

# **Supplemental Material**

**Table S1. Modeling Inputs – DELIVER Population Only.**

	<b>Value</b>	<b>Range</b>	<b>Source</b>
<b><u>Transition Probabilities*</u></b>			
Worsening HF events	0.022	0.016-0.032	Trial
Cardiovascular Mortality	0.004	0.003-0.006	Trial
<b><u>Effectiveness of Dapagliflozin vs. Placebo</u></b>			
Worsening HF events	0.73	0.62-0.87	Trial
Cardiovascular Mortality	0.88	0.74-1.05	Trial
<b><u>Proportions of Events (US Population)</u></b>			
Proportion of Worsening HF Events attributable to HF Hospitalization	0.92	0.87-0.95	Trial
Proportion of All-Cause Mortality attributable to CV Mortality	0.47	0.36-0.57	Trial
<b><u>Utilities</u></b>			
Dapagliflozin	0.825	0.821-0.829	Isaza et. al. & prior modeling <sup>6,29,30</sup> ; Trial
Placebo	0.811	0.806-0.815	Isaza et. al. & prior modeling <sup>6,29,30</sup> ; Trial

\*Transition probabilities derived from placebo event rates among US participants in DELIVER. Worsening HF events include hospitalization for HF and urgent HF visits.

“Trial” refers to participant-level data from the DELIVER trial. All other modeling parameters are consistent with those reported in Table 1.

**Table S2. Cost Effectiveness at Various Monthly Costs of Dapagliflozin using Pooled Data from DAPA-HF and DELIVER.**

	Mean # Worsening HF Events	Life- years	Costs (\$)	Effectiveness (QALYs)	Incremental Cost (\$)	Incremental Effectiveness (QALYs)	ICER (\$/QALY)
<b>Medicare Part-D (\$514.95/mo)<sup>23</sup></b>							
Standard of Care	2.38	7.47	109,003	6.04	45,509	0.53	85,554
Dapagliflozin	1.87	7.98	154,512	6.57			
<b>Medicare Part-D w/ 49% Rebate (\$262.62/mo)<sup>^8</sup></b>							
Standard of Care	2.38	7.47	111,561	6.04	21,321	0.53	40,081
Dapagliflozin	1.87	7.98	130,324	6.57			
<b>Wholesale Acquisition Cost (\$548.83/mo)<sup>24</sup></b>							
	2.38	7.47	109,003	6.04	48,765	0.53	91,675
	1.87	7.98	157,768	6.57			
<b>Federal Supply Schedule Big Four (\$396.14/mo)<sup>25</sup></b>							
	2.38	7.47	109,003	6.04	34,120	0.53	64,143
	1.87	7.98	151,913	6.57			
<b>Medicare Part-D w/ Reduced Rebate (\$314.08/mo)<sup>*8,21</sup></b>							
	2.38	7.47	109,003	6.04	26,257	0.53	49,362
	1.87	7.98	135,260	6.57			
<b>Canadian Estimate (\$68.25/mo)<sup>27</sup></b>							
	2.38	7.47	109,003	6.04	2,688	0.53	5,053
	1.87	7.98	111,691	6.57			

