	2,086,000	1,989,000	889,000	7,218,000	787,000	384,000	43,000	144,000	21,945,000	
Total enrollees in plans/policies subject to AB 97	8,405,000	2,086,000	1,989,000	889,000	7,218,000	787,000	384,000	43,470	144,000	21,945,470	
Premiums	, ,	· ·	, ,	,		,	,	<u> </u>	,		
Average portion of premium paid by employer	\$0.0335	\$0.1500	\$0.0000	\$0.0000	\$0.0000	\$0.0000	\$0.2560	\$0.4996	\$0.0000	\$8,574,000	
Average portion of premium paid by employee	\$0.0111	\$0.0725	\$0.4526	\$0.0000	\$0.0000	\$0.0000	\$0.0900	\$0.2512	\$0.5773	\$15,278,000	
Total premium	\$0.0446	\$0.2225	\$0.4526	\$0.0000	\$0.0000	\$0.0000	\$0.3460	\$0.7507	\$0.5773	\$23,853,000	
Enrollee expenses											
For covered benefits (deductibles, copays, etc.)	-\$0.0229	-\$0.1245	-\$0.2704	\$0.0000	\$0.0000	\$0.0000	-\$0.2167	-\$0.4329	-\$0.3422	-\$13,691,000	
For noncovered benefits (e)	\$0.0000	\$0.0000	\$0.0000	\$0.0000	\$0.0000	\$0.0000	\$0.0000	\$0.0000	\$0.0000	\$0	
Total expenditures	\$0.0217	\$0.0980	\$0.1822	\$0.0000	\$0.0000	\$0.0000	\$0.1293	\$0.3179	\$0.2351	\$10,161,000	
Percent change											
Premiums	0.0079%	0.0401%	0.0725%	0.0000%	0.0000%	0.0000%	0.0482%	0.1184%	0.1058%	0.0196%	
Total expenditures	0.0036%	0.0145%	0.0228%	0.0000%	0.0000%	0.0000%	0.0152%	0.0383%	0.0322%	0.0075%	

Source: California Health Benefits Review Program, 2021.

Notes: (a) Includes enrollees with grandfathered and nongrandfathered health insurance acquired outside or through Covered California (the state's health insurance marketplace).

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- (b) Approximately 54.1% of CalPERS enrollees in DMHC-regulated plans are state retirees, state employees, or their dependents. About one in five (20.5%) of these enrollees has a pharmacy benefit not subject to DMHC. CHBRP has projected no impact for those enrollees. However, CalPERS could, postmandate, require equivalent coverage for all its members (which could increase the total impact on CalPERS).
- (c) Medi-Cal Managed Care Plan expenditures for members over 65 include those who are also Medicare beneficiaries. This population does not include enrollees in COHS.
- (d) Enrollees in plans and policies regulated by DMHC or CDI aged 0 to 64 years as well as enrollees 65 years or older in employer-sponsored health insurance. This group includes commercial enrollees (including those associated with Covered California or CalPERS) and Medi-Cal beneficiaries enrolled in DMHC-regulated plans.
- (e) Includes only those expenses that are paid directly by enrollees or other sources to providers for services related to the mandated benefit that are not currently covered by insurance. This only includes those expenses that will be newly covered, postmandate. Other components of expenditures in this table include all health care services covered by insurance.

Key: CalPERS HMOs = California Public Employees' Retirement System Health Maintenance Organizations; CDI = California Department of Insurance; COHS = County Organized Health Systems; DMHC = Department of Managed Health Care; MCMC = Medi-Cal Managed Care.

PUBLIC HEALTH IMPACTS

As discussed in the *Policy Context* section, AB 97 would prohibit a deductible from being applied to insulin prescriptions. Other cost sharing (copayments, coinsurance) would still be permitted. The public health impact analysis includes estimated impacts in the short term (within 12 months of implementation) and in the long term (beyond the first 12 months postmandate).

Estimated Public Health Outcomes

Measurable health outcomes relevant to AB 97 included utilization of insulin and the associated effects of insulin adherence on health as measured by glycemic control (HbA1c levels), healthcare utilization (e.g., emergency department visits, hospitalizations), productivity (disability, absenteeism), and diabetes-related complications or comorbidities (e.g., amputations, ulcers, blindness, heart attack, stroke). As presented in the *Medical Effectiveness* section, there is a *preponderance of evidence* in the literature that cost sharing affects insulin use and adherence in patients with diabetes, and insufficient evidence on the effect of cost sharing for insulin on diabetes-related health and utilization outcomes listed above.

As presented in the *Benefit Coverage, Utilization, and Cost Impacts* section, 36,750 enrollees who use insulin and have a deductible would experience an average of a 3% reduction in cost sharing, reducing average monthly cost sharing from \$89 to \$876. Almost 10% of enrollees who use insulin would experience a decrease in cost sharing of more than \$20.

The segment of the insured population most impacted by AB 97 would be enrollees for whom a deductible applies before the copay, or for enrollees with high-deductible plans, which require the enrollee to pay list price for insulin until the deductible is met for the year. Also affected are enrollees with diabetes who are prescribed more than one type of insulin or a higher-tiered insulin (Cefalu et al., 2018). Enrollees with type 2 diabetes mellitus (T2DM) are more likely than those with type 1 diabetes mellitus (T1DM) to increase utilization owing to the inability of patients with T1DM to limit insulin intake without adverse effects on their health. However, the majority of these enrollees have prescription drug or other medical utilization that causes them to reach the annual deductible and out of pocket maximum.

