Special Section on Pharmacokinetics and ADME of Biological Therapeutics—Minireview

Absorption, Distribution, Metabolism, and Excretion of US Food and Drug Administration–Approved Antisense Oligonucleotide Drugs

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ABSTRACT

Absorption, distribution, metabolism, and excretion (ADME) are the key biologic processes for determination of a drug’s pharmacokinetic parameters, which have direct impacts on efficacy and adverse drug reactions (ADRs). The chemical structures, dosage forms, and sites and routes of administration are the principal determinants of ADME profiles and consequent impacts on their efficacy and ADRs. Newly developed large molecule biologic antisense oligonucleotide (ASO) drugs have completely unique ADME that is not fully defined. ASO-based drugs are single-stranded synthetic antisense nucleic acids with diverse modes of drug actions from induction of mRNA degradation, exon skipping and restoration, and interactions with proteins. ASO drugs have a great potential to treat certain human diseases that have remained untreatable with small molecule-based drugs. The ADME of ASO drugs contributes to their unique set of ADRs and toxicity. In this review, to better understand their ADME, the 10 US Food and Drug Administration (FDA)-approved ASO drugs were selected: fomivirsen, pegaptanib, mipomersen, nusinersen, inotersen, defibrotide, eteplirsen, golodirsen, viltolarsen, and casimersen. A meta-analysis was conducted on their formulation, dosage, sites of administration, local and systematic distribution, metabolism, degradation, and excretion. Membrane permeabilization through endocytosis and nucleolytic degradation by endonucleases and exonucleases are major ADME features of the ASO drugs that differ from small-molecule drugs. The information summarized here provides comprehensive ADME characteristics of FDA-approved ASO drugs, leading to a better understanding of their therapeutic efficacy and their potential ADRs and toxicity. Numerous knowledge gaps, particularly on cellular uptake and subcellular trafficking and distribution, are identified, and future perspectives and directions are discussed.

SIGNIFICANCE STATEMENT

Through a systematic analysis of the existing information of absorption, distribution, metabolism, and excretion (ADME) parameters for 10 US Food and Drug Administration (FDA)-approved antisense oligonucleotide (ASO) drugs, this review provides an overall view of the unique ADME characteristics of ASO drugs, which are distinct from small chemical drug ADME. This knowledge is useful for discovery and development of new ASO drugs as well as clinical use of current FDA-approved ASO drugs.

Introduction

Absorption, distribution, metabolism, and excretion (ADME) of a drug affect the biologic and physiologic processes that determine the drug concentrations reaching to its organ(s) and tissue(s) over time. ADME parameters influence whether desired therapeutic goals can be reached while undesired adverse drug reactions (ADRs) can be avoided. ADRs can often be reduced by choosing the correct administration dose and route of administration, including newly developed classes of biologic drugs, such as antisense oligonucleotide (ASO) drugs (Amantana and Iversen, 2005).

ASO drugs are single-stranded nucleic acid polymers intended for the treatment of common and rare human diseases (Smith and Zain, 2019). Their principles of drug action involve the modulation of the production or function of disease-causing proteins via alterations of pre-

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ABBREVIATIONS: ADME, absorption, distribution, metabolism, and excretion; ADR, adverse drug reaction; Apo B, apolipoprotein B; ASO, antisense oligonucleotide; AUC, area under the curve; CMV, cytomegalovirus; CNS, central nervous system; CSF, cerebrospinal fluid; DMD, Duchenne muscular dystrophy; EE, early endosome; FDA, US Food and Drug Administration; P450, cytochrome P450; TTR, transthyretin; VEGF, vascular endothelial growth factor.
mRNA splicing, mRNA degradation, regulation of protein translation, or direct interactions with proteins (Di Fusco et al., 2019). Because of the ability to directly alter the production of disease-causing proteins, the ASO drugs have shown significant potential to bring transformative treatments to a variety of human diseases (Bennett et al., 2019), particularly some diseases, such as Duchenne muscular dystrophy (DMD), for which small chemical drugs are not available (Quemener et al., 2020). ADME processes of ASO drugs have direct impacts on therapeutic efficacy and ADRs; therefore, the understanding of general ADME features of ASO drugs and their unique characteristics is vital. Because the ADME characteristics of ASO drugs differ substantially from small chemical drugs, understanding of ADME is critical for the discovery and development of new ASO drugs with greater efficacy and improved safety profiles as well as for the clinical practice of approved ASO drugs. By analyzing all ASO drugs approved by the US Food and Drug Administration (FDA), this article aims to provide an overview of both class-wide and ASO-specific ADME features of the 10 FDA-approved ASO drugs and how these features differ from those of small molecule drugs.