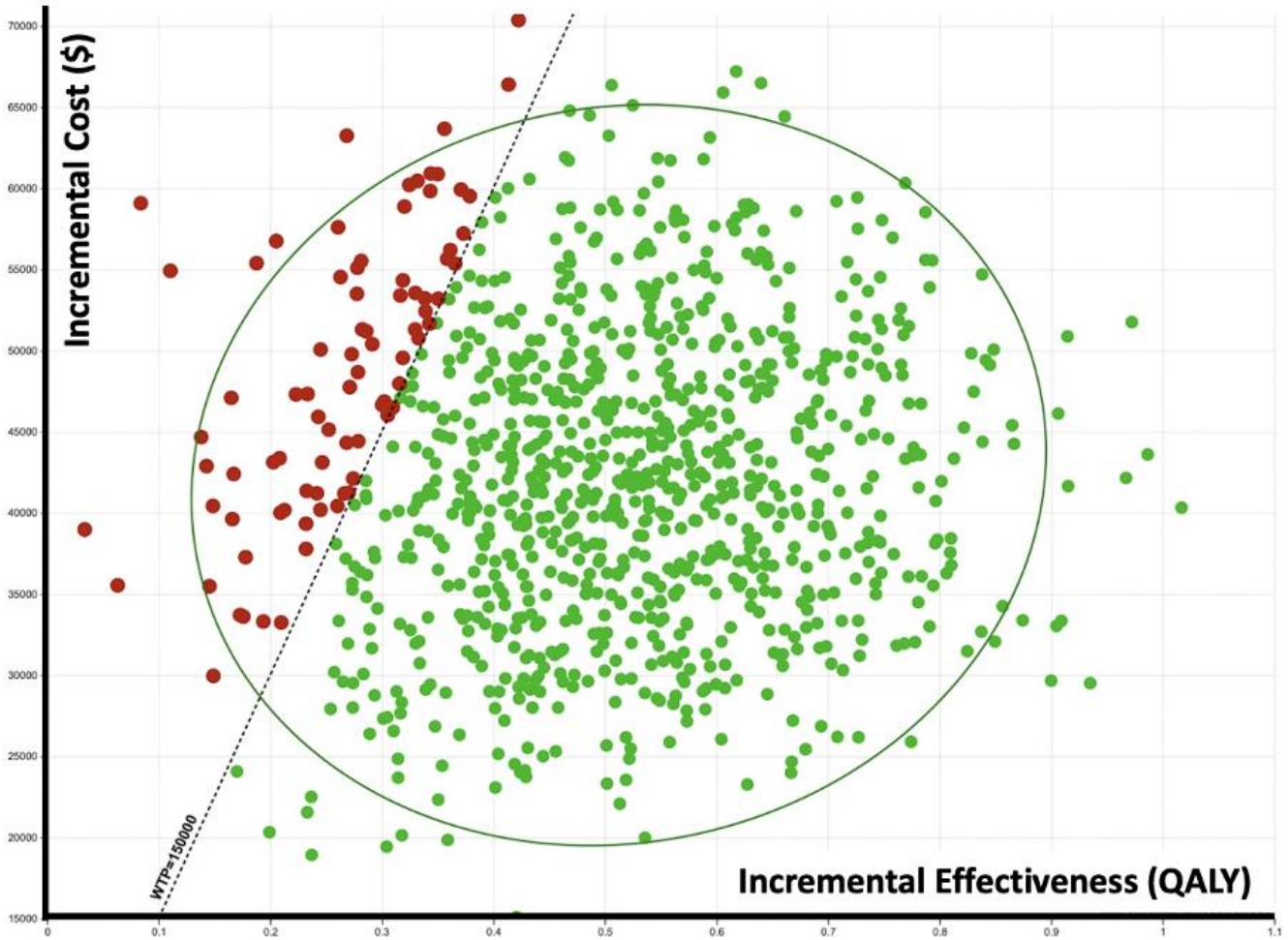
Worsening HF events = hospitalization for heart failure or urgent HF visit; QALYs = quality-adjusted life years; mo = month

<sup>^</sup> Based on published estimate of 49% rebates.

<sup>\*</sup>Based on published estimates of a reduced rebate in which 20% of the rebate is retained by entities.

Note: ICERs represent ratio of incremental costs and incremental QALY without rounding.