In the first year postmandate, 36,750 enrollees who use insulin and have a deductible at baseline would experience a small reduction in cost sharing. CHBRP projects that as a result, there would be a 0.2% increase in utilization of insulin. CHBRP found a *preponderance of evidence* that cost sharing for insulin is effective in improving adherence to insulin in patients with diabetes, and *insufficient evidence* on the effect of cost sharing for diabetes-related health outcomes. Due to the small decrease in cost sharing and small increase in utilization, CHBRP projects no measurable public health impact. However, at the person-level, for enrollees who would not otherwise meet their deductible or out of pocket maximum and would therefore experience a higher change in cost sharing, AB 97 may result in improved glycemic control, a reduction in healthcare utilization, a reduction in long-term complications attributable to DM, and improved quality of life for enrollees that experience a decrease in cost sharing and improved insulin adherence, or begin using insulin due to reduced costs.

Glycemic Control

For the population that would be impacted by AB 97, achieving stable blood glucose levels, measured as HbA1c, could reduce the frequency and severity of episodes of hyperglycemia and hypoglycemia. In the most severe cases, hyperglycemia can lead to ketoacidosis, followed by coma or death. Similarly, escalation of hypoglycemia can lead to cognitive dysfunction, seizures, coma, and death. Additionally, hypoglycemia unawareness occurs more frequently among those who are insulin dependent (Martin-Timon and Canizo-Gomez, 2015). Therefore, achievement of more stable HbA1c levels through increased utilization and adherence to insulin could avoid these serious health consequences associated with diabetes.

Healthcare Utilization

For the population that would be impacted by AB 97, impacts to healthcare utilization may include reduced hospitalizations and outpatient appointments, and measurable offsets from reductions in insulin-related ED visits. This would reduce costly emergency services and also have direct impacts on the patient. Reduced time in hospitals and EDs also reduces the exposure to hospital-acquired infections and infectious diseases that are prevalent in these settings. This may be a considerable positive health outcome for patients with diabetes who have a compromised immune system and possible other comorbidities, should these reductions in utilization materialize.

Long-Term Complications

Time spent in hyperglycemia and frequency and severity of hyper- and hypoglycemia over a lifetime are associated with serious morbidity and mortality outcomes. In the United States, diabetes is the leading cause of blindness, amputations, and kidney failure, and a key contributor to stroke, heart disease, dental disease, nerve damage, and premature death. To the extent that AB 97 can help individuals taking insulin afford their prescribed dose, it is possible that rates of these comorbid conditions attributable to diabetes could be reduced.

Quality of Life

CHBRP found no literature specifically addressing the impact of reduced cost sharing for insulin on health-related quality of life. However, quality-of-life improvements have been evaluated with regards to outcomes associated with AB 97. In one cross-sectional study, insulin utilization was found to be positively associated with quality of life: significant differences were observed for T2DM insulin users for diet, monitoring, disease-specific knowledge, and adherence to treatment as compared to oral antidiabetic medications (OAD) users (Gillani, 2019). Additionally, Hajós and colleagues (2012) found improvements in quality-of-life scores with improved HbA1c levels due to optimized insulin therapy for those with T2DM who had suboptimal glycemic control (Hajós et al., 2012). There is also evidence that quality of life in patients with diabetes is affected more so by the presence of complications, and not necessarily by the diagnosis itself (Venkataraman, 2013). Peripheral neuropathy was the complication most strongly associated with reduced quality of life (Venkataraman, 2013).

Impact on Disparities⁵¹

Insurance benefit mandates that bring more state-regulated plans and policies to parity may change an existing disparity. As described in the *Error! Reference source not found.* section, disparities in d iabetes exist by race/ethnicity, age, gender, education, income, and health literacy. CHBRP did not find evidence indicating differential use of insulin by any reported disparity within the first 12 months postmandate; therefore, it is projected that AB 97 would have no impact on these diabetes disparities statewide (for a discussion of potential impacts beyond the first 12 months of implementation [including SDOH], see *Long-Term Impacts*). For enrollees who have cost-related barriers to insulin use, AB 97 could improve disparities related to income by reducing the allowed cost-sharing amounts. However, it is worth noting that reduced cost sharing generally shifts the cost to premiums for all enrollees, and this shift could impact lower income enrollees disproportionately.

Despite AB 97 applying only to privately insured enrollees, AB 97 would not exacerbate racial or ethnic disparities due to differences in populations represented in private insurance and Medi-Cal, as Medi-Cal beneficiaries do not have cost sharing.

⁵¹ For details about CHBRP's methodological approach to analyzing disparities, see the *Benefit Mandate Structure* and *Unequal Racial/Ethnic Health Impact*s document here: http://chbrp.com/analysis_methodology/public_health_impact_analysis.php.

LONG-TERM IMPACTS

In this section, CHBRP estimates the long-term impact of AB 97, which CHBRP defines as impacts occurring beyond the first 12 months after implementation. These estimates are qualitative and based on the existing evidence available in the literature. CHBRP does not provide quantitative estimates of longterm impacts because of unknown improvements in clinical care, changes in prices, implementation of other complementary or conflicting policies, and other unexpected factors.

Long-Term Utilization and Cost Impacts

Utilization Impacts

CHBRP estimates annual insulin utilization after the initial 12 months from the enactment of AB 97 would likely stay similar to utilization estimates during the first 12 months postmandate. Utilization changes may occur if new diabetes products or medications change the landscape of insulin use for enrollees with diabetes, however CHBRP is unable to predict these types of changes. Similarly, health care utilization due to improved diabetes management may change in the long term. Reductions in significant complications or comorbidities may take years to develop, but are not trivial.

Cost Impacts

CHBRP estimates cost after the initial 12 months from the enactment of AB 97 are likely to remain similar in the subsequent years. While potential improvements in health outcomes due to better glycemic control among enrollees with diabetes may result in offsets in the future, the small change in utilization of insulin is unlikely to result in substantial cost savings. CHBRP is unable to estimate these changes quantitatively due to the lack of data on long-term utilization and cost due to improved insulin adherence.

Long-Term Public Health Impacts

Some interventions in proposed mandates provide immediate measurable impacts (e.g., maternity service coverage or acute care treatments), whereas other interventions may take years to make a measurable impact (e.g., coverage for tobacco cessation or vaccinations). When possible, CHBRP estimates the longterm effects (beyond 12 months postmandate) to the public's health that would be attributable to the mandate, including impacts on social determinants of health, premature death, and economic loss.

