## **Legend:** New Text Removed Text Unchanged Text Moved Text Section

As a large global health company operating in a complex industry, we encounter a variety of risks and uncertainties, which could have a material adverse effect on our business, liquidity, results of operations, financial condition or the trading price of our securities. You should carefully consider each of the risks and uncertainties discussed below, together with other information contained in this Form 10-K, including MD & A. These risks and uncertainties are not the only ones we face. Additional risks and uncertainties not presently known to us or that we currently believe to be immaterial may also adversely affect us. The following risk factors have been organized by category for ease of use; however many of the risks may have impacts in more than one category. These categories, therefore, should be viewed as a starting point for understanding the significant risks facing us and not as a limitation on the potential impact of the matters discussed. Risk factors are not necessarily listed in order of importance. Strategic and Operational Risks Future performance of our business will depend on our ability to execute our strategic and operational initiatives effectively. The future performance of our business will depend in large part on our ability to effectively implement and execute our strategic and operational initiatives. Successfully executing on these initiatives depends on a number of factors, including our ability to: • differentiate our products, services and solutions from those of our competitors; • develop and bring to market new and innovative products, solutions or programs that focus on improving patient outcomes and experiences and assist in controlling costs or in response to government regulation; • develop and create data and analytic solutions to support and improve outcomes for our products, services and solutions, including creating and developing solutions and services through partnerships with other industry participants; • grow and support our product portfolio, expand our addressable markets and identify and introduce the proper mix, coordination or integration of products that will be accepted by the marketplace; • evaluate drugs for efficacy, value and price to assist clients in selecting a cost- effective formulary; • offer cost- effective home delivery pharmacy and specialty services; • access or continue accessing key drugs and successfully penetrate key treatment categories in our specialty pharmacy business; • attract and retain sufficient numbers of qualified employees, particularly in <mark>a <del>an increasingly</del> competitive job market; • attract, develop and maintain</mark> collaborative relationships with a sufficient number of qualified partners; • attract new and maintain existing customer and client relationships; • leverage purchase volume to deliver discounts to health benefit providers; • transition health care providers from volume- based fee- for- service arrangements to a value- based system; • improve medical cost competitiveness in our targeted markets; • manage our medical, pharmacy, administrative and other operating costs effectively; and • contract with health care providers, pharmacy providers and pharmaceutical manufacturers on market competitive terms. For our strategic initiatives to succeed, we must effectively collaborate across our operations, integrate our acquired businesses, actively work to ensure consistency throughout the organization and promote a global mindset along with a focus on individual customers and clients. If we fail to do so, our business may be unable to grow as planned, or the result of expansion may be unsatisfactory. We will be unable to rapidly respond to competitive, economic and regulatory changes if we do not make important strategic and operational decisions quickly, define our appetite for risk, implement new governance, managerial and organizational processes smoothly and communicate roles and responsibilities clearly. If these initiatives fail or are not executed effectively, our consolidated financial position and results of operations could be negatively affected. We operate in a highly competitive, evolving and rapidly changing industry and our failure to adapt could negatively impact our business. The health service industry continues to be dynamic and rapidly evolving. Any significant shifts in the structure of the industry could alter industry dynamics and adversely affect our ability to attract or retain clients and customers. Industry shifts could result (and have resulted) from, among other things: • a large intra- or inter- industry merger or industry consolidation; • strategic alliances; • new or alternative business models or new government options or offerings; • continuing consolidation among physicians, hospitals and other health care providers, as well as changes in the organizational structures chosen by physicians, hospitals and health care providers; • new market entrants, including those not traditionally in the health service industry; • the ability of larger employers and clients to contract directly with providers; • technological changes and rapid shifts in the use of technology, such as telehealth and AI; • the impact or consequences of legislation or regulatory changes; • impacts to distribution channels, including changes <del>in to</del> the United States Postal Service or the consolidation of shipping carriers; • increased drug acquisition cost or unexpected changes to drug pricing trend; • changes in the generic / biosimilar drug market or the failure of new generic / biosimilar drugs to come to market; or • changes in utilization of health care, prescription drugs or other covered services and items, including under risk- based contracts in the health benefit management market and for those businesses that utilize risk adjustment methodology. Our failure to anticipate or appropriately adapt to changes in the industry could negatively impact our competitive position and adversely affect our business and results of operations. Our failure to compete effectively, to differentiate our products and services from those of our competitors and maintain or increase market share, including maintaining or increasing enrollments in businesses providing health benefits, could materially adversely affect our results of operations, financial position and cash flows. We operate in a highly competitive environment and an industry subject to significant market pressures brought about by customer and client needs, legislative and regulatory developments and other market factors. In particular markets, our competitors may have greater, better or more established capabilities, resources, market share, reputation or business relationships, or lower profit margin or financial return expectations. Our clients are well informed and organized and can easily move between our competitors and us. Our Express Scripts client contracts generally have three- year terms and may be subject to periodic renegotiation of pricing terms based on market factors. As described in greater detail in the description of our business in Item 1 of this Form 10- K, one of our key clients in the Evernorth Health

Services segment <del>is <mark>include</mark> t</del>he <del>United States Department of Defense-<mark>DoD, Prime and Centene</mark> . If one or more of our large</del> clients terminates or does not renew a contract for any reason, including as a result of being acquired, or if the provisions of a contract with a large client are modified, renewed or otherwise changed with terms less favorable to us, our results of operations could be adversely affected and we could experience a negative reaction in the investment community resulting in decreases in the trading price of our securities or other adverse effects. Our success depends, in part, on our ability to compete effectively in our markets, set prices appropriately in highly competitive markets to keep or increase our market share, increase customers as planned, differentiate our business offerings by innovating and delivering products and services that provide enhanced value to our customers, provide quality and satisfactory levels of service and retain accounts with favorable medical cost experience or more profitable products versus retaining or increasing our customer base in accounts with unfavorable medical cost experience or less profitable products. We must remain competitive to attract new customers, retain existing customers and further integrate additional product and service offerings. To succeed in this highly competitive marketplace, it is imperative that we maintain a strong reputation. Increasingly, our customers, clients and investors consider our efforts on a variety of matters that could impact our stakeholders, including our employees and the communities in which we operate, such as our efforts with respect to the environment and diversity, equity and inclusion. The negative reputational impact of a significant event, including a failure to execute on customer or client contracts or strategic or operational initiatives, failure to comply with applicable laws or regulations, or failure to innovate and deliver products and services that demonstrate greater value to our customers, could affect our ability to grow and retain profitable arrangements, which could have a material adverse effect on our business, results of operations, financial position and cash flows. We face price competition and other pressures that could compress our margins or result in premiums that are insufficient to cover the cost of services delivered to our customers. While we compete on the basis of many service and quality-related factors, we expect that price will continue to be a significant basis of competition and we may face pressure to contain premium rates. Our client contracts are subject to negotiation as clients seek to contain their costs, including by reducing benefits offered. Increasingly, our clients seek to negotiate performance guarantees that require us to pay penalties if the guaranteed performance standard is not met. Clients can easily move between our competitors and us. Our clients are well informed and typically have knowledgeable consultants that seek competing bids from our competitors before contract renewal. In addition, as brokers and benefit consultants seek to enhance their revenue streams, they look to take on services that we typically provide. Each of these events could negatively impact our financial results. Federal and state regulatory agencies may restrict or prevent entirely our ability to implement changes in premium rates. Fiscal or other concerns related to the government- sponsored programs in which we participate, such as Medicare Advantage plans and Medicare Part D plans, may cause decreasing