earth
enterprise

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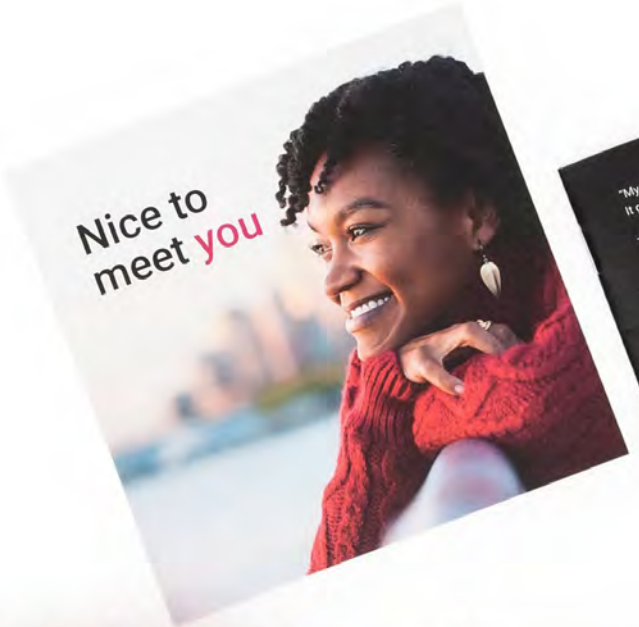
the human energy company™

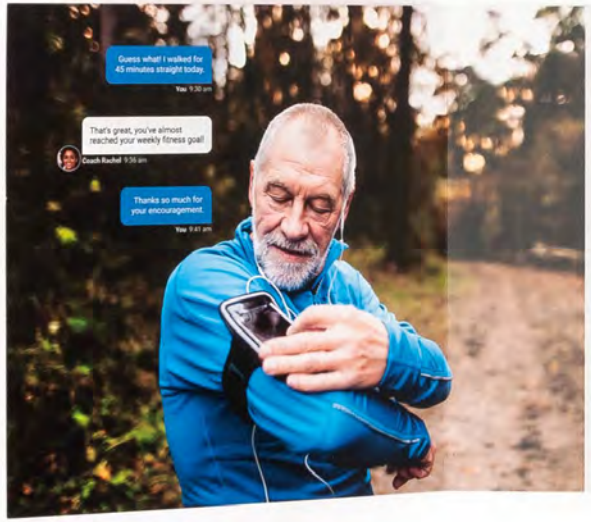
quick guide inside

the human energy company™

How to check your blood sugar

1. Wash your hands with soap and water and dry thoroughly with a clean towel.
2. Insert a test strip into the strip port.
3. Wait for "apply sample to test strip" animation to appear on meter screen.
4. Lance your fingertip, then gently massage finger from base to tip until blood drop appears.
5. Apply blood drop to the tip of inserted test strip.
6. Listen for beep. Remove your finger from test strip and wait for reading.





Turn-by-turn navigation for life

Whether you've been living with diabetes for a day or decades, One Drop adapts to you wherever you are on your journey. Our proven approach to getting healthy uses the power of data to help you make lasting changes.

An ally at every step

Connect with a personal coach who can answer your questions and offer encouragement when you need it most. Coaches use your real-time data to give you personalized feedback specific to your situation.

Customized transformation plan

Using proven techniques from behavioral science, your interactive educational experience helps you change your habits and stick with them. Master the topics that matter most for your health.

Know what to do next

Glucose predictions alert you to a sharp drop or unexpected rise in blood sugar levels ahead of time, available for people with type 2 diabetes. Predictions are paired with real-time advice, so you can prevent problems before they happen.

Set reminders, reach goals, and more

Remember to take medications, eat healthy, exercise, and more with personalized reminders. Set health goals and track your progress toward reaching them.