CHBRP estimates that some of the outcomes discussed may take longer than 12 months to observe. Specifically, reductions in significant complications or comorbidities may take years to develop, as would significant differences in disability and absenteeism. AB 97 is unlikely to impact these public health outcomes statewide, but at a person-level it could make a substantial difference in long-term healthcare spending, morbidity, and mortality.

Impacts on Disparities and the Social Determinants of Health⁵²

In the case of AB 97, evidence shows that although variances in education, income, and health literacy exist for the population with diabetes mellitus (DM) and contribute to differences in insulin adherence, CHBRP projects no statewide changes in these social determinants of health (SDOH) that would be attributable to AB 97. However, it is possible that at the person-level, a reduction in cost sharing for insulin therapy could reduce differences in adherence due to income and socioeconomic status.

http://chbrp.com/analysis_methodology/public_health_impact_analysis.php.

⁵² For more information about SDOH, see CHBRP's publication Incorporating Relevant Social Determinants of Health Into CHBRP Benefit Mandate Analyses at

In the long term, CHBRP estimates that AB 97 could improve disparities related to income for some enrollees who have cost-related barriers to insulin use and experience a substantial decrease in cost sharing for insulin. CHBRP is unable to estimate reductions in existing disparities. However, because the prevalence of diabetes is higher for Blacks than for Whites, and there is evidence that cost-related medication nonadherence is also more associated with Blacks, it is possible that this disparity may be reduced for the population AB 97 impacts.

Impacts on Premature Death and Economic Loss

Premature death is often defined as death occurring before the age of 75 years (NCI, 2019).⁵³ In California, it is estimated that there were nearly 5,300 years of potential life lost (YPLL) per 100,000 population each year between 2015 and 2017 (CDPH, 2019; County Health Rankings, 2019).54

Diabetes contributes significantly to premature death and economic loss in California. In addition to complications from diabetes, hypoglycemia is prevalent among those with T1DM and contributes to increased risk of death from diabetes (McCoy, et al., 2012). In addition, diabetes is the seventh leading cause of death in California, and an overall contributor to premature death (e.g., people with diabetes aged 50 years or older die almost 8 years earlier than those without diabetes) (Conroy et al., 2014). The CDC reports that almost 6,000 Californians with diabetes died prematurely in 2013. Despite the diabetes mortality rate decreasing since 1999 for Blacks and Hispanics, these groups still experience twice the mortality rate as non-Hispanic Whites, with Asian/Pacific Islanders remaining stable and American Indian and Alaskan Natives fluctuating over time (Conroy et al., 2014).

As discussed in the Background on Diabetes Mellitus and Insulin for Glycemic Control section, total direct medical expenses in California were estimated to be \$27.6 billion. An additional \$9.5 billion was spent on indirect costs due to lost productivity. Indirect costs have also been reported as high as \$32.6 billion when including morbidity and premature mortality costs (Shrestha et al., 2018). For non-Medicare or Medicaid payers (private insurance, other payers, and out-of-pocket from patients), medical costs related to diabetes are \$11.7 billion in California (Shrestha et al., 2018).

In the long term, the impact of AB 97 on premature mortality is unknown due to the lack of evidence that reduced cost sharing for insulin reduces mortality. However, well-controlled blood glucose results in fewer diabetes-related comorbidities (e.g. blindness, amputations, kidney disease, etc.). Therefore, for those patients who attain good glycemic control through increased adherence to insulin, these diabetes-related comorbidities that are known to lead to premature death could be prevented, delayed, or ameliorated.

The impact of AB 97 on economic loss is unknown due to the lack of literature on this topic. However, to the extent that better glycemic control is achieved, and comorbidities and lost productivity reduced, there is the potential for reduced economic loss.

⁵³ For more information about CHBRP's public health methodology, see http://chbrp.com/analysis_methodology/public_health_impact_analysis.php.

⁵⁴ The overall impact of premature death due to a particular disease can be measured in years of potential life lost prior to age 75 and summed for the population (generally referred to as "YPLL") (Gardner and Sanborn, 1990).

APPENDIX A TEXT OF BILL ANALYZED

On February 16, 2021, the California Assembly Committee on Health requested that CHBRP analyze AB 97.

ASSEMBLY BILL NO. 97

Introduced by Assembly Member Nazarian

December 08, 2020

An act-relating to insulin. to amend Section 1367.51 of the Health and Safety Code, and to amend Section 10176.61 of the Insurance Code, relating to health care coverage.

LEGISLATIVE COUNSEL'S DIGEST

AB 97, as amended, Nazarian. Insulin-Health care coverage: insulin affordability.

Existing law, the Knox-Keene Health Care Service Plan Act of 1975, provides for the licensure and regulation of health care service plans by the Department of Managed Health-Care. Care and makes a willful violation of the act's requirements a crime. Existing law provides for the regulation of health insurers by the Department of Insurance. Existing law requires a health care service plan contract or health disability insurance policy issued, amended, delivered, or renewed on or after January 1, 2000, that covers prescription benefits to include coverage for insulin, if it is determined to be medically necessary.

This bill would express the intent of the Legislature to enact legislation to make insulin more affordable for Californians.

This bill would prohibit a health care service plan contract or a health insurance policy issued, amended, delivered, or renewed on or after January 1, 2022, from imposing a deductible on an insulin prescription drug. Because a willful violation of these provisions by a health care service plan would be a crime, the bill would impose a state-mandated local program.

The California Constitution requires the state to reimburse local agencies and school districts for certain costs mandated by the state. Statutory provisions establish procedures for making that reimbursement.

This bill would provide that no reimbursement is required by this act for a specified reason.