reimbursement rates, delays in premium payments, restrictions on implementing changes in premium rates or insufficient increases in reimbursement rates. Any limitation on our ability to maintain or increase our premium or reimbursement levels, or a significant loss of customers or clients resulting from our need to increase or maintain premium or reimbursement levels, could adversely affect our business, cash flows, financial condition and results of operations. Premiums in the Cigna Healthcare segment are generally set for one- year periods and are priced well in advance of the date on which the contract commences or renews. Our revenue on Medicare Advantage plans, Individual and Family Plans (" IFP") and Medicare Part D plans is based on rates and bids submitted midyear in the year before the contract year. Although we base the premiums we charge and our Medicare Advantage, IFP and Medicare Part D rates and bids on our estimate of future health care costs over the contract period, actual costs may exceed what we estimate in setting premiums. Our participation in health insurance exchanges through our IFP offerings involves uncertainties associated with mix and volume of business and could adversely affect our results of operations, financial position and cash flows. Our health care costs also are affected by external events that we cannot forecast or project and over which we have little or no control, including changes in laws and regulations, as well as pandemics, costly new treatments, new treatment guidelines, provider billing practices, inflation and changes in customers' health care utilization patterns, which may, among other things, impact our ability to appropriately document their health conditions. Our profitability depends, in part, on our ability to accurately predict, price for and effectively manage future health care costs. Relatively small differences between predicted and actual medical costs or utilization rates as a percentage of revenue can result in significant changes in our financial results. Strong competition within the pharmacy benefit business has also generated greater demand for lower product and service pricing, increased revenue sharing and enhanced product and service offerings. These competitive factors have historically applied pressure on our operating margins and caused many companies, including us, to reduce the prices charged for products and services while sharing with clients a greater portion of the formulary fees and related rebates received from pharmaceutical manufacturers. Our inability to maintain positive trends, or failure to identify and implement new ways to mitigate pricing pressures, could negatively impact our ability to attract or retain clients or sell additional services, which could negatively impact our margins and have a material adverse effect on our business and results of operations. In addition, legislative reforms related to rebates, reporting, and other activities may adversely affect our competitive position, cash flows, financial condition and results of operations. The reserves we hold for expected medical claims are based on estimates that involve an extensive degree of judgment and are inherently variable. If actual claims exceed our estimates, our operating results could be materially adversely affected, and our ability to take timely corrective actions to contain future costs may be limited. We maintain and record medical claims reserves in our Consolidated Balance Sheets for estimated future payments. Our estimates of health care costs payable are based on a number of factors, including historical claim experience, but this estimation process requires extensive judgment. Considerable variability is inherent in such estimates, and the accuracy of the estimates is highly sensitive to changes in medical claims submission and processing patterns or procedures, changes in customer base and product mix, changes in the utilization of prescription drugs, medical or other covered items or services, changes in medical cost trends, changes in our health management practices, changes in regulations and the introduction of new benefits and products. If we are not able to accurately and promptly anticipate and detect medical cost trends, our ability to take timely corrective actions to limit future costs and reflect our current benefit cost experience in our

pricing process may be limited. Additionally, we must estimate the amount of rebates payable by us under the ACA's and CMS' minimum loss ratio rules and the amounts payable by us to, and receivable by us from, the United States federal government under the ACA's remaining premium stabilization program. Because establishing reserves is an inherently uncertain process involving estimates of future losses, there can be no certainty that ultimate losses will not exceed existing reserves which may adversely affect our results of operations, financial position and cash flows. If we fail to develop and maintain satisfactory relationships with health care payers, physicians, hospitals and other health service providers and with producers and consultants, our business and results of operations may be adversely affected. We contract with or employ physicians, hospitals and other health service providers and facilities to provide health services to our customers, as well as health care payers (as a service provider to those payers). Our results of operations are substantially dependent on our ability to contract for these services at competitive prices. In any particular market, physicians, hospitals and health service providers may enter into exclusive arrangements with competitors or simply refuse to contract with us, demand higher payments or take other actions that could result in higher medical costs or less desirable products or services for our customers. In some markets, certain providers, particularly hospitals, physician / hospital organizations and multispecialty physician groups, may have significant or controlling market positions that could result in a diminished bargaining position for us. If providers refuse to contract with us, use their market position to negotiate more favorable contracts or place us at a competitive disadvantage, our ability to market products or to be profitable in those areas could be materially adversely affected. Additionally, certain regulations may impact our ability to obtain competitive prices. Establishing collaborative arrangements with physician groups, specialist groups, independent practice associations, hospitals and health care delivery systems is key to our strategic focus to transition from volume-based fee- for- service arrangements to a value- based health care system. If such collaborative arrangements do not result in the lower medical costs that we project or if we fail to attract health care providers to such arrangements, or are less successful at implementing such arrangements than our competitors, our attractiveness to customers may be reduced and our ability to profitably grow our business may be adversely affected. Our ability to develop and maintain satisfactory relationships with providers may also be negatively impacted by other factors not associated with us, such as changes in Medicare or Medicaid reimbursement levels, increasing pressure on revenue and other pressures on health care providers and increasing consolidation activity among hospitals, physician groups and providers. Continuing consolidation among physicians, hospitals and other providers, the emergence of accountable care organizations, vertical integration of providers and other entities, changes in the organizational structures chosen by physicians, hospitals and providers, new market entrants, including those not traditionally in the health care industry, and the increased use of virtual care services (including telehealth) may affect the way providers interact with us and may change the competitive landscape in which we operate. In some instances, these organizations may compete directly with us, potentially affecting the way we price our products and services or causing us to incur increased costs if we change our operations to be more competitive. Out- of- network providers for non- Medicare services are not limited by any agreement with us in the amounts they bill . For Medicare Advantage, out- of- network providers can only receive the same rate that CMS pays for Medicare services. While benefit plans place limits on the amount of charges that will be considered for reimbursement and regulations seek to prescribe payment levels, establish methodologies and dispute resolution processes, providers are increasingly sophisticated and aggressive. As a result, the outcome of disputes where we do not have a provider contract may cause us to pay higher medical or other benefit costs than we projected. Additionally, certain of our products and services are sold in part through non-nonexclusive --- exclusive producers and consultants for whose services and allegiance we compete. Our sales could be materially adversely affected if we are unable to attract, retain and support such independent producers and consultants or if our sales strategy is not appropriately aligned across distribution channels. If we lose our relationship with one or more key pharmaceutical manufacturers, or if the payments made or discounts provided by pharmaceutical manufacturers decline, our business and results of operations could be adversely affected. We maintain relationships with numerous pharmaceutical manufacturers, which provide us with, among other things: • discounts for drugs we purchase to be dispensed from our home delivery and specialty pharmacies; • discounts, in the form of rebates, for drug utilization; • fees for administering rebate programs, including invoicing, allocating and collecting rebates; • fees for services provided to pharmaceutical manufacturers by our specialty pharmacies; and • access to limited distribution specialty pharmaceuticals by our specialty pharmacies. Our contracts with pharmaceutical manufacturers are typically non-nonexclusive --- exclusive and terminable on relatively short notice by either party. The consolidation of pharmaceutical manufacturers, the termination or material alteration of our relationships, or our failure to renew contracts on market competitive terms could have a material adverse effect on our business and results of operations. In addition, arrangements between payors and pharmaceutical manufacturers have been the subject of debate in federal and state legislatures and various other public and governmental forums. Adoption of new laws, rules or regulations or changes in, or new interpretations of, existing laws, rules or regulations, relating to any of these programs could materially adversely affect our business and results of operations. If significant changes occur within the pharmacy provider marketplace, or if other issues arise with respect to our pharmacy networks, including the loss of or adverse change in our relationship with one or more key pharmacy providers, our business and financial results could be adversely affected. More than 67,000 pharmacies participated in one or more of our networks as of December 31, 2022-2023 . The ten largest retail pharmacy chains represent approximately 60 % of the total number of stores in our largest network. In certain geographic areas of the United States, our networks may be comprised of higher concentrations of one or more large pharmacy chains. Contracts with retail pharmacies are generally non-nonexelusive--- exclusive and are terminable on relatively short notice by either party. If one or more of the larger pharmacy chains terminates its relationship with us, or is able to renegotiate terms substantially less favorable to us, our customers' access to retail pharmacies or our business could be materially adversely affected. The entry of one or more additional large pharmacy chains into the pharmacy benefit management business, the consolidation of existing pharmacy chains or increased leverage or market share by the largest pharmacy providers could increase the likelihood of negative changes in our relationship with such pharmacies. Changes in the overall composition