FLUORESCENT SENSOR
TRANSMITTER
MOBILE APPS
109
HERE'S WHAT THE EVERSENSE BRIDGE PROGRAM DOES
EVERSENSE® BRIDGE — A NEW PATIENT ACCESS PROGRAM IS HERE!
NOW YOU CAN TRY THE EVERSENSE® 90-DAY CGM SYSTEM FOR JUST
eversense
Continuous Glucose Monitoring System
90 DAYS TO BE UNSTOPPABLE
LIFT HERE
Little sensor. Big life.
NOT ACTUAL PRODUCTS

Helpful Information for People With Newly Diagnosed Advanced Non-Small Cell Lung Cancer and Their Families

Avastin, in combination with carboplatin and paclitaxel (chemotherapy), has been proven effective in clinical trials and is approved by the FDA for first-line treatment of patients newly diagnosed with a common form of NSCLC (unresectable, locally advanced, recurrent, or metastatic non-squamous, non-small cell lung cancer).

About Avastin

What is Avastin?

Avastin (pronounced ah-va'-stin) is an innovative targeted therapy that is approved by the FDA for use in combination with carboplatin and paclitaxel (chemotherapy) to help fight NSCLC. If you have been diagnosed with unresectable, locally advanced, recurrent, or metastatic non-squamous, NSCLC, then Avastin is an option for you.

Avastin is not chemotherapy, so the side effects and how it treats cancer are different. *Avastin is known as a targeted therapy, because it attacks a protein that is produced by many cells and overproduced by cancer cells.* In the next section, you will learn how Avastin is thought to work.

KEEP THIS VALUABLE RESOURCE ON HAND

Refer back to this brochure as you gather important information, support, and more. Also, take notes so that you can remember to discuss any questions you may have with your doctor or health care team.

In clinical trials, some people receiving Avastin in combination with carboplatin and paclitaxel (chemotherapy) experienced side effects, including some that were serious. To learn more about the possible side effects of Avastin, please see pages 14 to 20 and 32 to 33, and enclosed full Product Information, including **Boxed WARNINGS**.

AVASTIN® bevacizumab

Take An Active Role

Even though you have NSCLC, you still have a life to live. Taking an active role and managing your treatment goals and plans means doing everything you can to feel better, so you can focus on what matters most to you. Here's how you can start:

Please see pages 14 to 20 and 32 to 33, and enclosed full Product Information, including **Boxed WARNINGS**, for additional safety information.

1. **Understand your treatment.** Learn how each of your anticancer drugs work and what to expect from each one. Understand how targeted therapy, like Avastin, is different from chemotherapy, and know the effects of each therapy on your body. Keep a record of your medicine, any possible side effects, and your dosing schedule. Feel comfortable asking your doctor and health care team questions to help you understand how your medications work. See the "Questions to ask your doctor" section on the following page to get started.

2. **Talk to your doctor about your treatment goals.** You and your doctor know your medical history. You may also have discussed what you want to accomplish and gain, including daily goals, social support, and other important aspects of your life. Together, you should discuss treatment goals; this way, you'll know what to expect and be able to stay focused on what matters most to you.

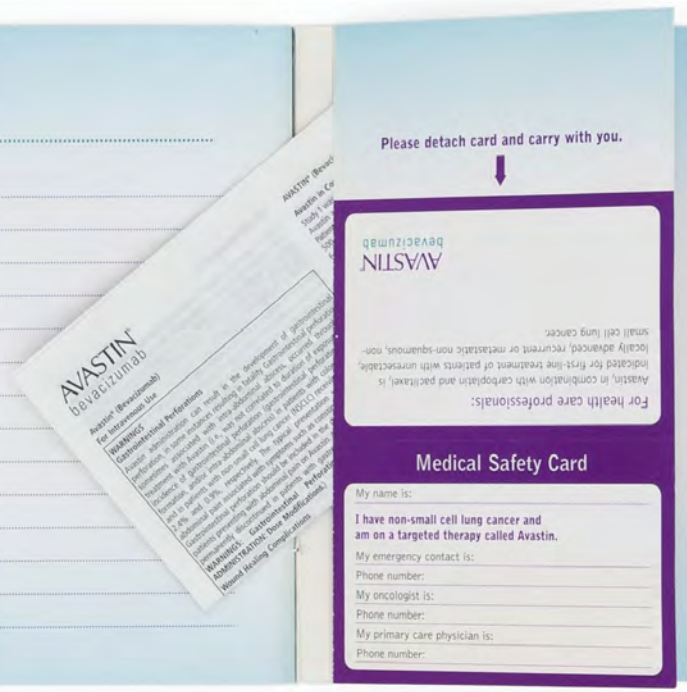
3. **Keep track of how you're feeling, and let your health care team know about it.** Get a journal to keep track of changes in your health. Use it to express how you feel physically, and write down questions you may have for your health care team. Include any side effects you notice and report them to your doctor or nurse. Knowing how you're doing while on treatment can help your health care team provide the best care for you.