Vote: majority Appropriation: no Fiscal Committee: noyes Local Program: noyes

THE PEOPLE OF THE STATE OF CALIFORNIA DO ENACT AS FOLLOWS:

SECTION 1. (a) The Legislature finds and declares all of the following:

- (1) Approximately 263,000 Californians are diagnosed with type 1 diabetes each year. Approximately 4,037,000 Californian adults have diabetes.
- (2) Every Californian with type 1 diabetes, and many with type 2 diabetes, rely on daily doses of insulin to survive.
- (3) Insulin prices have nearly tripled, creating financial hardships for people who rely on it to survive.
- (4) One in four people using insulin have reported insulin underuse due to the high cost of insulin.
- (5) Imposing a deductible on insulin, and requiring individuals to meet that deductible, creates a financial burden that presents a barrier to accessing insulin.
- (6) Diabetes is the seventh leading cause of death, and it is a leading cause of disabling and lifethreatening complications, including heart disease, stroke, kidney failure, amputation of the lower extremities, and new cases of blindness among adults.
- (7) Studies have shown that managing diabetes can prevent complications and medical emergencies associated with diabetes that result in emergency room visits, hospitalizations, and costly treatments.
- (b) Therefore, it is the intent of the Legislature to enact legislation on important policies to reduce the costs for Californians with diabetes to obtain life-saving and life-sustaining insulin.
- SEC. 2. Section 1367.51 of the Health and Safety Code is amended to read:
- **1367.51.** (a) Every A health care service plan contract, except a specialized health care service plan contract, that is issued, amended, delivered, or renewed on or after January 1, 2000, and that covers hospital, medical, or surgical expenses shall include coverage for the following equipment and supplies for the management and treatment of insulin-using diabetes, non-insulin-using diabetes, and gestational diabetes as medically necessary, even if the items are available without a prescription:
- (1) Blood glucose monitors and blood glucose testing strips.
- (2) Blood glucose monitors designed to assist the visually impaired.
- (3) Insulin pumps and all related necessary supplies.
- (4) Ketone urine testing strips.

- (5) Lancets and lancet puncture devices.
- (6) Pen delivery systems for the administration of insulin.
- (7) Podiatric devices to prevent or treat diabetes-related complications.
- (8) Insulin syringes.
- (9) Visual aids, excluding eyewear, to assist the visually impaired with proper dosing of insulin.
- (b) Every A health care service plan contract, except a specialized health care service plan contract, that is issued, amended, delivered, or renewed on or after January 1, 2000, that covers prescription benefits shall include coverage for the following prescription items if the items are determined to be medically necessary:
- (1) Insulin.
- (2) Prescriptive medications for the treatment of diabetes.
- (3) Glucagon.
- (c) The copayments and deductibles for the benefits specified in subdivisions (a) and (b) shall not exceed those established for similar benefits within the given plan.
- (d) (1) Notwithstanding subdivision (c), a health care service plan contract that is issued, amended, or renewed on or after January 1, 2022, shall not impose a deductible on an insulin prescription drug.
- (2) For purposes of this subdivision, "insulin prescription drug" means a prescription drug that contains insulin and is used to control blood glucose levels to treat diabetes.

(d) Every plan

(e) A health care service plan shall provide coverage for diabetes outpatient self-management training, education, and medical nutrition therapy necessary to enable an enrollee to properly use the equipment, supplies, and medications set forth in subdivisions (a) and (b), and additional diabetes outpatient self-management training, education, and medical nutrition therapy upon the direction or prescription of those services by the enrollee's participating physician. If a plan delegates outpatient self-management training to contracting providers, the plan shall require contracting providers to ensure that diabetes outpatient self-management training, education, and medical nutrition therapy are provided by appropriately licensed or registered health care professionals.

(e)

(f) The diabetes outpatient self-management training, education, and medical nutrition therapy services identified in subdivision-(d) (e) shall be provided by appropriately licensed or registered health care professionals as prescribed by a participating health care professional legally authorized to prescribe the service. These benefits shall include, but not be limited to, instruction that will enable diabetic patients and their families to gain an understanding of the diabetic disease process, and the daily management of diabetic therapy, in order to thereby avoid frequent hospitalizations and complications.

(f)

(g) The copayments for the benefits specified in subdivision-(d) (e) shall not exceed those established for physician office visits by the plan.

(g)Every

(h) A health care service plan governed by this section shall disclose the benefits covered pursuant to this section in the plan's evidence of coverage and disclosure forms.

(h)

- (i) A health care service plan-may shall not reduce or eliminate coverage as a result of the requirements of this section.
- (i) Nothing in this section shall be construed to
- (j) This section does not deny or restrict in any way the department's authority to ensure plan compliance with this chapter—when if a plan provides coverage for prescription drugs.
- SEC. 3. Section 10176.61 of the Insurance Code is amended to read:
- **10176.61.** (a) Every An insurer issuing, amending, delivering, or renewing a disability insurance policy on or after January 1, 2000, that covers hospital, medical, or surgical expenses shall include coverage for the following equipment and supplies for the management and treatment of insulinusing diabetes, non-insulin-using diabetes, and gestational diabetes as medically necessary, even if the items are available without a prescription:
- (1) Blood glucose monitors and blood glucose testing strips.
- (2) Blood glucose monitors designed to assist the visually impaired.
- (3) Insulin pumps and all related necessary supplies.
- (4) Ketone urine testing strips.
- (5) Lancets and lancet puncture devices.

- (6) Pen delivery systems for the administration of insulin.
- (7) Podiatric devices to prevent or treat diabetes-related complications.
- (8) Insulin syringes.
- (9) Visual aids, excluding eyewear, to assist the visually impaired with proper dosing of insulin.
- (b) Every An insurer issuing, amending, delivering, or renewing a disability insurance policy on or after January 1, 2000, that covers prescription benefits shall include coverage for the following prescription items if the items are determined to be medically necessary:
- (1) Insulin.
- (2) Prescriptive medications for the treatment of diabetes.
- (3) Glucagon.
- (c) The coinsurances and deductibles for the benefits specified in subdivisions (a) and (b) shall not exceed those established for similar benefits within the given policy.
- (d) (1) Notwithstanding subdivision (c), a health insurance policy that is issued, amended, or renewed on or after January 1, 2022, shall not impose a deductible on an insulin prescription drug.
- (2) For purposes of this subdivision, "insulin prescription drug" means a prescription drug that contains insulin and is used to control blood glucose levels to treat diabetes.