**Brief Historical Perspective.** The discovery and development of ASO drugs have a history of nearly half a century (Oberemok et al., 2018), dating back to 1978 when a specific oligonucleotide was first discovered to be able to inhibit Rous sarcoma viral RNA translation (Stephenson and Zamecnik, 1978). After nearly 20 years of effort by the pharmaceutical industry, the FDA approved the first ASO drug, fomiviren, for the treatment of cytomegalovirus (CMV) retinitis in 1998 (Fig. 1A) (Roehr, 1998). However, fomiviren was withdrawn from the market by the European Union in 2002 and in the US in 2006 due to the success of other highly active antiretroviral therapy and the reduced incidence of CMV infection (Shahryari et al., 2019). After a gap of more than a decade with only one drug (pegaptanib) approved in 2004, the FDA has approved eight ASO drugs since 2013, including mipomersen, nusinersen, inotersen, golodirsen, viltolarsen, and casimersen (Fig. 1A). However, when mipomersen was approved by the FDA for the treatment of familial hypercholesterolemia, a black box warning was placed in the drug label describing the risk of hepatotoxicity (Dixon et al., 2014). Because of this serious safety warning and unacceptable risk, the FDA withdrew mipomersen from the US market in 2019 only after 6 years of use. Recently, further refinement of the ASO drug discovery and development platform has led to a much faster pace in the emergence of new members of this class of drugs used in the treatments of muscular disorders, neurologic disorders, and various other diseases (Bennett et al., 2017; Dhuri et al., 2020) (Fig. 1). More than 100 ASO drugs have been enrolled in various phases of clinical trials, with some close to be approved within a few years (Croke et al., 2021). The annual market sales of ASO drugs have been estimated to be approximately $1 billion in 2021 and are expected to increase to $1.62 billion by 2025 with a steady annual growth rate of approximately 13%.

Based on their main modes of drug action, the 10 FDA-approved ASO drugs can be classified into five groups (Fig. 2). 1) Group 1 drugs induce RNase H-mediated mRNA degradation, including fomiviren targeting on CMV IE2 mRNA, mipomersen on apolipoprotein B (Apo B) mRNA, and inotersen on transthyretin (TTR) mRNA (Fig. 2A). 2) Group 2 drugs induce exon skipping, including eteplirsen for dystrophin exon 51 skipping, golodirsen for dystrophin exon 53 skipping, viltolarsen for dystrophin exon 53 skipping, and casimersen for dystrophin exon 45 skipping (Fig. 2B). 3) Nusinersen in Group 3 works as exon restoration. The target of nusinersen is the pre-mRNA of survival motor neuron 2 (SMN2) (Wurster and Ludolph, 2018). Upon ASO binding, SMN2 pre-mRNA then undergoes an ASO-induced splicing event, resulting in inclusion of exon 7 (Wurster and Ludolph, 2018). With exon 7 included, a fully functional SMN2 mRNA is transcribed and levels of SMN protein are increased (Fig. 2C). 4) Pegaptanib in Group 4 works via an aptamer-protein complex by interacting with vascular endothelial growth factor (VEGF) receptor (Fig. 2D). 5) Defibrotide in Group 5 targets on multiple targets. Defibrotide works on veno-occlusive disease by antagonistic mechanisms. Defibrotide is a polydisperse mixture of various single-stranded oligonucleotide particles. By binding to adenosine receptors A1 and A2, defibrotide reduces systemic levels of plasminogen activator inhibitor 1 and increases concentrations of endogenous prostaglandins (I2 and E2), which modulates thrombomodulin, platelets, and fibrinolysis and thus results in the anti-inflammatory, antiatherosclerotic, anti-ischemic, and antithrombotic property of defibrotide (Baker and Demaris, 2016).

![Timeline of the 10 FDA-approved antisense oligonucleotide (ASO) drugs with FDA approval dates, companies that developed the drugs, and target diseases. FH, familial hypercholesterolemia; PNA, polyneuropathy of amyloidosis; SMA, spinal muscular atrophy; VOD, veno-occlusive disease.](dmd.aspetjournals.org)
Fig. 2. The mode of drug action of the 10 FDA-approved ASO drugs. (A) ASOs with RNase H-mediated mRNA degradation. By binding to their target mRNAs, ASO drugs cause the targeted mRNAs to be broken down into smaller pieces within the cell. ASO drugs in this modality are fomivirsen targeting on CMV IE2 mRNA, mipomersen targeting on Apo B mRNA, and inotersen targeting on transthyretin (TTR) mRNA. (B) ASO-mediated exon skipping. By sterically blocking a splicing site in the exon with mutations, ASO drugs result in skipping of the mutated exon and increased production of protein with enhanced function due to the skipped exon. ASO drugs in this mode include eteplirsen for exon 51 skipping, golodirsen and viltolarsen for exon 53 skipping, and casimersen for exon 45 skipping.
As with any other drug under discovery and development, the information available on the ADME features of these ASO drugs has been part of the vast battery of studies of an investigational drug with the anticipation that some ADME principles would vary for ASO drug molecules from traditional drugs (Weidolf et al., 2021). ADME properties are useful for the generation of preclinical pharmacokinetic predictions of drug efficacy at specific doses and frequency of treatment in clinical studies (Reichel and Lienau, 2016; Wang et al., 2019b).

**Key Recent Advances.** The ADME features of the 10 FDA-approved ASO drugs were investigated to find common class-wide characteristics and unique features of individual ASO drugs. This analysis was based on available information from the FDA ASO drug labeling, the outcomes of clinical trials from ASO investigational new drug (IND) applications, and the published literature.