4. **Stay involved in all decision making.** Always make sure you get the answers you need from your doctor and health care team so that you can feel comfortable with your treatment plan. Understanding the reasons for the treatment your doctor suggests may help you feel more in control of your situation. Play an active role in helping your health care team make the best decision for your individualized care.

Always discuss with your doctor and health care team any questions you have about cancer therapy and treatment with Avastin.

AVASTIN®
bevacizumab

Notes

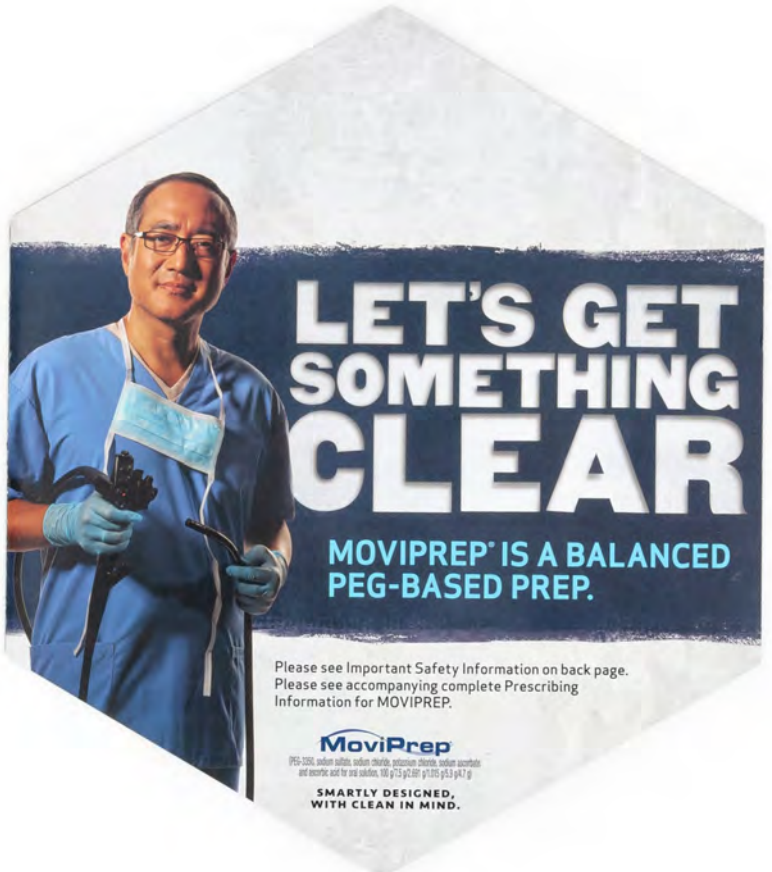


Medical Safety Card

Make sure any doctor or health care professional you visit knows that you're receiving Avastin. Simply fill out the information on the card below and carry this card with you at all times in your wallet, purse, or where it is most likely to be found when you're being treated by health care professionals other than your oncologist. Show it to any health care professional who may treat you so that he/she can provide the most appropriate care. This may include your dentist or an emergency room physician.

Please see pages 14 to 20 and 32 to 33, and enclosed full Product Information, including **Boxed WARNINGS**, for additional safety information.