(d)Every insurer

(e) An insurer shall provide coverage for diabetes outpatient self-management training, education, and medical nutrition therapy necessary to enable an insured to properly use the equipment, supplies, and medications set forth in subdivisions (a) and (b) and additional diabetes outpatient self-management training, education, and medical nutrition therapy upon the direction or prescription of those services by the insured's participating physician. If an insurer delegates outpatient self-management training to contracting providers, the insurer shall require contracting providers to ensure that diabetes outpatient self-management training, education, and medical nutrition therapy are provided by appropriately licensed or registered health care professionals.

(e)

(f) The diabetes outpatient self-management training, education, and medical nutrition therapy services identified in subdivision-(d) (e) shall be provided by appropriately licensed or registered health care professionals as prescribed by a health care professional legally authorized to prescribe the services.

(f)

(g) The coinsurances and deductibles for the benefits specified in subdivision (d) (e) shall not exceed those established for physician office visits by the insurer.

(g)

(h) Every disability insurer governed by this section shall disclose the benefits covered pursuant to this section in the insurer's evidence of coverage and disclosure forms.

(h)An insurer may

(i) An insurer shall not reduce or eliminate coverage as a result of the requirements of this section.

(i)

- (j) This section does not apply to vision-only, dental-only, accident-only, specified disease, hospital indemnity, Medicare supplement, long-term care, or disability income insurance, except that for accident-only, specified disease, and hospital indemnity insurance coverage, benefits under this section only apply to the extent that the benefits are covered under the general terms and conditions that apply to all other benefits under the policy. Nothing in this section may be construed as imposing This section does not impose a new benefit mandate on accident-only, specified disease, or hospital indemnity insurance.
- **SEC. 4.** No reimbursement is required by this act pursuant to Section 6 of Article XIII B of the California Constitution because the only costs that may be incurred by a local agency or school district will be incurred because this act creates a new crime or infraction, eliminates a crime or infraction, or changes the penalty for a crime or infraction, within the meaning of Section 17556 of the Government Code, or changes the definition of a crime within the meaning of Section 6 of Article XIII B of the California Constitution.

SECTION 1. It is the intent of the Legislature to enact legislation to make insulin more affordable for Californians.

APPENDIX B LITERATURE REVIEW METHODS

This appendix describes methods used in the literature review conducted for this report. A discussion of CHBRP's system for medical effectiveness grading evidence, as well as lists of MeSH Terms, publication types, and keywords, follows.

Studies of cost sharing related to insulin use and adherence for diabetes were identified through searches of PubMed, the Cochrane Library, Web of Science, and the Cumulative Index of Nursing and Allied Health Literature. Websites maintained by the following organizations that produce and/or index meta-analyses and systematic reviews were also searched: the Agency for Healthcare Research and Quality (AHRQ), the International Network of Agencies for Health Technology Assessment (INAHTA), the National Health Service (NHS) Centre for Reviews and Dissemination, the National Institute for Health and Clinical Excellence (NICE), and the Scottish Intercollegiate Guideline Network. The American Diabetic Association (ADA) website and available resources were also searched as pertinent to this bill and reviewed.

The search was limited to studies published from 2020 to present, because CHBRP had previously reviewed this literature using the same search terms in 2020 for the AB 2203 analysis. Articles were eliminated if they did not focus on a specific treatment, were from outside the United States, were of poor quality, or did not report findings from clinical research studies. The American Diabetic Association (ADA) website and available resources were also searched as pertinent to this bill and reviewed.

Reviewers screened the title and abstract of each citation retrieved by the literature search to determine eligibility for inclusion. The reviewers acquired the full text of articles that were deemed eligible for inclusion in the review and reapplied the initial eligibility criteria.

Medical Effectiveness Review

The medical effectiveness literature review returned abstracts for 122 articles, of which 19 were reviewed for inclusion in this report. A total of two new studies since 2020 were included in the medical effectiveness review for AB 97, as well as five studies that were included in the previous review for AB 2203.

Medical Effectiveness Evidence Grading System

In making a "call" for each outcome measure, the medical effectiveness lead and the content expert consider the number of studies as well the strength of the evidence. Further information about the criteria CHBRP uses to evaluate evidence of medical effectiveness can be found in CHBRP's *Medical Effectiveness Analysis Research Approach*. To grade the evidence for each outcome measured, the team uses a grading system that has the following categories:

- Research design;
- Statistical significance;
- Direction of effect;
- Size of effect; and
- Generalizability of findings.

The grading system also contains an overall conclusion that encompasses findings in these five domains. The conclusion is a statement that captures the strength and consistency of the evidence of an intervention's effect on an outcome. The following terms are used to characterize the body of evidence regarding an outcome:

Clear and convincing evidence;

⁵⁵ Available at: http://chbrp.com/analysis_methodology/medical_effectiveness_analysis.php.

- Preponderance of evidence;
- Limited evidence;
- Inconclusive evidence; and
- Insufficient evidence.

A grade of *clear and convincing evidence* indicates that there are multiple studies of a treatment and that the <u>large majority</u> of studies are of high quality and consistently find that the treatment is either effective or not effective.

A grade of *preponderance of evidence* indicates that the <u>majority</u> of the studies reviewed are consistent in their findings that treatment is either effective or not effective.

A grade of *limited evidence* indicates that the studies had limited generalizability to the population of interest and/or the studies had a fatal flaw in research design or implementation.