**Administration.** The FDA-approved ASO drugs can be divided into four groups based on their administration routes: (A) intravitreal injection for fomivirsen and pegaptanib (Fig. 3A), (B) intrathecal injection for nusinersen (Fig. 3B), (C) subcutaneous injection (s.c.) for mipomersen and inotersen (Fig. 3C), and (D) intravenous infusion (i.v.) for defibrotide, eteplirsen, golodirsen, viltolarsen, and casimersen (Fig. 3D). It should be noted that defibrotide may be administered orally for long-term usage, though this formulation is not available yet. Each ASO drug uses a specific route of administration to treat its target disease or condition.

Fomivirsen is used to treat CMV retinitis in the eye and is administered by direct intravitreal injection with a 50 μl solution containing 0.33 mg of the drug (de Smet et al., 1999). Fomivirsen has two phases of administration: loading and maintenance phases. The loading dose consists of two intravitreal injections every two weeks, and the maintenance dose is given only once every 4 weeks. Pegaptanib is used to treat neovascular macular degeneration in the back of the eye; therefore, it is administered by direct intravitreal injection of a 90 μl solution containing 0.3 mg pegaptanib once every 4 weeks.

Nusinersen was designed for the treatment of spinal muscular atrophy (Goodkey et al., 2018). A direct intrathecal injection of a 5 ml solution containing 12 mg drug is used to deliver the drug to the spinal cord. Nusinersen also has two phases of administration: loading and maintenance phases. In the loading phase, the first 3 doses are given at 14-day intervals with the fourth dose given 30 days after the third dose. Maintenance doses are subsequently provided once every 4 months. Special instructions for sedating patients must be followed when performing the intrathecal injections of nusinersen and are guided by ultrasound or other imaging techniques. A removal of 5 ml cerebrospinal fluid over

<table>
<thead>
<tr>
<th>Administration path</th>
<th>ASO Drug</th>
<th>Formulation</th>
<th>Dosing</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>Fomivirsen</td>
<td>0.33 mg in 50 μl</td>
<td>Loading dosing: twice other week; Maintenance dosing: once every 4 weeks</td>
</tr>
<tr>
<td></td>
<td>Pegaptanib</td>
<td>0.30 mg in 90 μl</td>
<td>Dosing: once every 6 weeks</td>
</tr>
<tr>
<td>B</td>
<td>Nusinersen</td>
<td>12 mg in 5 mL</td>
<td>Loading dosing: first 3 doses with 14 days interval, the 4th dose 30 days after the 3rd dose; Maintenance dosing: once every 4 months</td>
</tr>
<tr>
<td></td>
<td>Mipomersen</td>
<td>200 mg in 1 mL</td>
<td>Dosing: once weekly</td>
</tr>
<tr>
<td></td>
<td>Inotersen</td>
<td>284 mg in 1 mL</td>
<td>Dosing: once weekly</td>
</tr>
<tr>
<td>C</td>
<td>Defibrotide</td>
<td>6.25 mg/kg</td>
<td>Dosing: 2-hour infusion every 6 hours for 21 days</td>
</tr>
<tr>
<td></td>
<td>Eteplirsen</td>
<td>30 mg/kg</td>
<td>Dosing: 35-60 min infusion once weekly</td>
</tr>
<tr>
<td></td>
<td>Golodirsen</td>
<td>30 mg/kg</td>
<td>Dosing: 35-60 min infusion once weekly</td>
</tr>
<tr>
<td></td>
<td>Casimersen</td>
<td>30 mg/kg</td>
<td>Dosing: 35-60 min infusion once weekly</td>
</tr>
<tr>
<td></td>
<td>Viltolarsen</td>
<td>80 mg/kg</td>
<td>Dosing: 60 min infusion once weekly</td>
</tr>
<tr>
<td>D</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Fig. 3. Administration paths, formulation, and dosing information of the FDA-approved ASO drugs. The ASO drugs are divided based on their administration paths with (A) intravitreal injection, (B) intrathecal injection, (C) subcutaneous injection, and (D) intravenous infusion. The information for formulation and dosing is extracted from the FDA drug labels at Drugs@FDA: FDA-Approved Drugs (https://www.accessdata.fda.gov/scripts/cder/daf/index.cfm).
1–3 minutes using a spinal anesthesia needle is recommended prior to the nusinersen administration.

Mipomersen is the ASO drug developed for the treatment of familial hypercholesterolemia. It works by targeting Apo B mRNA and stimulating its degradation in the liver. The ensuing reduction in Apo B protein is responsible for the lipid-lowering effect of mipomersen (Parham and Goldberg, 2019). An s.c. injection of 200 mg mipomersen in 1 ml solution is applied once weekly for up to 6 months to achieve the therapeutic goal. Mipomersen should not be administered by intramuscular or intravenous injection according to the FDA drug label (Wong and Goldberg, 2014). Inotersen was developed to treat polynuropathy of amyloidosis caused by over-deposits of TTR protein in tissues, which is mainly produced in the liver (Benson et al., 2018; Keam, 2018). Inotersen is administered by an s.c. injection with a 1 ml solution containing 284 mg drug once weekly. Just like mipomersen, inotersen must be injected subcutaneously.

Defibrotide was approved to treat hepatic veno-occlusive disease, and the mechanism of action of defibrotide involves an enhancement of the enzymatic activity of plasmin to hydrolyze fibrin clots (Richardson et al., 2017). Defibrotide is given as a 2-hour i.v. infusion every 6 hours at a dose of 6.25 mg/kg for a total of 21 days (Richardson et al., 2021).