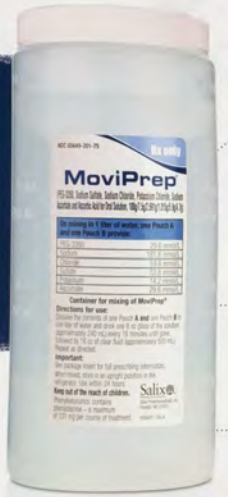
AVASTIN®
bevacizumab


LET'S GET SOMETHING CLEAR
MOVIPREP® IS A BALANCED PEG-BASED PREP.
Please see Important Safety Information on back page.
Please see accompanying complete Prescribing Information for MOVIPREP.
MoviPrep
SMARTLY DESIGNED, WITH CLEAN IN MIND.

NOT EVERY PATIENT IS ALIKE.
That's exactly why MoviPrep® is uniquely designed with the safety of PEG and the built-in hydration of osmotic/electrolyte solution.[1]

MoviPrep®
(PEG-3350, sodium sulfate, sodium chloride, potassium chloride, sodium ascorbate, and ascorbic acid for oral solution. 100 g/7.5 g/2.691 g/1.015 g/5.9 g/4.7 g)

SMARTLY DESIGNED, WITH CLEAN IN MIND.



Please see Important Safety Information on back page. Please see accompanying complete Prescribing Information for MOVIPREP.

HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use MOVIPREP® safely and effectively. See full prescribing information for MOVIPREP.

aspartat
potassiu
(3)
• Each po
acid. US

THE SAFETY OF PEG

- PEG + ELS solutions are well trusted and have been extensively used by physicians for nearly 30 years[2]
- Over 8 million MoviPrep prescriptions written in the US since 2006[3]
- No differences in safety and tolerability between geriatric and younger adult patients[1]

89% EXCELLENT OR GOOD CLEANSING OF THE COLON when split-dosed[1]

THE BENEFIT OF BUILT-IN HYDRATION

- Built-in osmotic and electrolyte balance to help keep patients hydrated during the prep process[1]

CAN **OTHER PREPS** SAY THAT?

INDICATION
MOVIPREP® (PEG-3350, sodium sulfate, sodium chloride, potassium chloride, sodium ascorbate and ascorbic acid for oral solution) is an osmotic laxative indicated for cleansing of the colon as a preparation for colonoscopy in adults 18 years of age or older.

CONTRAINDICATION
MOVIPREP® (PEG-3350, sodium sulfate, sodium chloride, potassium chloride, sodium ascorbate and ascorbic acid for oral solution) is contraindicated in patients with gastrointestinal (GI) obstruction, bowel perforation, gastric retention, ileus, toxic colitis or toxic megacolon, and patients who have had a hypersensitivity reaction to any of its components.

DISEASE STATE OVERVIEW

Pathogenesis, Diagnosis, and Treatment of Pulmonary Arterial Hypertension (PAH)

LUNG BIOTECHNOLOGY

What is PAH?

HEALTHY HEART AND LUNGS

The heart and lungs work together to make sure that every part of your body gets the oxygen it needs. Blood that is pumped from the right side of your heart goes to your lungs. Just like the air you inhale moves through passageways, the blood that flows through your lungs moves through a set of branching pipes called pulmonary vessels. If there is trouble with the pulmonary vessels, you may have difficulty getting oxygen into the blood.