A grade of *inconclusive evidence* indicates that although some studies included in the medical effectiveness review find that a treatment is effective, a similar number of studies of equal quality suggest the treatment is not effective.

A grade of *insufficient evidence* indicates that there is not enough evidence available to know whether or not a treatment is effective, either because there are too few studies of the treatment or because the available studies are not of high quality. It does not indicate that a treatment is not effective.

Search Terms (* indicates truncation of word stem)

- A1c
- Adherence
- Age
- Age Factors
- American
- Amputation
- Birth Injuries
- Birth Weight
- Birth/Newborn/Infant Hypoglycemia
- Birth/Newborn/Infant Size
- Bisexual
- Blindness
- Cardiovascular Diseases
- Cardiovascular Disorders
- Coinsurance
- Co-Insurance
- Coma
- Comorbidity
- Compliance
- Complications
- Copay
- Co-Pay
- Copayments (Insurance)
- Cost
- Cost Control
- Cost Effectiveness
- Cost Sharing
- Cost Shifting

- Cost Shifting in Employer-Based Health Insurance
- · Costs and Cost Analysis
- Deductible
- Deductibles (Insurance)
- Diabetes
- Diabetes Complications
- Diabetes Management
- Diabetes Mellitus
- Diabetes Mellitus Complications
- Diabetic Angiopathies
- Diabetic Cardiomyopathies
- Diabetic Coma
- Diabetic Ketoacidosis
- Diabetic Neuropathies
- Diabetic Patients
- Disease Management
- Disparities
- Disparity
- Donohue Syndrome
- Drug Utilization
- Drug Utilization Review
- Dystocia
- Economics
- Emergency Service, Hospital
- Emergency Service, Hospital
- Emergency Services
- Emergency Visits/Admissions
- ESRD

- Ethnic
- Ethnic Groups
- Facilities and Services Utilization
- Fetal Outcome
- Gender
- Gender Equality
- Gender Equity
- Gender Gap
- Gender Specific Care
- Gestational diabetes
- Glycated Hemoglobin A
- Glycated Hemoglobin levels
- Glycemic Control
- Glycohemoglobin A levels
- HbA1c
- Health Insurance
- Health Status Disparities
- Healthcare Disparities
- Heart Attack
- Homosexual
- Hospital Admission
- Hospital Discharge
- Hospitalization
- Hospitalization
- Hypoglycemia
- Incidence
- Income
- Income (Economic)
- Income Level
- Infant, Low Birth Weight
- Infant, Premature
- Infant, Premature, Diseases
- Insulin
- Insurance Costs
- Ketoacidosis
- Kidney Diseases
- Kidney Failure
- Kidney Failure, Chronic
- Lesbian
- LGBTQ
- Medication Adherence
- Minorities
- Minority
- Minority Groups
- Minority Health
- · Mortality, Premature
- Myocardial Infarction
- Newborn Respiratory Distress Syndrome
- Nonadherence
- Non-Compliance
- Office Visits
- Office Visits

- Outcome
- Outcome Assessment
- Out-Of-Pocket
- Patient Compliance
- Patient-Reported Outcomes
- Peripheral Neuropathy
- Pregnancy Complications
- Pregnancy in Diabetes
- Pregnancy Outcome
- Premature Death/Mortality
- Prevalence
- Race and Ethnic Discrimination
- Race Factors
- Racial
- Racial and Ethnic Attitudes
- Racial and Ethnic Differences
- Racial and Ethnic Groups
- Racial Disparities
- Renal Insufficiency, Chronic
- SDOH
- Sex Factors
- Sexual Behavior
- Sexual Orientation
- Sexuality
- Social Determinants
- Social Determinants of Health
- Stroke
- Transgender
- Transsexual
- Treatment Outcomes
- Treatment Refusal
- Type 2 Diabetes
- U.S.
- United States
- US
- Usage
- Utilization
- Utilization Review

APPENDIX C COST IMPACT ANALYSIS: DATA SOURCES, CAVEATS, AND ASSUMPTIONS

The cost analysis in this report was prepared by the members of the cost team, which consists of CHBRP task force members and contributors from the University of California, Los Angeles, and the University of California, Davis, as well as the contracted actuarial firm, Milliman, Inc.⁵⁶

Information on the generally used data sources and estimation methods, as well as caveats and assumptions generally applicable to CHBRP's cost impacts analyses are available at CHBRP's website.⁵⁷

This appendix describes analysis-specific data sources, estimation methods, caveats, and assumptions used in preparing this cost impact analysis.

Analysis-Specific Caveats and Assumptions

This subsection discusses the caveats and assumptions relevant specifically to an analysis of AB 97.

Assumptions for Baseline Benefit Coverage

- The population subject to the mandated offering includes enrollees in DMHC-regulated commercial insurance plans, CDI-regulated policies, and CalPERS plans subject to the requirements of the Knox-Keene Health Care Service Plan Act.
- CHBRP conducted a carrier survey to determine the percentage of enrollees that are enrolled in HSA qualified plans by regulator, line of business, and deductible or metal tier.
- CHBRP assumed all non-HSA plans have separate medical and pharmacy deductibles. Non-HSA
 plans with combined deductibles exist but they are not common.

Assumptions for Baseline Utilization and Cost

- National Drug Codes (NDCs) for insulin were identified using the MediSpan[®] Master Drug Data Base v2.5.
- Once identified, these NDCs for insulin were used to extract data from Milliman's 2019
 Consolidated Health Cost Guidelines Sources Database (CHSD). CHBRP limited its data pull to
 California only. These data were used to develop prevalence, utilization, baseline allowed cost,
 and enrollee cost-sharing information by commercial market segment for insulin users. In
 addition, CHBRP developed this information separately for two distinct groups of insulin users:
 - Enrollees who did not have any claims that exceeded the mandated cost-sharing cap;
 and
 - Enrollees who had at least one claim that exceeded the mandated cost-sharing cap.
- 2019 allowed cost for insulin was trended 1.5% per year from 2019 to 2022 based on recent and projected annual increases in net insulin prices.