```
of our pharmacy networks, or reduced pharmacy access under our networks, could have a negative impact on our claims volume
or our competitiveness in the marketplace, which could cause us to fall short of certain guarantees in our contracts with clients or
otherwise impair our business or results of operations. Changes in drug pricing or industry pricing benchmarks could materially
impact our financial performance. Contracts in the prescription drug industry, including our contracts with retail pharmacy
networks and our pharmacy and specialty pharmacy clients, generally use pricing metrics published by third parties as
benchmarks to establish pricing for prescription drugs. If these benchmarks are no longer published by third parties, we, or our
contractual partners, adopt other pricing benchmarks for establishing prices within the industry, legislation or regulation
requires the use of other pricing benchmarks, or future changes in drug prices substantially deviate from our expectations, the
short- or long- term impacts may have a material adverse effect on our business and results of operations. Additionally, laws
such as the Inflation Reduction Act have granted CMS the ability to negotiate drug prices for certain Part D and Part B
drugs, and other federal and state legislative proposals may lead to changes in drug pricing for federal health care
programs. Our business depends on our ability to effectively invest in, implement improvements to and properly maintain the
uninterrupted operation, availability and data integrity of our information technology and other business systems. Our business
is highly dependent on maintaining effective information systems as well as the integrity and timeliness of the data we use to
serve our customers and health care providers and to operate our business. If our data were found to be inaccurate or unreliable
due to fraud or other error, or if we, or any of the third- party providers or subcontractors that we or they engage, were to fail
to maintain information systems and data integrity effectively, we could experience operational disruptions that may impact our
clients, customers and health care providers and hinder our ability to provide or establish appropriate pricing for products and
services, retain and attract clients and customers, establish reserves and report financial results timely and accurately and
maintain regulatory compliance, among other things. Our information technology strategy and execution are critical to our
continued success. We must continue to invest in and maintain long- term solutions that will enable us to anticipate customer
needs and expectations, enhance the customer experience, act as a differentiator in the market and protect against cybersecurity
risks and threats or other events that could disrupt our information technology systems such as man-made or natural disasters
(including those as a result of climate change). Our success is dependent, in large part, on maintaining the effectiveness of
existing technology systems and continuing to deliver and enhance technology systems that support our business processes in a
cost- efficient and resource- efficient manner. Increasing regulatory and legislative changes will place additional demands on
our infrastructure that could have a direct impact on resources available for other projects tied to our strategic initiatives. In
addition, recent trends toward greater consumer engagement in health care require new and enhanced technologies, including
more sophisticated applications for mobile devices. Connectivity among technologies is becoming increasingly important. We
must also develop new systems to meet current market standards and keep pace with continuing changes in information
processing technology, evolving industry and regulatory standards and customer needs. Failure to do so may present compliance
challenges and impede our ability to deliver services in a competitive manner. Further, because system development projects are
long- term in nature, they may be more costly than expected to complete and may not deliver the expected benefits upon
completion. Our failure to effectively invest in, implement improvements to and properly maintain the uninterrupted operation,
availability and data integrity of our systems could adversely affect our results of operations, financial position and cash flow.
As a large global health company, we and our vendors are subject to cyberattacks or other privacy or data security incidents. If
we are unable to prevent or contain the effects of any such attacks, or fail to ensure vendors do the same, we may suffer
exposure to substantial liability, reputational harm, loss of revenue or other damages. Our business depends on our clients' and
customers' willingness to entrust us with their health- related and other sensitive-personal information ("PI"), including
Protected Health information Information ("PHI") that is subject to privacy, security or data breach notification laws.
Computer networks or systems may be vulnerable to intrusion physical break- ins., computer viruses or malware,
programming errors, attacks by third parties or similar disruptive problems. We have been, and will likely continue to be, the
target of computer viruses or other malicious codes, unauthorized access, cyberattacks or other computer- related penetrations.
There have been, and will likely continue to be, large scale cyberattacks within the health service industry. Additionally,
hardware, software or applications we develop or procure from third parties may contain defects in design, manufacturer defects
or other problems that could unexpectedly compromise information technology. Human or technological error has and could in
the future result in, for example, unauthorized access to, acquisition, disclosure, modification, misuse, loss, or destruction of
company, customer, or other third- party data or systems; theft of sensitive, regulated, or confidential data including PI personal
information and intellectual property; the loss of access to critical data or systems through ransomware, destructive attacks or
other means; and business delays, service or system disruptions or denials of service. As we increase the amount of PI personal
information that we store and share digitally, our exposure to unauthorized uses and disclosures, and data privacy and related
cybersecurity risks increases, including the risk of undetected attacks, damage, loss or unauthorized access or acquisition or
misappropriation of proprietary or personal information, and the cost of attempting to protect against these risks also increases.
The health care data ecosystem is complex and requires data exchange with vendors, business partners, health care
professionals, the government and others. If disruptions, data disclosures, security incidents or breaches are not detected
quickly, their effect could be compounded. We have dedicated significant resources to implement privacy and security
technologies, processes and procedures to protect PI consumer identity and provide employee awareness training around
phishing, malware and other cyber risks; however, there are no assurances that such measures will be effective against all types
of security incidents or breaches. Further, we depend on many vendors to support and assist our business, which requires such
vendors to generate, store and use PI sensitive personal information. Cybersecurity threats are rapidly evolving and those
threats and the means for obtaining access to our proprietary systems are becoming increasingly sophisticated. Cyberattacks can
originate from a wide variety of sources including terrorists, nation states, internal actors, or third parties, such as external
service providers, and the techniques used change frequently or are often not recognized until after they have been launched. For
```

```
example, there <del>has been <mark>continues to be</mark> an increase in new financial fraud schemes akin to ransomware attacks on large</del>
companies whereby a cybercriminal installs a type of malicious software, or malware, that prevents a user or enterprise from
accessing computer files, systems or networks and demands payment of a ransom for their return. Those parties may also
attempt to fraudulently induce employees, customers or other users of our systems to disclose sensitive information or
inadvertently provide access to systems in order to gain access to our data or that of our customers. In addition, while we have
certain standards for all vendors that provide us services, our vendors, and in turn, their own service providers, may become
subject to the same types of security breaches. Finally, our offices may be vulnerable to security incidents or security attacks,
acts of vandalism or theft, misplaced or lost data, human error or similar events that could negatively affect our systems and our
customers' and clients' data. The costs to eliminate or address security threats and vulnerabilities before or after a cyber-incident
could be significant. Our remediation efforts may not be successful and could result in interruptions, delays, or cessation of
service and loss of existing or potential customers. In addition, the unauthorized access, acquisition, use, disclosure or
dissemination of sensitive-personal information, proprietary information or confidential information about us, our customers or
other third parties could expose our customers' and their private information to the risk of financial or medical identity theft.