DISEASE STATE OVERVIEW

LUNG
BIOTECHNOLOGY
2016

Pathogenesis, Diagnosis, and Treatment of Pulmonary Arterial Hypertension (PAH)

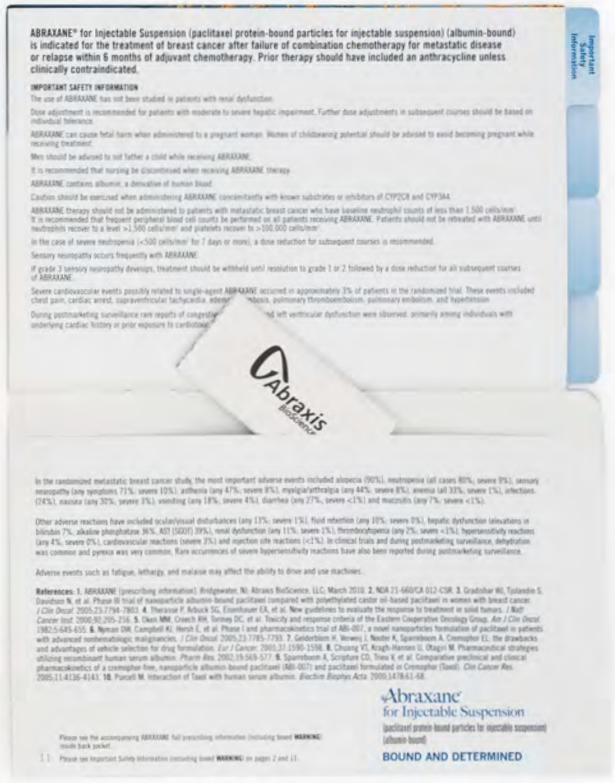
1 What is PAH?
2 Pathophysiology
3 Clinical Classification
4 Signs & Symptoms
5 Diagnostic Approach
6 Treatment Approach

PAH HEART AND LUNGS
PAH is high blood pressure in the arteries (blood vessels) of the lungs, which is different than ordinary high blood pressure that can occur throughout the body. Whereas ordinary high blood pressure often causes no apparent symptoms, PAH makes it more difficult for your body to get the oxygen it needs, which can cause symptoms such as shortness of breath, feeling tired, and more. Over time, PAH affects the condition of the right side of the heart, which decreases the ability of the heart and lungs to work properly.
Enlarged right ventricle
Enlarged right ventricle
Thickened walls
Healthy Heart
PAH Affected Heart
Progression of the disease
Increased PVR
Myocardial hypertrophy
Progressive contractile impairment
RV dilation to preserve stroke volume
RV failure
2 Pathophysiology
3 Clinical Classification
4 Signs & Symptoms
5 Diagnostic Approach
6 Treatment Approach

TYVASO
(treprostinil)
(I inhale) because I have things to focus on
Adding Tyvaso, an inhaled treatment for pulmonary arterial hypertension (PAH), to background oral therapy may help you improve exercise ability.
INDICATION
Tyvaso is a prescription medicine used in adults to treat pulmonary arterial hypertension (PAH) (WHO Group 1), which is high blood pressure in the arteries of your lungs. Tyvaso can improve exercise ability in people who also take bosentan (an endothelin receptor antagonist, (ERA)) or sildenafil (a phosphodiesterase-5 (PDE-5) inhibitor). Your ability to do exercise decreases 4 hours after taking Tyvaso.
The effects of Tyvaso are unknown in patients under 18 years of age.
Please look inside for Indication and complete Important Safety Information.
Please visit www.tyvaso.com to see the Full Prescribing Information.
Powered by videoplusprint.com

Deliver the albumin-bound advantage
nab® technology delivers paclitaxel to tumor cells by exploiting the favorable tumor-targeting properties of albumin[6,7]
ABRAXANE® particle
albumin-bound paclitaxel
ABRAXANE uses proprietary nab® technology
Paclitaxel is bound to albumin particles[1]
Albumin facilitates the delivery of paclitaxel, which is poorly water soluble, to tumor cells[6,7]
The nab® technology of ABRAXANE exploits the high albumin-uptake property of tumor cells to deliver paclitaxel without the use of solvents[1,6,7]
ABRAXANE for Injectable Suspension (paclitaxel protein-bound particles for injectable suspension) (albumin-bound) is indicated for the treatment of breast cancer after failure of combination chemotherapy for metastatic disease or relapse within 6 months of adjuvant chemotherapy. Prior therapy should have included an anthracycline unless clinically contraindicated.
Please see the accompanying ABRAXANE full prescribing information (including boxed WARNING) inside back pocket.
Please see Important Safety Information (including boxed WARNING) on pages 2 and 11.
Abraxane® for Injectable Suspension (paclitaxel protein-bound particles for injectable suspension) (albumin-bound)
BOUND AND DETERMINED
Convenience
Safety
ABC of Support®