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⁵⁶ CHBRP's authorizing statute, available at http://chbrp.com/CHBRP authorizing statute 2018 FINAL.pdf, requires that CHBRP use a certified actuary or "other person with relevant knowledge and expertise" to determine financial impact.

⁵⁷ See method documents posted at http://chbrp.com/analysis_methodology/cost_impact_analysis.php; in particular, see 2019 Cost Analyses: Data Sources, Caveats, and Assumptions.

Assumptions for Baseline Cost Sharing

- CHBRP summarized non-HSA pharmacy deductible information by regulator, line of business, and deductible or metal tier levels.
 - CHBRP assumed 14% of large group plans have a pharmacy deductible based on the large group percentage of workers with a separate pharmacy deductible from Kaiser Family Foundation's 2019 Employer Health Benefit Survey. CHBRP assumed large group plans with a pharmacy deductible have a pharmacy deductible of \$190 based on the average large group pharmacy deductible from Kaiser Family Foundation's 2019 Employer Health Benefit Survey.
 - CHBRP assumed 10% of the small group and individual grandfathered plans have a pharmacy deductible based on the small group percentage of workers with a separate pharmacy deductible from Kaiser Family Foundation's 2015 Employer Health Benefit Survey. CHBRP assumed the small group and individual grandfathered plans with a pharmacy deductible have a pharmacy deductible of \$160 based on the average small group pharmacy deductible from Kaiser Family Foundation's 2015 Employer Health Benefit Survey.
 - For all non-grandfathered small group plans, CHBRP reviewed the 2021 Covered California plan offerings. CHBRP assumed the platinum and gold plans have no pharmacy deductible. CHBRP assumed the silver and bronze plans have \$300 and \$500, respectively.
 - o For all non-grandfathered individual plans, CHBRP reviewed the 2022 Covered California plan offerings and assumed the non-HSA plan pharmacy deductible at each tier.
- CHBRP summarized HSA pharmacy deductibles using the 2021 individual and 2020 small group Covered California plans for individual and small group non-grandfathered plans. Large group and grandfathered plans are assumed to have a \$2,500 deductible from Kaiser Family Foundation's 2019 Employer Health Benefit Survey.
- CHBRP summarized the cost and utilization of insulin, all other drugs, and all other medical services by user using claims data from CHSD. CHBRP then used this distribution of costs by member to evaluate the value of each deductible. CHBRP was able to determine how many members would satisfy the deductible using other drugs or medical services, and would therefore be unaffected by this change.

Assumptions for Postmandate Utilization

• CHBRP assumes reduced cost sharing for insulin increases the utilization of outpatient prescription insulin based on literature that establishes evidence of price elasticity of demand for prescription drugs (Goldman et al., 2004). The authors found use of insulin decreased by 8% when co-payments doubled. Applying this to CHBRP's estimated reduction in cost sharing for those enrollees impacted by AB 97, CHBRP estimates about a 0.3% - 1.7% increase in utilization of insulin for enrollees whose cost sharing decreases as a result of the removal of a deductible depending on the plan design.

Assumptions for Postmandate Cost

- CHBRP did not assume service costs would increase as a result of AB 97.
- CHBRP did not assume any offsets associated with removing deductibles from the insulin benefit.

Assumptions for Postmandate Cost Sharing

- For all services, the postmandate cost-share for enrollees without deductibles did not change.
- CHBRP summarized the cost and utilization of insulin, all other drugs, and all other medical
 services by user. CHBRP then used this distribution of costs by member to evaluate the value of
 each deductible. CHBRP calculated the cost sharing for each member using the copay or
 coinsurance for the corresponding drug tier. For some enrollees who would satisfy the deductible
 using other drugs or medical services, their postmandate cost sharing was equal to their
 premandate cost sharing.

Determining Public Demand for the Proposed Mandate

This subsection discusses public demand for the benefits AB 97 would mandate. Considering the criteria specified by CHBRP's authorizing statute, CHBRP reviews public demand for benefits relevant to a proposed mandate in two ways. CHBRP:

- Considers the bargaining history of organized labor; and
- Compares the benefits provided by self-insured health plans or policies (which are not regulated by the DMHC or CDI and therefore not subject to state-level mandates) with the benefits that are provided by plans or policies that would be subject to the mandate.

Among publicly funded self-insured health insurance policies, the preferred provider organization (PPO) plans offered by CalPERS currently have the largest number of enrollees. The CalPERS PPOs currently provide benefit coverage different to what is available through group health insurance plans and policies that would be subject to the mandate, by specifying that cost sharing for insulin for CalPERS enrollees is below the proposed threshold.

To further investigate public demand, CHBRP used the bill-specific coverage survey to ask carriers who act as third-party administrators for (non-CalPERS) self-insured group health insurance programs whether the relevant benefit coverage differed from what is offered in group market plans or policies that would be subject to the mandate. The responses indicated that there were no substantive differences.

In 2020, CVS Health, a pharmacy benefit manager, began offering a plan design with zero out-of-pocket costs for diabetes medications and supplies.⁵⁸ This indicates there is a demand from employers and other purchasers of pharmacy benefits for plan designs with no or low cost sharing for insulin prescriptions.

Second Year Impacts on Benefit Coverage, Utilization, and Cost

CHBRP has considered whether continued implementation during the second year of the benefit coverage requirements of AB 97 would have a substantially different impact on utilization of either the tests, treatments, or services for which coverage was directly addressed, the utilization of any indirectly affected utilization, or both. CHBRP reviewed the literature and consulted content experts about the possibility of varied second year impacts and determined the second year's impacts of AB 97 would be substantially the same as the impacts in the first year (see Table 1). Minor changes to utilization and expenditures are due to population changes between the first year postmandate and the second year postmandate.