Unauthorized access, acquisition, use, disclosure or dissemination of confidential and proprietary information about our business
and strategy could also negatively affect the achievement of our strategic initiatives. Such events could cause us to breach our
contractual obligations and violate applicable laws. These events would negatively affect our ability to compete, our reputation,
customer base and revenues and expose us to mandatory disclosure requirements, government investigations, litigation and other
enforcement proceedings, material fines, penalties or remediation costs and compensatory, special, punitive and statutory
damages, consent orders and other adverse actions, any of which could adversely affect our business, results of operations,
financial condition or liquidity. Our use The scale, scope and duration of artificial intelligence the ongoing COVID-19
pandemic continues to be unknown and the overall impact on machine learning present regulatory and legal challenges that
<mark>could negatively affect</mark> our business <mark>and , operating results, eash flows or our reputation financial condition has been and may</mark>
continue to be material. Our use The COVID-19 pandemic has adversely affected, and is continuing to affect, global
economics, financial markets and the overall environment for our business, and the extent to which it may impact our future
results-of artificial intelligence ("AI") operations and overall financial performance remains uncertain. While vaccination
rates continue to rise, the COVID-19 pandemie, including vaccination efficacy machine learning ("ML") technologies, the
implementation of and reaction to vaccination and testing mandates and the occurrence of new variants, could continue to effect
such economics and financial markets as well as the health more recent technological advances in AI / ML, pose risks to us
and availability of our workforce subject us to new and existing laws and regulations. As While we are committed to
responsible use of AI / ML and following applicable laws and regulations, and while we have made progress developing
governance as to use of AI / ML by our organization, any failure to use AI / ML responsibly and to adhere to such laws,
regulations and governance could have a material unfavorable effect on our business, result-results of operations, and
financial condition. Depending on how existing laws and regulations are interpreted, and as new laws are passed, we
may experience new disruptions have to make changes to our business practices to comply with such obligations. These
obligations may make it harder for us to conduct our business using AI / ML, lead to regulatory fines or penalties,
require us retrain our AI / ML, or prevent or limit our use of AI / ML. Our use of AI / ML technologies could also result
in additional compliance costs, regulatory investigations and actions, and consumer or other lawsuits. If we are unable to
use AI / ML, or if regulators restrict our ability to use AI / ML for certain purposes, it could make our business less
efficient, result in competitive disadvantages, and subject us to potential unfavorable business impacts. To the extent that
we rely on or use the output of AI / ML, any inaccuracies, biases or errors could have unfavorable impacts on us, our
business and our results of operations and our business could be adversely affected further, directly or indirectly, by the
ongoing pandemic. The COVID-19 pandemic has in some instances, and may continue to, heighten the potential adverse effects
on our business, operating results, eash flows or financial condition. The as described below or in other risk factors within this
section of the Form 10-K including, but not limited to, the likelihood of and impact from: • unfavorable economic conditions on
our clients and customers (both employers and individuals), health care and pharmacy providers, pharmaccutical manufacturers
and third-party vendors, as well as federal and state entities and programs; • changes in medical claims submission and
processing patterns or procedures; changes in customer base and product mix; changes in utilization of regulatory prescription
drugs, medical or other covered items or services, including increased behavioral health services utilization; changes in medical
eost trends; changes in our health management practices; and legal the introduction of new benefits and products causing actual
elaims to exceed our estimates; • changes in health care utilization patterns, provider billing practices and other external events
that we cannot forecast or project and over which we have little or no control impacting our ability to accurately predict, price
for and manage health care costs and ultimately our profitability, including impacts from care deferral on, among other things,
risk risks adjustment revenue and acuity of future care; * increased costs or reductions in revenue, including costs for COVID-
19-related care, testing and treatment; vaccine and other coverage mandates; inflation; and support for employees, clients,
eustomers and providers; • compliance with substantial government regulation, including privacy and security requirements
associated with AI / ML providing telehealth and remote care options and new laws or regulations or changes in existing laws or
regulations, such as vaccine, testing and coverage mandates and premium deferrals, which laws or regulations may vary
significantly by jurisdiction; • cyberattacks or other privacy or data security incidents, including as a result of the transition to a
hybrid work environment by substantially all of our workforce and the workforces of third parties with whom we contract; •
significant shifts in the structure of the industry which could alter dynamics and, if we fail to adapt, negatively impact our
business; • risks inherent in foreign operations, including political, legal, operational, regulatory, economic and other risks; •
economic and market conditions affecting the value of our financial instruments and the value of particular assets and liabilities;
and • fluctuations in equity market prices, interest rates and credit spreads limiting our ability to raise or deploy capital and
```

```
affecting our overall liquidity. We believe COVID-19 and its is variants adverse impact on our business, operating results,
eash flows or financial condition will be driven primarily by the severity and duration of the pandemic, including the impact of
the breadth and timing of implementation and the efficacy and costs of vaccination programs, the pandemic's continued impact
on our employees, clients, customers, suppliers and partners, as well as the U. S. and global economics and the continued
actions taken by governmental authorities and other third parties in response to the pandemic. Those primary drivers are largely
unknown beyond our knowledge and control, and may be more adverse than our current expectations. Given these
uncertainties, we cannot estimate the full impact COVID-19 will have on our business, operating results, cash flows or financial
condition, but the adverse impact could be material. As a global company, we face political, legal, operational, regulatory,
economic and other risks that present challenges and could negatively affect our multinational operations or our long-term
growth. As a global company, our business is increasingly exposed to risks inherent in foreign operations. These risks can vary
substantially by market, and include political, legal, operational, regulatory, economic and other risks, including government
intervention that we do not face in our U. S. operations. The global nature of our business and operations may present challenges
including, but not limited to, those arising from: • geopolitical business conditions and demands; • regulation that may
discriminate against U. S. companies, favor nationalization or expropriate assets; • price controls or other pricing issues and
exchange controls; restrictions that prevent us from transferring funds out of the countries in which we operate; foreign currency
exchange rates and fluctuations and restrictions on converting currencies from foreign operations into other currencies;
uncertainty with respect to the adoption of new tax laws and the interpretation of tax positions; • reliance on local employees
and interpretations of labor laws in foreign jurisdictions; • managing our partner relationships in countries outside of the United
States; • providing data protection on a global basis and sufficient levels of technical support in different locations; • the global
trend for companies to enact local data residency requirements; • acts of civil unrest, war and terrorism, including the ongoing
conflict in the Middle East as well as other political and economic conflicts such as through imposition of economic or political
sanctions; • man- made disasters, natural disasters (including those arising as a result of climate change) and pandemics , such as
the COVID-19 pandemie, in locations where we operate; and • general economic and political conditions, including
conditions that may become unpredictable during a U. S. presidential election year. These factors may increase in
significance as we continue to expand globally and operating in new foreign markets may require considerable management
time before operations generate any significant revenues and earnings. Any one of these challenges could negatively affect our
operations or long- term growth. International operations also require us to devote significant resources to implement controls
and systems in new markets to comply with, and to ensure that our vendors and partners comply with, U. S. and foreign laws
prohibiting bribery, corruption and money laundering, in addition to other regulations regarding, among other things, our
products, direct- to- consumer communications, customer privacy, data protection and data residency. Violations of these laws
and regulations could result in fines, criminal sanctions against us, our officers or employees, restrictions or outright prohibitions
on the conduct of our business and significant reputational harm. Our success depends, in part, on our ability to anticipate these
risks and manage these challenges. Our failure to comply with laws and regulations governing our conduct outside of the United
States or to establish constructive relations with non- U. S. regulators could have a material adverse effect on our business,
results of operations, financial condition, liquidity and long-term growth. Please see" — Legal and Compliance Risks"
below. Strategic transactions involve risks and we may not realize the expected benefits because of integration or separation
difficulties, underperformance relative to our expectations and other challenges. As part of our strategy, we regularly consider
and enter into strategic transactions, including mergers, acquisitions, joint ventures, licensing arrangements, divestitures and
other relationships (collectively referred to as" strategic transactions"). There is significant competition for attractive targets and
opportunities and we may be unable to identify and successfully complete strategic transactions in the future. In addition, from
time to time, we evaluate alternatives for our businesses that do not meet our strategic, growth or profitability objectives, and we
may divest or wind down such businesses. We may be unable to complete any such divestiture on terms favorable to us, within
the expected timeframes, or at all . For example, in January 2024 we announced the HCSC transaction, which is subject to
regulatory approvals and other closing conditions. We may be unable to satisfy the closing conditions in a timely manner
to complete the HCSC transaction, or we may otherwise fail to receive the anticipated benefits from the transaction, even
if it is completed. We may have continued financial exposure to divested businesses following the completion of any such