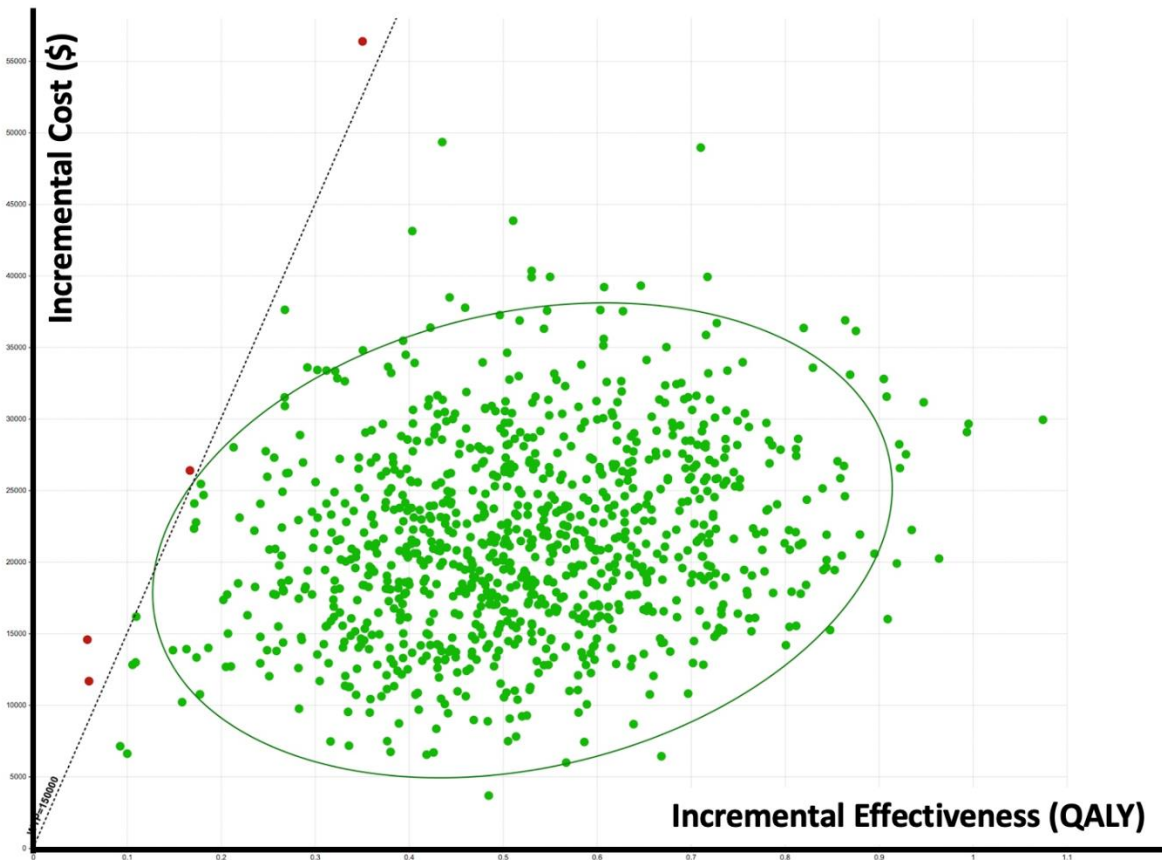
**Figure S1. Probabilistic Sensitivity Analysis of the Addition of Dapagliflozin to Standard of Care in Chronic HF.**



Model parameters were independently varied across their distributions in a probabilistic sensitivity analysis for 100,000 iterations using the full (undiscounted) Medicare cost, with each iteration displayed as a dot in this scatter plot. The dashed black line represents a willingness to pay threshold of \$150,000 per QALY gained. The green oval represents points falling in the 95% credible interval. Green dots represent iterations at an ICER <\$150,000 per QALY gained; red dots represent iterations at an ICER  $\geq$ \$150,000 per QALY gained.