Eteplirsen, golodirsen, viltolarsen, and casimersen were all developed to treat DMD. They all target the dystrophin gene pre-mRNA, resulting in the skipping of specific mutation-containing exons. Their administration is via i.v. infusion for approximately 30–60 minutes once weekly at a dose of 30 mg/kg for eteplirsen, golodirsen, and casimersen and 80 mg/kg for viltolarsen. In addition, viltolarsen is infused via i.v. using a peripheral or central venous catheter and cannot be infused concomitantly with other medications through the same i.v. access line.

Small chemical drugs typically have a high oral availability compared to large molecule therapeutics despite being metabolized by phase I/II enzymes in the liver, intestine, and kidney. Due to oral dosing, the half-life is normally very short and requires daily dosing, sometimes multiple times daily. The longer half-life of ASO drugs allows less frequent dosing requirements, such as weekly, monthly, and even yearly.

Absorption. Absorption is a key parameter that affects pharmacokinetics and defines the movement of the unmetabolized drug from the site of administration to the local or systematic circulation (Alagga and Gupta, 2021). Different factors can affect the absorption of drugs into the body. These include drug administration routes and first-pass metabolism. Both factors have an impact on drug bioavailability, measured as percentage of the bioactive form of drug out of the total amount of the parent drug administered. Absorption will result in a peak time of plasma drug concentration. The peak plasma time is the time needed to achieve the highest level of drug present in plasma reflective of absorption of the drug into the body. The 10 FDA-approved ASO drugs investigated in this review have various absorption patterns related to the disease state intended to treat, how the drug is administered, and peak plasma concentration time (Table 1).

Both fomivirsen and pegaptanib are intended to treat ocular diseases, which require the drug target and optimization approach to be more local than systematic. Based on animal studies on rabbits, absorption of fomivirsen is highest in the retina and iris (Henry et al., 2001). Fomivirsen was detectable in the retina within hours of injection, and concentrations still increased over 3–5 days. Due to the low doses used for intravitreal administration and slow disposition of fomivirsen from the eye, there is limited systemic drug exposure. Similarly, animal studies indicated that the highest concentration of pegaptanib was found in the eyes of rabbits, especially in the vitreous fluids, retina, and aqueous fluid (Foy et al., 2007). Pegaptanib is slowly absorbed into the systemic circulation after intravitreal administration and, as for many ASO drugs, results in ADRs involving the kidney.

Nusinersen administration methods circumvent the blood-brain barrier through intrathecal injection, which then affects the central nervous system (CNS). Nusinersen is distributed and absorbed in the CNS with the help of cerebrospinal fluid after intrathecal administration. The plasma concentration is relatively low compared with cerebrospinal fluid concentration (Luu et al., 2017). The median terminal half-life of nusinersen in the cerebrospinal fluid was determined to be 163 days, which supported infrequent dosing of once every 4–6 months in patients with spinal muscular atrophy. After administration, the peak plasma nusinersen concentrations can be detected within a few hours (Chiriboga et al., 2016).

The remaining ASO drugs, namely mipomersen, inotersen, defibrotide, eteplirsen, golodirsen, viltolarsen, and casimersen, have primarily systemic absorption and affect a range of organs in the body. After subcutaneous administration, peak plasma concentrations of mipomersen and inotersen were observed at 2–4 hours (Li et al., 2014; Yu et al., 2020). With intravenous infusion, peak plasma concentrations of defibrotide, eteplirsen, golodirsen, viltolarsen, and casimersen can be reached within 1–2 hours. Defibrotide stands alone, reaching its peak plasma concentration at 2 hours. It is important to mention that the effects of the ASO drugs on the kidneys are prominent with most ASO drugs investigated in this review.

Distribution. Distribution refers to the movement of a drug between intravascular (blood/plasma) and extravascular compartments of the