transaction, including increased costs due to potential litigation, contingent liabilities and indemnification of the buyer related to,
among other things, lawsuits, regulatory matters or tax liabilities. Our ability to achieve the anticipated benefits of strategic
transactions, including synergies, cost savings, innovation and operational efficiencies, is subject to numerous uncertainties and
risks, including our ability to successfully combine or separate business operations, resources and systems, including data
security systems and internal financial control standards, in an efficient and effective manner. Integration and separation
activities may result in additional and unforeseen expenses, and the anticipated benefits may not be fully realized or may take
longer to realize than expected. These activities are complex, costly and time- consuming and may divert management's
attention from ongoing business concerns. Delays or issues encountered in these activities could have a material adverse effect
on the revenues, expenses, operating results and financial condition of the Company. Additionally, the benefits of strategic
transactions and the related timing could be impacted by various factors, including political instability, natural disasters,
fluctuations in currency exchange rates, delays in obtaining regulatory approval and changes in regulations. Strategic
transactions could result in increased costs, including facilities and systems consolidation or separation costs and costs to retain
key employees, decreases in expected revenues, earnings or cash flows and goodwill or other intangible asset impairment
charges. As of December 31, <del>2022-2023</del>, our goodwill and other intangible assets had a carrying value of approximately $ 78-75
billion, representing 54.49 % of our total consolidated assets. The value of our goodwill may be materially and adversely
impacted if the businesses we acquire do not perform in a manner consistent with our assumptions. Future evaluations requiring
an impairment to goodwill and other intangible assets could materially affect our results of operations and shareholders' equity
```

in the period in which the impairment occurs. A material decrease in shareholders' equity could negatively impact our debt ratings or potentially impact our compliance with existing debt covenants. See Note 19-20 to the Consolidated Financial Statements for more information on goodwill and intangibles. In addition, the trading price of our securities may decline if, among other things, we are unable to achieve our estimates of earnings growth and operational cost savings, or the transaction costs are greater than expected. The trading price also may decline if we do not achieve the perceived benefits of a transaction as rapidly or to the extent anticipated by financial or industry analysts. Additionally, joint ventures and equity investments present risks that are different from acquisitions, including risks related to: specific operations and finances of the businesses we invest in; selection of appropriate parties; differing objectives of the various parties; competition between and among parties; compliance activities (including compliance with applicable CMS requirements); growing the business in a manner acceptable to all the parties; maintaining positive relationships among the parties, clients and customers; initial and ongoing governance of joint ventures and customer and business disruption that may occur upon a joint venture termination. Further, we may finance strategic transactions by issuing common stock for some or all of the purchase price that could dilute the ownership interests of our shareholders, or by incurring additional debt that could increase costs and impact our ability to access capital in the future. In addition, effective internal controls are necessary to provide reliable and accurate financial reports and to mitigate the risk of fraud. The integration of businesses is likely to cause increasing complexity in our systems and internal controls and make them more difficult to manage. Any difficulties in assimilating businesses into our control system could cause us to fail to meet our financial reporting obligations. We also rely on the internal controls and financial reporting controls of joint venture entities and other entities in which we invest and their failure to maintain effectiveness or comply with applicable standards may materially and adversely affect us. Ineffective internal controls could also cause investors to lose confidence in our reported financial information that could negatively impact the trading price of our securities and our access to capital. We are dependent on the success of our relationships with third parties for various services and functions. To improve operating costs, productivity and efficiencies, we contract with third parties for the provision of specific services. Our operations may be adversely affected if a third party fails to satisfy its obligations, if the arrangement is terminated in whole or in part or if there is a contractual dispute between us and the third party. Even though contracts are intended to provide certain protections, we have limited control over the actions of third parties. For example, noncompliance with any privacy or security laws and regulations, any security breach involving one of our third- party vendors or a dispute between us and a third- party vendor related to our arrangement could have a material adverse effect on our business, results of operations, financial condition, liquidity and reputation. Outsourcing also may require us to change our existing operations, adopt new processes for managing these service providers or redistribute responsibilities to realize the potential productivity and operational efficiencies. If there are delays or difficulties in changing business processes or our third- party vendors do not perform as expected, we may not realize, or not realize on a timely basis, the anticipated economic and other benefits of these relationships. This could result in additional costs or regulatory compliance issues or create other operational or financial problems for us. Terminating or transitioning, in whole or in part, arrangements with key vendors could result in additional costs or penalties, risks of operational delays or potential errors and control issues during the termination or transition phase. We may not be able to find an alternative vendor in a timely manner or on acceptable terms. If there is an interruption in business or loss of access to data resulting from a security breach, termination or transition in services, we may not be able to meet the demands of our customers and, in turn, our business and results of operations could be adversely impacted. A significant disruption in service within our operations or among our key suppliers or other third parties could materially adversely affect our business and results of operations. Our business is highly dependent upon our ability to perform, in an efficient and uninterrupted fashion, necessary business functions, such as claims processing and payment, internet support and customer call centers, data centers and corporate facilities, processing new and renewal business, maintaining appropriate shipment and storage conditions for prescriptions (such as temperature and protection from contamination) and home delivery processing. In some instances, our ability to provide services or products (including processing and dispensing prescriptions) depends on the availability of services and products provided by suppliers, providers, pharmaceutical manufacturers, vendors or shipping carriers. A disruption, or threat of disruption, in our supply chain, including as a result of future the COVID-19 pandemic pandemics or public health emergencies, or inability to access or deliver products that meet requisite quality safety standards and patient needs in a timely and efficient manner could adversely impact our business. Increasing natural disasters in connection with climate change could also be a direct threat to us and our third- party vendors, service providers or other stakeholders. Natural disasters, such as wildfires, hurricanes and snow and ice storms, have impacted and may continue to impact our customers and pose a risk to our employees and facilities located in the impacted region. Responses to such scenarios have and may include, among other things, making temporary policy changes, such as waiving various medical requirements, assisting with replacement medications, transferring prescriptions and expanding our help line. In addition, there is a risk that actions taken to respond to climate change could increase the cost of energy, fuel and other commodities, which would increase our operating costs. We are also subject to risk as a result of information technology disruptions. Any failure or disruption of our performance of, or our ability to perform, key business functions, including through unavailability or cyberattack of our information technology systems or those of third parties (including cloud service providers), could cause slower response times, decreased levels of service satisfaction and harm to our reputation. Our systems interface with and depend on third- party systems and we could experience service denials if demand for such service exceeds capacity or a third- party system fails or experiences an interruption. While we have adopted, and continue to enhance, business continuity and disaster recovery plans and strategies, there is no guarantee that such plans and strategies will be effective, which could interrupt the functionality of our information technology systems or those of third parties. Our failure to implement adequate business continuity and disaster recovery strategies could significantly reduce our ability to provide products and services to our customers and clients, which could have material adverse effects on our business and results of operations. In managing medical practices and operating pharmacies, onsite clinics and other types of medical facilities, we may be subject to additional liability

that could result in significant time and expense. In addition to contracting with physicians and other health care providers for services, we employ physicians, pharmacists, nurses and other health care providers at our home delivery and specialty pharmacies, onsite low acuity and primary care practices and infusion clinics that we manage and operate for our customers, as well as certain clinics for our employees. We also provide in-home care through health care providers that we employ, as well as through third- party contractors. As such, we may be subject to liability for certain acts, omissions, or injuries caused by our employees or agents, or occurring at one of these practices, pharmacies or clinics. The defense of any actions may require diverting personnel and other resources and incurring significant costs that could have a material adverse effect on our business, results of operations, financial condition, liquidity and reputation. Legal and Compliance Risks-Our business is subject to substantial government regulation, as well as new laws or regulations or changes in existing laws or regulations that could have a material adverse effect on our business, results of operations, financial condition and liquidity. Our business is regulated at the federal, state, local and international levels. The laws and rules governing our business and related interpretations are increasing in number and complexity, are subject to frequent change and can be inconsistent or in conflict with each other. Noncompliance with applicable regulations by us or our third- party vendors could have material adverse effects on our business, results of operations, financial condition, liquidity and reputation. We must identify, assess and respond to new trends in the legislative and regulatory environment, as well as comply with the various existing regulations applicable