WTP = willingness-to-pay

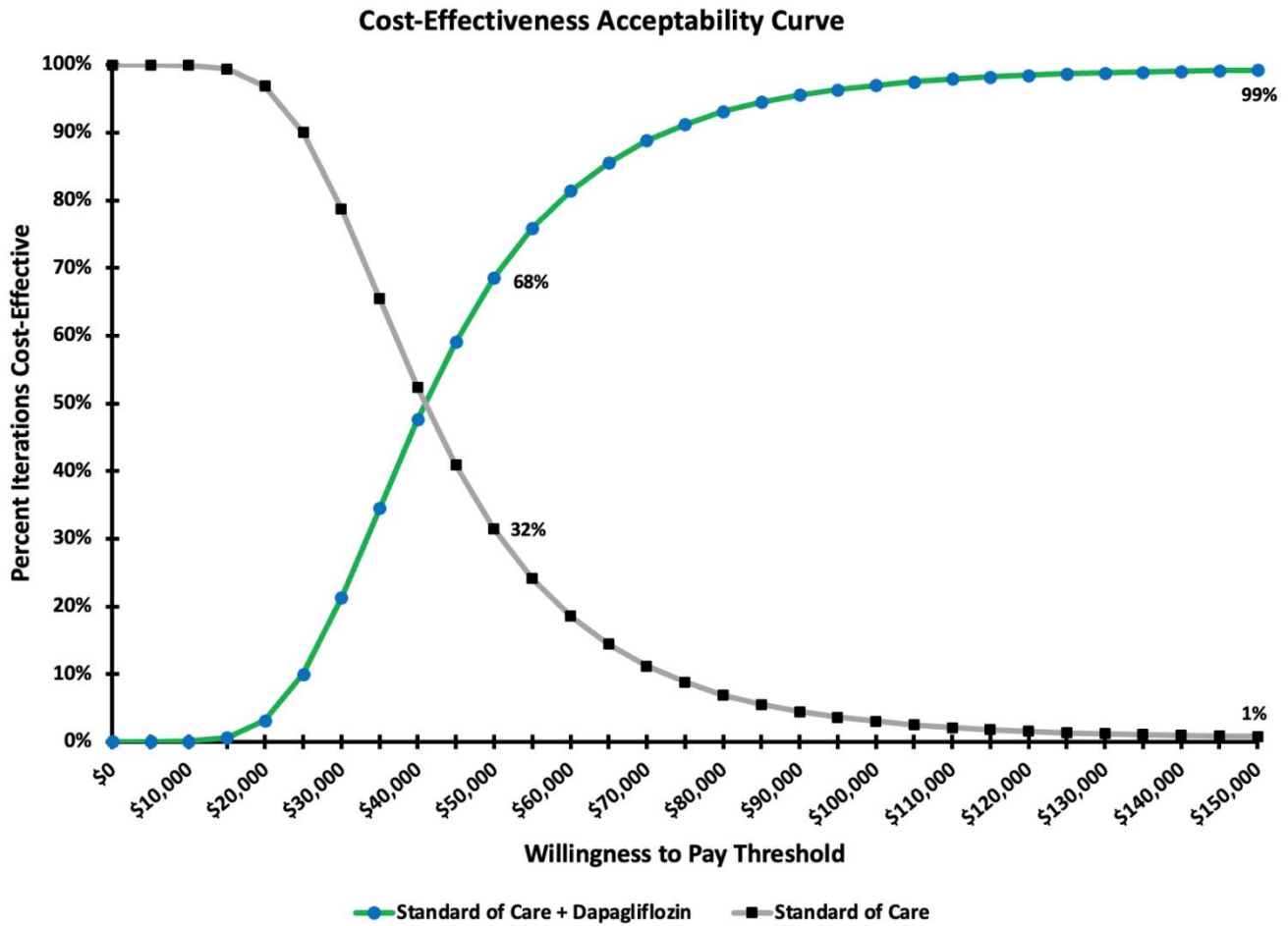
**Figure S2. Probabilistic Sensitivity Analysis of the Addition of Dapagliflozin to Standard of Care in Chronic HF.**



Model parameters were independently varied across their distributions in a probabilistic sensitivity analysis for 100,000 iterations using the discounted Medicare cost (\$262/mo), with each iteration displayed as a dot in this scatter plot. The dashed black line represents a willingness to pay threshold of \$150,000 per QALY gained. The green oval represents points falling in the 95% credible interval. Green dots represent iterations at an ICER <\$150,000 per QALY gained; red dots represent iterations at an ICER ≥\$150,000 per QALY gained.

WTP = willingness-to-pay

Figure S3. Cost-effectiveness Acceptability Curve based on Probabilistic Sensitivity Analysis.



All model parameters were independently varied across their distributions in a probabilistic sensitivity analysis for 100,000 iterations using the discounted Medicare cost (\$262/mo). The percentage of iterations that were cost-effective is plotted across various willingness-to-pay thresholds.