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### TABLE 1

Local and systematic absorption of the FDA-approved ASO drugs

*The data are extracted from the FDA drug labels of each ASO drug.*

<table>
<thead>
<tr>
<th>Drug</th>
<th>AdministrationSite and Route</th>
<th>Local vs. Systematic Absorption</th>
<th>Ability to Cross Blood-Brain Barrier</th>
<th>Tissues with Highest Drug Concentrations</th>
<th>Peak Plasma Concentration Time (Hours)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fomivirsen</td>
<td>Intravitreal injection</td>
<td>Local eye</td>
<td>No</td>
<td>Retina, iris; systemic exposure: minimal</td>
<td>Very low</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>(below limits of quantitation)</td>
<td></td>
</tr>
<tr>
<td>Pegaptanib</td>
<td>Intravitreal injection</td>
<td>Local eye</td>
<td>No</td>
<td>Vitreous fluid, retina, aqueous fluid; systemic exposure: kidney</td>
<td>Slowly absorbed into systemic circulation</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>CNS, plasma, peripheral tissue (skeletal muscle, liver, kidney), fat, bone marrow, spleen</td>
<td>1.7–6</td>
</tr>
<tr>
<td>Nusinersen</td>
<td>Intrathecal injection</td>
<td>CNS and systematic</td>
<td>Yes</td>
<td>Liver, kidney, bone marrow, adipose tissue, lymph nodes</td>
<td>3–4</td>
</tr>
<tr>
<td>Mipomersen</td>
<td>S.C. injection</td>
<td>Systematic</td>
<td>No</td>
<td>Liver, kidney, broad tissues</td>
<td>2–4</td>
</tr>
<tr>
<td>Inotersen</td>
<td>S.C. injection</td>
<td>Systematic</td>
<td>No</td>
<td>Cells lining blood vessel in liver</td>
<td>2</td>
</tr>
<tr>
<td>Defibrotide</td>
<td>I.V. infusion</td>
<td>Systematic</td>
<td>No</td>
<td>Kidney, skeletal muscles</td>
<td>1–2</td>
</tr>
<tr>
<td>Eteplirsen</td>
<td>I.V. infusion</td>
<td>Systematic</td>
<td>No</td>
<td>Kidney, all tissue except CNS</td>
<td>1</td>
</tr>
<tr>
<td>Golodirsen</td>
<td>I.V. infusion</td>
<td>Systematic</td>
<td>No</td>
<td>Kidney, skeletal muscle</td>
<td>1</td>
</tr>
<tr>
<td>Viltolarsen</td>
<td>I.V. infusion</td>
<td>Systematic</td>
<td>No</td>
<td>Kidney, skeletal muscle</td>
<td>1</td>
</tr>
<tr>
<td>Casimersen</td>
<td>I.V. infusion</td>
<td>Systematic</td>
<td>No</td>
<td></td>
<td>1</td>
</tr>
</tbody>
</table>
body. Protein bound and unbound forms of drugs affect their distribution in the body and absorption to tissues and organs (Manzoor and Mahabadi, 2021). Chemical properties of the ASO drugs affect their distribution. This includes factors such as acid-base characteristics, lipophilicity, size, and protein binding. Biodistribution of the 10 ASO drugs is quite different (Table 2). Of these ASO drugs, inotersen, nusinersen in plasma, defibrotide, and mipomersen have high protein-binding potential (>90%), whereas viltolarsen, golodirsen, nusinersen, eteplirsen, and defibrotide exhibit relatively lowered protein-binding potential (<40%). The freeer form pool of the drug in circulation, the more likely the drug will be absorbed into tissues for a drug response. Bioavailability of nusinersen, defibrotide, eteplirsen, golodirsen, viltolarsen, and capsiomersen after administration through intrathecal or intravenous routes is 100% because the drugs are directly administered into the bloodstream, thus bypassing first-pass metabolism in the liver. Bioavailability is also related to the range of distribution (Price and Patel, 2021). A drug with a wider distribution is more likely to leave the plasma and enter extravascular compartments of the body (the tissues). Comparing the availability data on the range of distribution at steady state, the rank ordering of the ASO drugs is golodirsen > eteplirsen > capsiomersen, viltolarsen, inotersen, and nusinersen. The extent of exposure of the drug in the body is demonstrated by the area under the curve (AUC) of a specific drug. Specific AUC data for golodirsen, viltolarsen, and capsiomersen were not found, but intersubject variability of AUC is 34%–44%, 16%–27%, and 16%–34%, respectively. AUC values for defibrotide showed a 50%–60% increase in patients with renal impairment and hemodialysis-dependent end-stage renal disease. Peak concentration was also 35%–37% higher in patients with renal impairment after both single and multdose regimen. This variability is important to be addressed in the monitoring of ADRs and toxicity in patients with increased AUC.

Metabolism. Metabolism is the process in which each drug becomes chemically modified and broken down by the body. All 10 FDA-approved ASO drugs have mechanisms of degradation in the bloodstream and target cells. These mechanisms are quite different from the conventional metabolic pathways that are used for small molecule drugs.