to our business. There are currently pending, and in the future there will likely be, legislative or regulatory proposals which seek to manage the health services industry, including managing prescription drug costs and health records, as well as regulating drug distribution. Federal and state governments have enacted and we expect federal and state governments to continue to enact and seriously consider many broad-based legislative and regulatory proposals that will or could materially impact various aspects of the health care and related benefits system. In addition, changes to government policies not specifically targeted to the health services industry, such as a change in tax laws and the corporate tax rate or government spending cuts, could have significant impacts on our business, results of operations, financial condition and liquidity. The trading price of our securities may react to the announcement of such proposals. As disclosed in Part II, Item 5 of this Form 10- K, we have an active share repurchase program authorized by our board of directors. Regulators, customers, investors, employees and other stakeholders are increasingly focusing on ESG matters and related disclosures. These changing rules, regulations and stakeholder expectations have resulted in, and are likely to continue to result in, increased general and administrative expenses and increased management time and attention spent complying with such regulations or meeting such expectations. For example, the European Union's ("EU's") Corporate Sustainability Reporting Directive ("CSRD") will require expansive disclosures on various sustainability topics such as climate change, biodiversity, workforce, supply chain, and business ethics by in-scope EU entities and certain non- EU entities with significant cross- border business in EU markets. In addition, California's recently- enacted Climate Corporate Data Accountability Act will require annual disclosures of covered companies' Scope 1, 2 and 3 greenhouse gas emissions. We are assessing our obligations under CSRD and other enhanced reporting requirements, and expect that compliance could require substantial effort in the future. Overall, ESG matters and related stakeholder reaction may impact our reputation and have other business impacts which could adversely affect our business. Existing or future laws, rules, U. S. Presidential Executive Orders, regulatory interpretations or judgments could force us to change how we conduct our business, affect the products and services we offer and where we offer them, restrict revenue and enrollment growth, increase our costs, including medical, operating, health care technology and administrative costs, and require enhancements to our compliance infrastructure and internal controls environment. For example, health care reforms or the invalidation, modification, repeal or replacement of the ACA or portions thereof could result in material changes to the way we conduct our business, as well as the loss of subsidies related to our IFP offerings and could impact the market for our products. We are required to obtain and maintain insurance and other regulatory approvals to, among other things, market many of our products, expand into additional geographic or product markets, increase prices for certain regulated products and consummate some of our acquisitions and dispositions. Delays in obtaining or failure to obtain or maintain these approvals could reduce our revenue or increase our costs. Additionally, we must maintain licenses and registrations in the jurisdictions in which we conduct business, and the suspension, material adverse modification or termination of such license and registrations could adversely affect our operations. Such licensure subjects many of our businesses to state regulation of our operations and products, as well as risks associated with doing business in those jurisdictions. Existing or future laws and rules could also require or lead us to take other actions such as changing our business practices, and could increase our liability. Further, failure to effectively implement or adjust our strategic and operational initiatives, such as by reducing operating costs, adjusting premium pricing or benefit design or transforming our business model in response to regulatory changes may have a material adverse effect on our results of operations, financial condition and cash flows. For more information on regulations affecting our business, see" Business - Regulation" in Part I, Item 1 of this Form 10- K. There are various risks associated with participating in government- sponsored programs, such as Medicare, including dependence upon government funding, compliance with government contracts and increased regulatory oversight and enforcement. Through our U. S. Government Healthcare business, we contract with CMS and various state governmental agencies to provide managed health care services including Medicare Advantage plans and Medicare Part D plans. Additionally, our Evernorth Health Services business provides services to government entities and payors participating in government health care programs and our relationships with these government entities is subject to laws and regulations regarding government contracts. Our revenues from government-funded programs, including our Medicare programs and our government clients, are dependent, in whole or in part, upon annual funding from the federal government or applicable state or local governments. Funding for these programs is dependent on many factors outside our control, including general economic conditions, continuing government efforts to contain health care costs, budgetary constraints at the federal or applicable state or local level and general political issues and priorities. These entities generally have the right to not renew or to cancel their contracts with us on short notice without cause or if funds

are not available. Unanticipated changes in funding, such as the application of sequestration by the federal or state governments, retroactive rate adjustments, a delay by Congress in raising the federal debt ceiling, or the failure to provide for continued appropriations or regular ongoing scheduled payments to us, could substantially reduce our revenues or profitability or impact our liquidity. The Medicare program has been the subject of regulatory reform initiatives. The premium rates paid to Medicare Advantage plans and Medicare Part D plans are established by contract, although the rates differ depending on a combination of factors, some of which are outside our control. For example, the base premium rate paid differs depending upon a combination of various factors such as defined upper payment limits, a member's health status, age, gender, county or region, benefit mix, member eligibility category and risk scores. Additionally, a portion of each Medicare Advantage plan's reimbursement is tied to the plan's Star Rating, with those plans receiving a rating of four or more stars eligible for quality-based bonus payments. A plan's Star Rating affects its image in the market and plans that perform well are able to offer enhanced benefits, market more effectively and for longer periods of time than other plans. The Star Rating system is subject to change annually by CMS, which may make it more difficult to achieve four stars or greater. Our Medicare Advantage plans' and Medicare Part D plans' operating results, premium revenue and benefit offerings are likely to continue to be significantly determined by their Star Ratings. There can be no assurances that we will be successful in maintaining or improving our Star Ratings in future years. In addition, audits of our performance for past or future periods may result in downgrades to our Star Ratings. If we do not maintain or improve our Star Ratings or if the quality- based bonus payments are reduced or eliminated, we may experience a negative impact on our revenue and the marketability of our plans may be adversely affected. Accordingly, our plans may not be eligible for full level quality bonuses, which could adversely affect the benefits such plans can offer, reduce membership or impact our financial performance. See the" Executive Overview- Key Transactions and Business Developments" section of MD & A in Part II, Item 7 of this Form 10- K for additional information on our Star Ratings. Additionally, if we fail to comply with CMS' contractual requirements, including data submission, enrollment and marketing, provider network adequacy, provider directory accuracy, quality measures, claims payment, continuity of care, timely and accurate processing of appeals and grievances, adverse findings under RADV audits, oversight of first tier downstream and related entities and call center performance, we may be subject to administrative actions, including enrollment sanctions or contract termination, fines or other penalties or enforcement actions that could materially impact our profitability. Any failure, or alleged failure,....., results of operations and reputation. We face risks related to litigation, regulatory audits and investigations. We are routinely involved in numerous claims, lawsuits, regulatory audits, investigations and other legal matters arising, for the most part, in the ordinary course of business. These legal matters could include benefit claims, breach of contract actions, tort claims (including claims related to the delivery of health care services, such as medical malpractice by staff at our affiliates' facilities, or by health care practitioners who are employed by us, have contractual relationships with us, or serve as providers to our managed care networks, including as a result of a failure to adhere to applicable clinical, quality and / or patient safety standards), claims arising from consumer protection laws, false claims act laws, claims disputes under federal or state laws and disputes regarding reinsurance arrangements, employment and employment discrimination- related suits, antitrust claims (including as a result of changes in the enforcement of antitrust laws), employee benefit claims, wage and hour claims, tax, privacy, intellectual property and whistleblower claims, shareholder suits and other securities law claims, real estate disputes, claims related to disclosure of certain business practices and claims arising from customer audits and contract performance, including government contracts. In addition, we have incurred and likely will continue to incur liability for practices and claims related to our health care business, such as marketing misconduct, failure to timely or appropriately pay for or provide health care, provider network structure, poor outcomes for care delivered or arranged, provider disputes including disputes over compensation or contractual provisions, ERISA claims, allegations related to calculations of cost sharing and claims related to our administration of self-funded business. We are also routinely involved in legal matters arising from our health services business, including without limitation claims related to the dispensing of pharmaceutical products by our home delivery and specialty pharmacies, pharmacy benefit management services, such as formulary management services, health benefit management services and provider services. Our pharmacy services operations are subject to the clinical quality, patient safety and other risks inherent in the dispensing, packaging and distribution of drugs, including claims related to purported dispensing and other operational errors. There are currently, and may be in the future, attempts to bring class action lawsuits against the Company and other companies in our industry; individual plaintiffs also may bring multiple claims regarding the same subject matter against us and other companies in our industry. Court decisions and legislative activity may increase our exposure for any of these types of claims. In some cases, substantial noneconomic or punitive damages