## CHEERS 2022 Checklist

Topic	No.	Item	Location where item is reported
<b>Title</b>			
	1	Identify the study as an economic evaluation and specify the interventions being compared.	Title, Page 1
<b>Abstract</b>			
	2	Provide a structured summary that highlights context, key methods, results, and alternative analyses.	Abstract, Page 5
<b>Introduction</b>			
<b>Background and objectives</b>	3	Give the context for the study, the study question, and its practical relevance for decision making in policy or practice.	Introduction, Paragraph 1-2
<b>Methods</b>			
<b>Health economic analysis plan</b>	4	Indicate whether a health economic analysis plan was developed and where available.	Methods, Paragraph 1
<b>Study population</b>	5	Describe characteristics of the study population (such as age range, demographics, socioeconomic, or clinical characteristics).	Methods, Paragraph 1
<b>Setting and location</b>	6	Provide relevant contextual information that may influence findings.	Methods, Paragraph 1
<b>Comparators</b>	7	Describe the interventions or strategies being compared and why chosen.	Methods, Paragraph 1
<b>Perspective</b>	8	State the perspective(s) adopted by the study and why chosen.	Methods, Paragraph 1
<b>Time horizon</b>	9	State the time horizon for the study and why appropriate.	Statistical Analysis, Paragraph 1
<b>Discount rate</b>	10	Report the discount rate(s) and reason chosen.	Statistical Analysis, Paragraph 1
<b>Selection of outcomes</b>	11	Describe what outcomes were used as the measure(s) of benefit(s) and harm(s).	Baseline assumptions and modeling inputs,

Topic	No.	Item	Location where item is reported
<b>Measurement of outcomes</b>	12	Describe how outcomes used to capture benefit(s) and harm(s) were measured.	Baseline assumptions and modeling inputs,
<b>Valuation of outcomes</b>	13	Describe the population and methods used to measure and value outcomes.	Baseline assumptions and modeling inputs,
<b>Measurement and valuation of resources and costs</b>	14	Describe how costs were valued.	Costs, Paragraph 1-2
<b>Currency, price date, and conversion</b>	15	Report the dates of the estimated resource quantities and unit costs, plus the currency and year of conversion.	Costs, Paragraph 3
<b>Rationale and description of model</b>	16	If modelling is used, describe in detail and why used. Report if the model is publicly available and where it can be accessed.	Statistical Analysis, Paragraph 1
<b>Analytics and assumptions</b>	17	Describe any methods for analysing or statistically transforming data, any extrapolation methods, and approaches for validating any model used.	Baseline assumptions and modeling inputs
<b>Characterising heterogeneity</b>	18	Describe any methods used for estimating how the results of the study vary for subgroups.	Statistical Analysis, Paragraph 2
<b>Characterising distributional effects</b>	19	Describe how impacts are distributed across different individuals or adjustments made to reflect priority populations.	Baseline assumptions and modeling inputs
<b>Characterising uncertainty</b>	20	Describe methods to characterise any sources of uncertainty in the analysis.	Not Statistical Analysis, Paragraph 2
<b>Approach to engagement with patients and others affected by the study</b>	21	Describe any approaches to engage patients or service recipients, the general public, communities, or stakeholders (such as clinicians or payers) in the design of the study.	N/A
<b>Results</b>			
<b>Study parameters</b>	22	Report all analytic inputs (such as values, ranges, references) including uncertainty or distributional assumptions.	Results, Paragraph 1



Topic	No.	Item	Location where item is reported
<b>Summary of main results</b>	23	Report the mean values for the main categories of costs and outcomes of interest and summarise them in the most appropriate overall measure.	Results, Paragraph 2
<b>Effect of uncertainty</b>	24	Describe how uncertainty about analytic judgments, inputs, or projections affect findings. Report the effect of choice of discount rate and time horizon, if applicable.	Paragraph 3
<b>Effect of engagement with patients and others affected by the study</b>	25	Report on any difference patient/service recipient, general public, community, or stakeholder involvement made to the approach or findings of the study	Not reported
<b>Discussion</b>			
<b>Study findings, limitations, generalisability, and current knowledge</b>	26	Report key findings, limitations, ethical or equity considerations not captured, and how these could affect patients, policy, or practice.	Discussion
<b>Other relevant information</b>			
<b>Source of funding</b>	27	Describe how the study was funded and any role of the funder in the identification, design, conduct, and reporting of the analysis	Disclosures, Page 3
<b>Conflicts of interest</b>	28	Report authors conflicts of interest according to journal or International Committee of Medical Journal Editors requirements.	Disclosures, Page 2-3

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