During clinical trials, defibrotide did not undergo biotransformation through oxidative metabolic pathways; instead, the main pathway was found to be nuclease degradation, including both endonuclease and exonuclease activities in the bloodstream and target cells (Geary et al., 2002). Exonucleases are a class of enzymes that are responsible for cleaving the phosphodiester bonds between adjacent nucleotides at the ends of the nucleic acid chains. Endonucleases are similar in function, but nucleolytic cleavage occurs at an internal location in the nucleic acid chain. Nuclease degradation of defibrotide begins once it has been distributed throughout the retina. The resulting metabolites have not been detected in the blood plasma, thus suggesting that there is no significant systemic absorption of the metabolites (Grillone and Lanz, 2001). Pegaptanib also does not undergo cytochrome P450 (P450)-mediated metabolism. Instead, its primary degradation pathway is endonuclease and exonuclease activity in blood (Kourlas and Schiller, 2006). Mipomersen has the same degradation pathway as well. After metabolic degradation, the metabolites of mipomersen are referred to as “chain shortened” because they range between 7 and 14 nucleotides and have no pharmacological function (Bell et al., 2011). Although the parent drug reaches plasma concentrations within hours of administration, the metabolites have not been found in the plasma, either free or bound to plasma proteins (Crooke and Geary, 2013). During the in vitro testing for eteplirsen, hepatic microsomes did not have a significant effect on the metabolism of the drug. It was determined that metabolism of the drug does not occur in the liver (Lim et al., 2017). There was also no accumulation of the drug or its metabolites, and minimal drug-drug interactions with plasma proteins or other coadministered drugs were detected (Syed, 2016). Nusinersen has an exonuclease-dependent metabolic pathway. The exonuclease activity specifically involves hydrolysis reactions as each individual nucleotide is cleaved off from the ends of the ASO molecule. There are no known interactions with the P450 enzymes, and thus the parent drug is not considered a substrate for hepatic oxidative metabolism (Neil and Bisaccia, 2019). Inotersen is metabolized by both endonucleases and exonucleases. The parent drug is first subject to endonuclease degradation, where the phosphodiester bonds toward the middle of the nucleic acid are cleaved. The metabolites of this reaction have been found to be inactive, thus suggesting that it does not have another pharmacological use as a metabolite. Then, these short metabolites are subject to even more nuclease degradation in the form of exonuclease activity. There is little or no interaction with coadministered drugs that are subject to oxidative metabolism because inotersen is not a substrate of the hepatic metabolic pathway (Gertz et al., 2019). Golodirsen, viltolarsen, and capsiomersen are all FDA-approved ASO drugs developed for the treatment of DMD. These drugs are not subject to hepatic metabolism and can be considered metabolically stable. The metabolites were not found in the plasma or urine. In vitro studies showed that these drugs had little to no interactions with...
eteplirsen is 339 ml/h per kg after 12 weeks of drug therapy with 30
mg/kg per week (Baker, 2017). The peak time is 1.1–1.2 hours, and the
half-life is around 3–4 hours. Nusinersen also primarily utilizes the urin-
yary route to excretion. It is excreted by the kidneys as pharmacologi-
ically inactive chain-shortened oligonucleotides (Li, 2020). The half-life
is expected to be 135–177 days in cerebrospinal fluid (CSF) and 63–87
days in plasma (Shorrock et al., 2018). For CSF clearance, it is assumed
that the transport from the CSF to the plasma compartment is the pre-
dominant mechanism (Luu et al., 2017). Inotersen is metabolized and
excreted through the urine. The duration of the 284 mg/1.5 ml dose is 1
week. The total body clearance rate and half-life are 3.18 l/h and 32.3
days. Since one of the target organs of inotersen is the kidney, the urin-
yary route to excretion is an efficient method of drug delivery to its tar-
get. During the first 24 hours, less than 1% of the drug is excreted un-
changed. Golodirsen also uses the kidney for excretion. However,
golodirsen is mainly excreted unchanged by the renal system. Vito-
larsen is mainly excreted in the urine unchanged as well. The elimina-
tion half-life is approximately 2.5 hours (%), and the plasma clearance
is 217 ml/h per kg (22%). Patients taking 1.25, 5, or 20 mg/kg vol-
larsen weekly for 12 weeks had a total clearance of between 149 ± 21
and 239 ± 97 ml/h per kg (Komaki et al., 2020). Lastly, casimersen also
primarily uses the urinary route for excretion. Its excretion is completed
by 336 hours postdose. The excretion method for casimersen is similar
to that of eteplirsen and golodirsen. Collectively, the commonality in
excretion for all ASO drugs is the predominant use of the urinary sys-
tem. Even fomivirsen, which is administered intravitreally, has trace
amounts detected in urine. This suggests a prominent preference for
excretion via the urinary route for the ASO drugs.

Differences in ADME between Small Molecule Drugs and
ASO Drugs. Based on the above analysis on the information from the
FDA drug label, clinical studies, and published literature, ADME features
of the FDA-approved ASO drugs are quite different from small chemical
drugs with their unique characteristics for ADME. The ADME behavior
of small chemical drugs is mediated by uptake and efflux transporters,
phase I and II drug-metabolizing enzymes, transcription factors, and
xenobiotic-activated nuclear receptors (Fig. 4A). The ADME of ASO
drugs is determined by completely different pathways (Fig. 4B). Absorp-
tion of ASOs into target cells involves uptake via endocytic internaliza-
tion into early endosomes (EEs). ASOs are then moved within the cell by
subcellular trafficking. Release of ASOs from endosomes has significant
impact on therapeutic efficacy and the occurrence of ADRs. Metabolism
of ASO drugs is mainly catalyzed by exonucleases and endonucleases,
resulting in various sizes of shortened oligonucleotides. Either intact or
shortened ASOs are excreted by membrane leakage, vesicle release, or
exosome release to the urinary tract. These features determine the half-
life of ASO drugs in the circulatory system and frequency of administra-
tion. To optimize therapeutic efficacy, studies on the ADME features for
ASO drugs should continue in real-world settings to monitor therapeutic
efficacy and ADRs.