may be sought. We procure insurance coverage to cover some of these potential liabilities, however we also self- insure a significant portion of our litigation risks. While we maintain some third- party insurance coverage, including excess liability insurance with third- party insurance carriers, certain liabilities or types of damages, such as punitive damages, may not be covered by insurance, insurers may dispute coverage or the amount of insurance may be insufficient to cover the entire damages awarded. Resolving disputes is often expensive and disruptive, regardless of the outcome. Additionally, it is possible that the resolution of current or future legal matters and claims could result in changes to our industry and business practices, losses material to our results of operations, financial condition and liquidity or damage to our reputation. We are frequently the subject of regulatory market conduct and other reviews, audits and investigations by state insurance and health and welfare and pharmacy departments, attorneys general, DOJ, CMS, DOL and the HHS- OIG and comparable authorities in foreign jurisdictions. Additionally, we are have in the past been, and may in the future be, subject to qui tam actions in which the government may or may not intervene. With respect to our Medicare Advantage and Medicare Part D businesses, CMS and HHS- OIG perform audits to determine a health plan's compliance with federal regulations and contractual obligations, including compliance with proper coding practices and fraud and abuse enforcement practices through audits designed to detect and correct improper payments, Certain of our contracts are currently have subject to RADV audits by CMS and the HHS-OIG that are awaiting CMS finalization. These audits could result in <del>significant adjustments in payments repayments made t</del>o the

government our health plans, which could adversely affect our results of operations. There also continues to be heightened review by federal and state regulators of business and reporting practices within the health services industry, including with respect to claims payment and related escheat practices, and increased scrutiny by other federal and state governmental agencies (such as state attorneys general) empowered to bring criminal actions in circumstances that could have previously given rise only to civil or administrative proceedings. In addition, various government agencies have conducted investigations and audits into certain pharmacy benefit management practices. For example, the FTC is conducting an ongoing study of the pharmacy benefit manager industry and the impact of pharmacy benefit managers on the accessibility and affordability of prescription drugs. In June 2022, the FTC issued an enforcement policy statement indicating the FTC would scrutinize the impact of rebates and fees paid by pharmaceutical manufacturers to pharmacy benefit managers and other intermediaries to determine if laws such as the FTC Act, the Clayton Act, the Robinson- Patman Act and the Sherman Act may have been violated. In July 2023, the FTC voted to issue a statement cautioning against reliance on prior advocacy letters that advocated against proposals to increase regulatory oversight and transparency of pharmacy benefit managers. The FTC previously required three group purchasing organizations to provide information and records on business practices and the six largest pharmacy benefit managers to provide information and records on topics including rebate contracts and ancillary agreements, documents related to strategies, conditions and plans for formulary placement, formulary exclusion, formulary tier assignment, and prior authorization regarding rebated drug products, <mark>and annual pharmacy reimbursement data for drugs on specialty drug lists and for rebated drug products.</mark> Many <del>of these</del> investigations and audits have resulted in other companies being subject to civil penalties, including the payment of money and entry into corporate integrity agreements Any failure, or alleged failure, to comply with various state and federal health care laws and regulations, including those related to the CIA or otherwise directed at preventing fraud and abuse in government funded programs, has resulted in and could in the future result in investigations or litigation, such as actions under the federal False Claims Act and similar whistleblower statutes under state laws. A successful action or claim against us could subject us to damage awards, including treble damages, fines, penalties or other enforcement actions, restrictions on our ability to market or enroll new customers, limits on expansion, restrictions or exclusions from programs or other agreements with federal or state governmental agencies, which could adversely impact our business, cash flows, financial condition, results of operations and **reputation.** We cannot predict what effect, if any, such government investigations and audits may ultimately have on us or on the industry in general. However, we will likely continue to experience government scrutiny and audit activity, which has and may in the future result in civil penalties. Regulatory audits, investigations, litigation or reviews or actions by other government agencies have resulted in and could result in changes to our business practices, retroactive adjustments to certain premiums, significant fines, penalties, civil liabilities, criminal liabilities or other sanctions, including corporate integrity agreements, restrictions on our ability to participate in government programs or exclusion from such programs, market certain products or engage in business- related activities, that could have a material adverse effect on our business, results of operation, financial condition and liquidity. In addition, disclosure of an adverse investigation or audit or the imposition of fines or other sanctions could negatively affect our reputation in certain markets and make it more difficult for us to sell our products and services. A description of material pending legal actions and other legal and regulatory matters is included in Note 23-24 to the Consolidated Financial Statements included in this Form 10- K. The outcome of litigation and other legal or regulatory matters is always uncertain. If we fail to comply with applicable privacy, security and data laws, regulations and standards, our business and reputation could be materially adversely affected. Most of our activities involve the receipt, use, storage or transmission of a substantial amount of individuals' **PI, including** PHI and personally identifiable information. We also use aggregated and or anonymized data for research and analysis purposes, and in some cases, provide access to such anonymized de-identified data, or analytics created from such data, to pharmaceutical manufacturers and third- party data aggregators and analysts. We may also use such information to create analytic models designed to predict, and potentially improve, outcomes and patient care. The collection, dissemination, receipt, maintenance, protection, use, transmission, disclosure, privacy, confidentiality, security, availability, integrity, creation, processing, and disposal of PI sensitive personal information are regulated at the federal, state, international and industry levels and requirements are imposed on us by contracts with clients. In some cases, such laws, rules, regulations and contractual requirements also apply to our vendors and require us to obtain written assurances of their compliance with such requirements. We are also subject to various other consumer protection laws that regulate our communications with customers , such as the FTC Act and the Telephone Consumer Protection Act . Certain of our businesses are also subject to the Payment Card Industry Data Security Standard, which is designed to protect credit card account data as mandated by payment card industry entities. International laws, rules and regulations governing the use and disclosure of such information, such as the GDPR, can be more stringent than similar laws in the United States, and they vary across jurisdictions. In addition, more jurisdictions are regulating the transfer of data across borders and domestic privacy and data protection laws are generally becoming more onerous. These laws, rules and contractual requirements are subject to change and the regulatory environment surrounding data security and privacy is increasingly demanding. Compliance with existing or new privacy, security and data laws, regulations and requirements may result in increased operating costs, and may constrain or require us to alter our business model or operations. For more information on privacy regulations to which we are subject, see" Business - Regulation" in Part I, Item 1 of this Form 10- K. HIPAA requires covered entities and business associates to comply with the HIPAA privacy, security and breach rules. While we endeavor to provide appropriate protections through our contracts with our third- party service providers and in certain cases assess their security controls, we have limited oversight or control over their actions and practices. Several of our businesses act as business associates to their covered entity eustomers-clients and, as a result, collect, receive, use, disclose, transmit and maintain **PHI** sensitive personal information in order to provide services to these customers. HHS administers an audit program to assess HIPAA compliance efforts by covered entities and business associates. In addition, HHS continues to exercise its enforcement authority to bring enforcement actions resulting from

```
complaints, compliance reviews, audits and investigations brought on by notification to HHS of a breach <mark>or other HIPAA</mark>
violation. An audit resulting in findings or allegations of noncompliance or the implementation of an enforcement action could
have an adverse effect on our results of operations, financial position, cash flows and reputation. Noncompliance or findings of
noncompliance with applicable laws, regulations or requirements, or the occurrence of any privacy or security breach involving
the misappropriation, loss or other unauthorized disclosure of PI protected personal information, whether by us or by one of our
third- party service providers, could materially adversely affect our business and reputation, including our results of operations,
financial position and cash flows. Effective prevention, detection and control systems are critical to maintain regulatory
compliance and prevent fraud and; failure of these systems could adversely affect us. Federal and state governments have made
investigating and prosecuting health care and other insurance fraud and abuse a priority. Fraud and abuse prohibitions
encompass a wide range of activities including kickbacks for referral of customers, billing for unnecessary medical services,
improper marketing and violations of patient privacy rights. Some of our businesses are also subject to federal and state laws and
regulations that may impact our relationships with health care providers and customers, including laws on self- referrals,
beneficiary inducements, false claims, fee-splitting, telemedicine, corporate practice of medicine, dispensing, packaging,
fulfillment, and distribution of controlled substances, other pharmaceutical products and medical devices, medical malpractice,
consumer protection, product liability, narrow networks, provider tiering programs, provider contracts, overpayments,
reimbursement of out- of- network claims, and licensure. The regulations and contractual requirements applicable to us are
complex and subject to change and may affect our ability to market or provide our products or services. In addition, ongoing
vigorous law enforcement, a highly technical regulatory scheme and the Dodd- Frank Act and related regulations enhance
regulators' enforcement powers and whistleblower incentives and protections. Our compliance efforts in this area will continue