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⁵⁸ CVS Health, A prescription for better diabetes management: RxZERO plan design eliminates member out-of-pocket costs. January 2020.

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CALIFORNIA HEALTH BENEFITS REVIEW PROGRAM COMMITTEES AND STAFF

A group of faculty, researchers, and staff complete the analysis that informs California Health Benefits Review Program (CHBRP) reports. The CHBRP **Faculty Task Force** comprises rotating senior faculty from University of California (UC) campuses. In addition to these representatives, there are other ongoing researchers and analysts who are **Task Force Contributors** to CHBRP from UC that conduct much of the analysis. The **CHBRP staff** coordinates the efforts of the Faculty Task Force, works with Task Force members in preparing parts of the analysis, and manages all external communications, including those with the California Legislature. As required by CHBRP's authorizing legislation, UC contracts with a certified actuary, **Milliman**, to assist in assessing the financial impact of each legislative proposal mandating or repealing a health insurance benefit.

The **National Advisory Council** provides expert reviews of draft analyses and offers general guidance on the program to CHBRP staff and the Faculty Task Force. CHBRP is grateful for the valuable assistance of its National Advisory Council. CHBRP assumes full responsibility for the report and the accuracy of its contents.

Faculty Task Force

Timothy T. Brown, PhD, University of California, Berkeley

Janet Coffman, MA, MPP, PhD, Vice Chair for Medical Effectiveness, University of California, San Francisco

Sylvia Guendelman, PhD, LCSW, University of California, Berkeley

Gerald Kominski, PhD, University of California, Los Angeles

Sara McMenamin, PhD, Vice Chair for Medical Effectiveness and Public Health, University of California, San Diego

Joy Melnikow, MD, MPH, Vice Chair for Public Health, University of California, Davis

Jack Needleman, PhD, University of California, Los Angeles

Nadereh Pourat, PhD, Vice Chair for Cost, University of California, Los Angeles

Marilyn Stebbins, PharmD, University of California, San Francisco

Task Force Contributors

Danielle Casteel, MA, University of California, San Diego

Shana Charles, PhD, MPP, University of California, Los Angeles, and California State University, Fullerton

Margaret Fix, MPH, University of California, San Francisco

Naomi Hillery, MPH, University of California, San Diego

Jeffrey Hoch, PhD. University of California, Davis

Julia Huerta, MPH, University of California, Davis

Michelle Keller, PhD, MPH, University of California, Los Angeles

Connie Kwong, University of California, San Francisco

Elizabeth Magnan, MD, PhD, University of California, Davis

Jacqueline Miller, University of California, San Francisco

Marykate Miller, MS, University of California, Davis

Dominique Ritley, MPH, University of California, Davis

Dylan Roby, PhD, University of California, Los Angeles, and University of Maryland, College Park

Emily Shen, University of California, San Francisco

Riti Shimkhada, PhD, University of California, Los Angeles

Meghan Soulsby Weyrich, MPH, University of California, Davis

Steven Tally, PhD. University of California, San Diego

Sara Yoeun, MPH, University of California, San Diego

National Advisory Council

Lauren LeRoy, PhD, Strategic Advisor, L. LeRoy Strategies, Chair

Stuart H. Altman, PhD, Professor of National Health Policy, Brandeis University, Waltham, MA

Deborah Chollet, PhD, Senior Fellow, Mathematica Policy Research, Washington, DC

Allen D. Feezor, Former Deputy Secretary for Health Services, North Carolina Department of Health and Human Services, Raleigh, NC

Charles "Chip" Kahn, MPH, President and CEO, Federation of American Hospitals, Washington, DC Jeffrey Lerner, PhD, President Emeritus, ECRI Institute Headquarters, Plymouth Meeting, PA; Adjunct Senior Fellow, Leonard Davis Institute of Health Economics, University of Pennsylvania

Donald E. Metz, Executive Editor, Health Affairs, Bethesda, MD

Dolores Mitchell, (Retired) Executive Director, Group Insurance Commission, Boston, MA
 Marilyn Moon, PhD, Senior Fellow, Retired, American Institutes for Research, Washington, DC
 Carolyn Pare, (Retired) President and CEO, Minnesota Health Action Group, Bloomington, MN
 Richard Roberts, MD, JD, Professor Emeritus of Family Medicine, University of Wisconsin-Madison, Madison, WI

Alan Weil, JD, MPP, Editor-in-Chief, Health Affairs, Bethesda, MD

CHBRP Staff

Garen Corbett, MS, Director John Lewis, MPA, Associate Director Adara Citron, MPH, Principal Policy Analyst Karen Shore, PhD, Contractor* An-Chi Tsou, PhD, Contractor* California Health Benefits Review Program MC 3116
Berkeley, CA 94720-3116
info@chbrp.org

info@chbrp.org (510) 664-5306

*Karen Shore, PhD, and An-Chi Tsou, PhD, are Independent Contractors who work with CHBRP to support legislative analyses and other special projects on a contractual basis.

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Danielle Casteel, MA, of the University of California, San Diego, prepared the medical effectiveness analysis. Stephen L. Clancy, MLS, AHIP, of the University of California, Irvine, conducted the literature search. Adara Citron, MPH, of CHBRP staff, prepared the prepared the policy Context, public health impact analysis and the cost impact analysis, and synthesized the individual sections into a single report. Barbara Dewey, FSA, MAAA, of Milliman, provided actuarial analysis. A member of the CHBRP Faculty Task Force, Sara McMenamin, PhD, of the University of California, San Diego, reviewed the analysis for its accuracy, completeness, clarity, and responsiveness to the Legislature's request.

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CHBRP assumes full responsibility for the report and the accuracy of its contents. All CHBRP bill analyses and other publications are available at www.chbrp.org.

Garen Corbett, MS Director

Please direct any questions concerning this document to: California Health Benefits Review Program; MC 3116; Berkeley, CA 94720-3116, info@chbrp.org, or www.chbrp.org