Current Challenges, Knowledge Gaps, and Future Perspec-
tive. Several knowledge gaps for the ADME features of the FDA-
approved ASO drugs were identified, including cellular uptake, subcell-
ular trafficking, and cellular efflux. Almost no information is available
on how these FDA-approved drugs are taken up into cells after local or
systematic distribution. Existing data on uptake mechanisms of ASO
molecules are mainly based on in vitro models of cultured cells. Com-
monly accepted theories are that ASOs undergo cellular uptake by two
competing pathways: the productive pathway bringing the drugs into
their targets and the nonproductive pathway directing the drugs into sat-
urable sinks (Khorkova and Wahlestedt, 2017). Both pathways take up
ASOs by clathrin-coated endocytic vesicles from plasma membrane.
Few factors have been strongly identified as key players for these path-
ways, such as TCP1 complex, Hsp90, SSB, NPM1, and AP2M1 in the

By analyzing the metabolic pathways, there are some commonalities
that can be identified between all FDA-approved ASO drugs. The primary
method of metabolism is nuclease degradation via endonuclease and exo-
nuclease activities. For the majority of these drugs, endonuclease activity
occurs first, thus cleaving the oligonucleotides into fragments. Then exo-
nuclease activity can take place on the 3′ and 5′ ends via hydrolysis reac-
tions. To increase stability of ASO drugs from nuclease degradation, they
have been modified with chemical groups that allow them to remain sta-
ble enough to reach their target tissues. In general, ASO drugs can have
phosphorothioate linkages in the phosphodiester backbone, 2′-O-methox-
yethyl ribose gapmer modifications, and 2′-4′ methylene-bridged nucleic
acids, which are also referred to as locked nucleic acids (Swayze et al.,
2007). These modifications allow the molecules to have better resistance
to nuclease degradation. The FDA-approved ASO drugs summarized
above also have such modifications that likely contribute to their stability
and longer residency at their target tissues. Fomivirsen is considered a
first-generation oligonucleotide, as it has phosphorothioate linkages in the
backbone. This means that one of the oxygen atoms in the phosphate
group at the phosphodiester bond is replaced by a sulfur atom (de Smet
et al., 1999). Mipomersen and inotersen, on the other hand, are considered
second-generation ASOs because of the 2′-O-methoxethyl ribose modi-
fication (Crooke and Geary, 2013). Second-generation ASO drugs have
been found to have better resistance against nuclease degradation; hence,
these drugs have longer half-lives as well and increased stability within
the target cell (Swayze et al., 2007). Another similarity between these
drugs is that they are not subject to metabolism by the liver or hepatic
microsomes. This is due to the drugs not being substrates for the major
class of drug-metabolizing enzymes, P450s. There is also little to no
potential for most of these drugs of having interactions with other proteins
or drug transporters because of their distinct metabolic pathways. The
metabolites for these drugs have not been found to be pharmacologically
active or to impact cellular processes in a manner that could be of safety
concern.

Although this information about the metabolism of ASO drugs pro-
vides insight into the primary pathways of drug degradation, there are
some areas that deserve more research. There is no detailed information
on the chemical and physical properties of ASO drug metabolites that
are the products of nuclease degradation. This topic could be studied
more to better understand the role of metabolites. The specific rates of
the hydrolysis reactions for exonuclease activity could be useful in
improving the stability of the oligonucleotides via chemical modifica-
tions. A possible future direction would be to investigate how the stabil-
ity of ASO drugs can be increased and how it would affect resistance to
nuclease degradation and therapeutic efficacy. The frequency and differ-
ent combinations of chemical modifications can be studied to investi-
gate whether the dosing intervals could be changed as well.

Excretion. Each individual ASO drug has its excretion features.
While conducting preclinical studies of fomivirsen on rabbits, a small
amount of administered fomivirsen was shown to be eliminated
through urine (16%) and feces (3%). Fomivirsen is administered intravi-
treally, and the half-life of fomivirsen is approximately 55 hours in
humans. In monkeys, the half-life is estimated to be 78 hours (Perry
and Balfour, 1999). This is different from the route of excretion for
mipomersen, which is primarily through renal filtration (Visser et al.,
2012) and urine by “nuclease digested metabolites” (Crooke and Geary,
2013). This route of excretion seems to be constant among species.
Eteplirsen is also excreted via the renal system. The renal system
accounts for two-thirds of the excretion of the administered dose within
24 hours post-intravenous administration. The total body clearance for
eteplirsen is 339 ml/h per kg after 12 weeks of drug therapy with 30

productive pathway and paraspeckle proteins, Ku70, and Ku80 in the nonproductive pathway (Li, 2020). However, very limited information is available for identifying key determinant factors in the intracellular endosomal packing of ASO drugs and what proportion of these drugs go into the productive versus nonproductive pathway. These are important considerations since these variables dictate in part the amount of an ASO drug that reaches its therapeutic targets. Furthermore, some surface receptors may be critical for this endocytosis process. For example, adenosine receptors in endothelial cells may have a potential role in the endocytic uptake of defibrotide (Palomo et al., 2016). Future studies