to require significant resources. Failure of our prevention, detection or control systems related to regulatory compliance or the
failure of employees to comply with our internal policies, including data systems security or unethical conduct by managers and
employees, could adversely affect our reputation and also expose us to litigation and other proceedings, fines and penalties. In
addition, provider or customer fraud that is not prevented or detected could impact our medical costs or those of our self-insured
clients. Further, during an economic downturn, we may experience increased fraudulent claims volume that may lead to
additional costs due to an increase in disputed claims and litigation. Economic Risks Economic and market conditions affect the
value of our financial instruments and the value of particular assets and liabilities, investment income and interest expense. As
an insurer, we have substantial investment assets that support insurance and contractholder deposit liabilities and surplus
requirements in our regulated companies. The market values of our investments vary depending on economic and market
conditions with no offsetting change in the value of a portion of our liabilities. A substantial portion of our investment assets are
in fixed interest- yielding debt securities of varying maturities and commercial mortgage loans. The value of these investment
assets can fluctuate significantly with changes in market conditions. A rise in interest rates would likely reduce the value of our
investment portfolio, increase interest expense on our indebtedness and increase investment income as investment assets mature
and are replaced. In addition, an economic contraction could result in delay in payment of principal or interest by issuers, or
defaults by issuers, reducing our investment income and requiring us to write down the value of our investments. Significant
stock market or interest rate declines could result in unfunded pension obligations resulting in the need for additional plan
funding by us and increased pension expenses. We currently have overfunded obligations in our frozen pension plans. A
significant decline in the value of the plans' equity and fixed income investments or unfavorable changes in applicable laws or
regulations could materially increase our expenses and change the timing and amount of required plan funding. This could
reduce the cash available to us, including our subsidiaries. We are also exposed to interest rate and equity risk associated with
our pension obligations. Sustained declines in interest rates could have an adverse impact on the funded status of our pension
plans and our reinvestment yield on new investments. See Note <del>17-18</del> to the Consolidated Financial Statements for more
information on our obligations under the pension plans. A downgrade in the financial strength ratings of our insurance
subsidiaries could adversely affect new sales and retention of current business, and a downgrade in our debt ratings would
increase the cost of borrowed funds and could negatively affect our ability to access capital. Financial strength, claims paying
ability and debt ratings by recognized rating organizations are each important factors in establishing the competitive position of
insurance and health benefits companies. Ratings information by nationally recognized <del>ratings</del> - <mark>rating</mark> agencies is broadly
disseminated and generally used throughout the industry. We believe that the claims paying ability and financial strength ratings
of our principal insurance subsidiaries are important factors in marketing our products to certain customers. Our debt ratings
impact both the cost and availability of future borrowings and, accordingly, our cost of capital. Each of the rating agencies
reviews ratings periodically and there can be no assurance that current ratings will be maintained in the future. A downgrade of
any of these ratings in the future could make it more difficult to either market our products successfully or raise capital to
support business growth. We maintain significant indebtedness in the ordinary course of business and may incur further
indebtedness in the future. Our indebtedness could adversely affect our financial condition, our ability to react to changes in the
economy or our industry and could divert our cash flow from operations for debt service costs, leaving us with less cash flow
from operations available to fund growth, stock repurchases, dividends and other corporate purposes. The total indebtedness of
The Cigna Group was approximately $ 31-30.1-9 billion as of December 31, 2022-2023. Carrying indebtedness: • requires us
to dedicate a portion of our cash flow from operations to debt payments, thereby reducing the availability of cash flow to fund
our operations and growth strategy, including investments, acquisitions and capital expenditures, make stock repurchases, pay
dividends and for general corporate purposes; • increases our vulnerability to general adverse economic and industry conditions,
which may require us to dedicate an even greater percentage of our cash flow from operations to the payment of principal and
interest on our debt and limit our access to capital markets such that additional capital may not be available or may be available
only on unfavorable terms; • exposes us to increases in interest rates to the extent increased interest expense is not offset by
increased income from our investment assets; and • limits our flexibility in planning for, or reacting to, changes in or challenges
```

relating to our business and industry. The covenants in our debt instruments may have the effect, among other things, of restricting our financial and operating flexibility to respond to significant changes in business and economic conditions. We may incur or assume significantly more debt in the future which may subject us to additional restrictive covenants and increase the risks described above. If our cash flow and capital resources are insufficient to service our debt obligations, we may be forced to seek additional dividends from our subsidiaries, sell assets, seek additional equity or debt capital or restructure our debt. Unfavorable developments in economic conditions may adversely affect our business, results of operations and financial condition, Many factors, including geopolitical issues, future economic downturns, man-made disasters, natural disasters (including those as a result of climate change) and pandemics, availability and cost of credit and other capital and consumer spending can negatively impact the U. S. and global economies. Our results of operations could be materially adversely affected by the impact of unfavorable economic conditions on our clients and customers (both employers and individuals), health care providers, pharmacy manufacturers, pharmacy providers and third- party vendors. For example: • Employers may take action to reduce their operating costs by modifying, delaying or canceling plans to purchase our products or making changes in the mix of products purchased that are unfavorable to us. • Higher unemployment rates, employee attrition (including challenges filling open positions in light of a an increasingly competitive job market) and workforce reductions could result in lower enrollment in our employer- based plans (including an increase in the number of employees who opt out of employer- based plans) or our individual plans. • Because of unfavorable economic conditions or the ACA, employers may stop offering health care coverage to employees or elect to offer this coverage on a voluntary, employee- funded basis as a means to reduce their operating costs. • If clients are not successful in generating sufficient funds or are precluded from securing financing, they may not be able to pay, or may delay payment of, accounts receivable that are owed to us. • Our clients or potential clients may force us to compete more vigorously on factors such as price and service to retain or obtain their business. • Our clients may be acquired, consolidated, or otherwise fail to successfully maintain or grow their business or workforce, which could reduce the number of customers we serve or otherwise result in lower than anticipated utilization of our services. • A prolonged unfavorable economic environment could adversely impact the financial position of hospitals and other health care providers, potentially increasing our medical costs. • Our third- party vendors could significantly and quickly increase their prices or reduce their output to reduce their operating costs. Our business depends on our ability to perform necessary business functions in an efficient and uninterrupted fashion. • Other insurers' financial condition may be weakened, increasing the risk that we will receive significant assessments for obligations of insolvent insurers pursuant to guaranty associations, indemnity funds or other similar laws and regulations. Certain of the foregoing events have occurred and may continue to occur, and the occurrence of these events may, individually or in the aggregate, lead to a decrease in our customer base, revenues or margins or an increase in our operating costs. In addition, during and following a prolonged unfavorable economic environment, federal and state budgets could be materially adversely affected, resulting in reduced or delayed reimbursements or payments in government programs such as Medicare and Social Security or under contracts with government entities. These budgetary pressures also could cause the government to impose new or a higher level of taxes or assessments on us, such as premium taxes on insurance companies and HMOs and surcharges or fees on select fee- for- service and capitated medical claims. Although we could attempt to mitigate or cover our exposure from such increased costs through, among other things, increases in premiums, there can be no assurance that we will be able to mitigate or cover all of such costs, which may have a material adverse effect on our business, results of operations, financial condition and liquidity. We are subject to the credit risk of our reinsurers. We enter into reinsurance arrangements with other insurance companies, primarily in connection with acquisition or divestiture transactions when the underwriting company is not being acquired or sold. Under all reinsurance arrangements, reinsurers assume insured losses, subject to certain limitations or exceptions that may include a loss limit. These arrangements also subject us to various obligations, representations and warranties with the reinsurers. Reinsurance does not relieve us of liability as the originating insurer. We remain liable to the underlying policyholders if a reinsurer defaults on obligations under the reinsurance arrangement. Although we regularly evaluate the financial condition of reinsurers to minimize exposure to significant losses from reinsurer insolvencies, reinsurers may become financially unsound. If a reinsurer fails to meet its obligations under the reinsurance contract or if the liabilities exceed any applicable loss limit, we will be forced to cover the claims on the reinsured policies. The collectability of amounts due from reinsurers is subject to uncertainty arising from a number of factors, including whether the insured losses meet the qualifying conditions of the reinsurance contract, whether reinsurers or their affiliates have the financial capacity and willingness to make payments under the terms of the reinsurance contract and the magnitude and type of collateral supporting our reinsurance recoverable, such as holding sufficient qualifying assets in trusts or letters of credit issued. Although a portion of our reinsurance exposures are secured, the inability to collect a material recovery from a reinsurer could have a material adverse effect on our results of operations, financial condition and liquidity. 47-44