ABRAXANE® delivers efficacy to make more possible
nab® delivers. nab® technology exploits the natural properties of albumin, facilitating the delivery of water-insoluble compounds to the tumor.
In a pivotal phase III clinical trial in metastatic breast cancer, ABRAXANE delivered nearly double the objective response rate vs solvent-based paclitaxel*
ABRAXANE for Injectable Suspension (paclitaxel protein-bound particles for injectable suspension) (albumin-bound) is indicated for the treatment of breast cancer after failure of combination chemotherapy for metastatic disease or relapse within 6 months of adjuvant chemotherapy. Prior therapy should have included an anthracycline unless clinically contraindicated.
Please see the accompanying ABRAXANE full prescribing information (including boxed WARNING) inside back pocket.
Please see Important Safety Information (including boxed WARNING) on pages 2 and 11.
*Please see next page for full details.
Abraxane® for Injectable Suspension (paclitaxel protein-bound particles for injectable suspension) (albumin-bound)
BOUND AND DETERMINED
nab®
Linear PK
Higher Dose
Efficacy

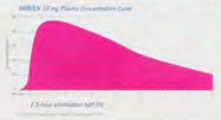

ABRAXANE® for Injectable Suspension (paclitaxel protein-bound particles for injectable suspension) (albumin-bound) is indicated for the treatment of breast cancer after failure of combination chemotherapy for metastatic disease or relapse within 6 months of adjuvant chemotherapy. Prior therapy should have included an anthracycline unless clinically contraindicated.
IMPORTANT SAFETY INFORMATION
The use of ABRAXANE has not been studied in patients with renal dysfunction.
Dose adjustment is recommended for patients with moderate to severe hepatic impairment. Further dose adjustments in subsequent courses should be based on individual tolerance.
ABRAXANE can cause fetal harm when administered to a pregnant woman. Women of childbearing potential should be advised to avoid becoming pregnant while receiving treatment.
Men should be advised to not father a child while receiving ABRAXANE.
It is recommended that nursing be discontinued when receiving ABRAXANE therapy.
ABRAXANE contains albumin, a derivative of human blood.
Caution should be exercised when administering ABRAXANE concomitantly with known substrates or inhibitors of CYP2C8 and CYP3A4.
ABRAXANE therapy should not be administered to patients with metastatic breast cancer who have baseline neutrophil counts of less than 1,500 cells/mm³.
It is recommended that frequent peripheral blood cell counts be performed on all patients receiving ABRAXANE. Patients should not be retreated with ABRAXANE until neutrophils recover to a level >1,500 cells/mm³ and platelets recover to >100,000 cells/mm³.
In the case of severe neutropenia (<500 cells/mm³ for 7 days or more), a dose reduction for subsequent courses is recommended.
Sensory neuropathy occurs frequently with ABRAXANE.
If grade 3 sensory neuropathy develops, treatment should be withheld until resolution to grade 1 or 2 followed by a dose reduction for all subsequent courses of ABRAXANE.
Severe cardiovascular events possibly related to single-agent ABRAXANE occurred in approximately 3% of patients in the randomized trial. These events included chest pain, cardiac arrest, supraventricular tachycardia, edema, dyspnea, hypotension, pulmonary thromboembolism, pulmonary embolism, and hypertension.
During postmarketing surveillance rare reports of congestive heart failure and left ventricular dysfunction were observed, primarily among individuals with underlying cardiac history or prior exposure to cardiotoxic
Abraxis BioScience