Fig. 4. Comparison of absorption, distribution, metabolism, and excretion (ADME) of small chemical drugs and ASO drugs. (A) ADME of an orally administrated small chemical drug. After absorption, a small chemical drug is normally distributed to metabolic organs of intestine, liver, and kidney via the bloodstream. In the metabolic organs, such as the liver, the drug is taken up by hepatocytes through either passive or active transport against concentration gradients with the help of uptake transporters. Normally, the lipophilic drug is biotransformed by either Phase I or II metabolizing enzymes to more polar water-soluble metabolites in rough endoplasmic reticulum (ER) membrane. The parental drug and metabolites are excreted from hepatocyte cells via efflux transporters into the bloodstream and then become available as an active form to target organs. Therapeutic concentrations of the drug in its target organs are determined by uptake and efflux transporters and Phase I and II metabolizing enzymes, which are tightly regulated at transcriptional and translation levels by transcription factors and nuclear receptors. (B) ADME of an ASO drug. After an intravenous infusion, subcutaneous injection, or direct injection, ASOs travel through the bloodstream, where they can be broken down via endogenous exo- and endonucleases. Once ASOs reach their target organ/cell, such as the liver, they enter cells through various endocytic pathways, including clathrin-mediated, caveolin-mediated, and micropinocytosis endocytosis, into early endosomes (EEs). After being internalized into EEs, ASO drugs go through subcellular trafficking to the late endosomes. A proportion of ASO drugs (maybe very small proportion) is released from late endosomes to cytoplasm to target mRNAs or pre-mRNAs in either the cytoplasm or the nucleus, where they execute their therapeutic actions. Many small cellular proteins including COPII can assist subcellular trafficking to the nucleus. However, the mechanism of nuclear trafficking is not fully understood. Another proportion of ASO drugs undergo into lysosomes or further are released out of the cell via three proposed mechanisms: membrane leakage, back-fusion mediated release, and vesicle-mediated release. Back-fusion mediated release is believed to help endogenous exo- and endonucleases break the ASO down further. ASO drugs may also leave the cell completely via the exosome release from the cell.
should employ FDA-approved ASO drugs as experimental tools to understand cellular uptake mechanisms for this class of drugs, which is completely different from the transporter-mediated uptake of small chemical drugs. Significant variations on the cellular uptake of ASO drugs may exist in the patient population, resulting in differences on therapeutic efficacy and safety.

Another knowledge gap is the lack of information on subcellular trafficking and distribution of the FDA-approved ASO drugs and their impact on therapeutic efficacy. Unlike small chemical drugs, which can quickly reach different subcellular regions after uptake by transporters or non–transporter-dependent diffusion across the cell membrane, ASO drugs are normally first encapsulated in endosomes after uptake into cells by endocytosis. Different ASO drugs have their targets located at different subcellular locations. For RNase H-mediated mRNA degradation, the ASO drugs need to reach either the cytoplasm (ribosomes) or nucleus (Liang et al., 2017). For exon skipping, the ASO drugs should be present in spliceosomes in the nucleus (Echevarria et al., 2018). Endosomal release and intracellular trafficking are key determinants of the pharmacological actions of the ASO drugs (Juliano, 2018). Numerous factors have been identified as mediators of these events, which can enhance endosomal release and intracellular trafficking (Ono et al., 2020). For example, Hsc70 (Liang et al., 2021), M6PR (Liang et al., 2020), and lipid conjugates (Wang et al., 2019a) are proposed contributors of endosomal release and ASO trafficking. Future studies should pay particular attention to how ASO drugs are taken up and released from endosomes and what assists them in reaching their therapeutic targets. Equally important is the identification of mechanisms that can lead to the optimization of intracellular trafficking of ASO drugs toward pharmacological targets to enhance therapeutic outcomes. It is also imperative to know which intracellular locations may be associated with ASO-mediated cell damage and toxicity to avoid ASO drugs from reaching and accumulating in these intracellular locations, which are distinct from their therapeutic targets.

Regarding ASO drugs biotransformation, the mechanism by which the remaining parent and shortened ASO drugs are effluxed out from cells after breakdown by endonucleases and exonucleases is completely unknown. It is also unknown whether efflux is a critical component. Unlike many small chemical drugs that are eliminated from the cells by efflux transporters, ASOs use completely different cellular exit mechanisms, which may have considerable impact on plasma concentrations and duration of ASO drugs in their target cells. It can be hypothesized that lack of efflux activity could have substantial effects on accumulation of ASO drugs in liver and kidney. Hepatotoxicity and kidney toxicity are a major safety concern for the FDA-approved ASO drugs (Alhamadani et al., 2022). Determination of efflux mechanisms may provide better understanding of interindividual variability of ASO-mediated drug toxicity.

The conclusions and development of ASO drugs have moved at a much faster pace in recent years. Numerous new ASO drugs are expected to be approved in the next decade, and the use of ASO drugs for a variety of indication will become more commonplace. However, before this relatively new class of drugs is more widely accepted, a better understanding of the ADME characteristic of these drugs is needed to optimize their therapeutic efficacy and safety profiles. The current review summarizes administration, absorption, distribution, metabolism (degradation), and excretion of the 10 FDA-approved ASO drugs. Specifically, this review explores the unique set of ADME features for ASO drugs that differs quite dramatically from the more traditional ADME profiles of small chemical drugs. Several knowledge gaps are identified on cellular uptake by endocytosis, intracellular endosomal release, intracellular/nuclear trafficking, and cellular efflux for these ASO drugs, and future directions are discussed.

Authorship Contributions

Participated in research design: Bahal, Rasmussen, Manautou, Zhong. Performed data analysis: Migliorati, S. Liu, A. Liu, Gogate, Nair. Wrote or contributed to the writing of the manuscript: Migliorati, S. Liu, A. Liu, Gogate, Nair, Bahal, Rasmussen, Manautou, Zhong.

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