In the randomized metastatic breast cancer study, the most important adverse events included alopecia (90%), neutropenia (all cases 80%; severe 9%), sensory neuropathy (any symptoms 71%; severe 10%), asthenia (any 47%; severe 8%), myalgia/arthralgia (any 44%; severe 8%), anemia (all 33%; severe 1%), infections (24%), nausea (any 30%; severe 3%), vomiting (any 18%; severe 4%), diarrhea (any 27%; severe <1%) and mucositis (any 7%; severe <1%).
Other adverse reactions have included ocular/visual disturbances (any 13%; severe 1%), fluid retention (any 10%; severe 0%), hepatic dysfunction (elevations in bilirubin 7%, alkaline phosphatase 36%, AST (SGOT) 39%), renal dysfunction (any 11%; severe 1%), thrombocytopenia (any 2%; severe <1%), hypersensitivity reactions (any 4%; severe 0%), cardiovascular reactions (severe 3%) and injection site reactions (<1%). In clinical trials and during postmarketing surveillance, dehydration was common and pyrexia was very common. Rare occurrences of severe hypersensitivity reactions have also been reported during postmarketing surveillance.
Adverse events such as fatigue, lethargy, and malaise may affect the ability to drive and use machines.
References: 1. ABRAXANE (prescribing information). Bridgewater, NJ: Abraxis BioScience, LLC; March 2010. 2. NDA 21-660/CA 012-CSR. 3. Gradishar WJ, Tjulandin S, Davidson N, et al. Phase III trial of nanoparticle albumin-bound paclitaxel compared with polyethylated castor oil-based paclitaxel in women with breast cancer. J Clin Oncol. 2005;23:7794-7803. 4. Therasse P, Arbuck SG, Eisenhauer EA, et al. New guidelines to evaluate the response to treatment in solid tumors. J Natl Cancer Inst. 2000;92:205-216. 5. Oken MM, Creech RH, Tormey DC, et al. Toxicity and response criteria of the Eastern Cooperative Oncology Group. Am J Clin Oncol. 1982;5:649-655. 6. Nyman DW, Campbell KJ, Hersh E, et al. Phase I and pharmacokinetics trial of ABI-007, a novel nanoparticles formulation of paclitaxel in patients with advanced nonhematologic malignancies. J Clin Oncol. 2005;23:7785-7793. 7. Gelderblom H, Verweij J, Nooter K, Sparreboom A. Cremophor EL: the drawbacks and advantages of vehicle selection for drug formulation. Eur J Cancer. 2001;37:1590-1598. 8. Chuang VT, Kragh-Hansen U, Otagiri M. Pharmaceutical strategies utilizing recombinant human serum albumin. Pharm Res. 2002;19:569-577. 9. Sparreboom A, Scripture CD, Trieu V, et al. Comparative preclinical and clinical pharmacokinetics of a cremophor-free, nanoparticle albumin-bound paclitaxel (ABI-007) and paclitaxel formulated in Cremophor (Taxol). Clin Cancer Res. 2005;11:4136-4143. 10. Purcell M. Interaction of Taxol with human serum albumin. Biochim Biophys Acta. 2000;1478:61-68.
Abraxane® for Injectable Suspension (paclitaxel protein-bound particles for injectable suspension) (albumin-bound)
BOUND AND DETERMINED
Please see the accompanying ABRAXANE full prescribing information (including boxed WARNING) inside back pocket.
Please see Important Safety Information (including boxed WARNING) on pages 2 and 11.
Important Safety Information

Breakthrough technology to break through opioid addiction
INDICATION
Prescription
14 BUNAVAIL.COM
Appropriate for a range of patients
Bunavail buccal film
DELIVERING CHANGE
PATIENT TYPES
Patients in various treatment situations may benefit from BUNAVAIL™
Starting
Those initiating maintenance treatment for the first time
Restarting
Those who have previously been treated for opioid dependence
Converting
Those who face obstacles with their current medication
USE IN SPECIAL POPULATIONS
Please see Important Safety Information on pages 21-22 and enclosed full Prescribing Information to use BUNAVAIL safely and effectively.

A good night's sleep and

refreshed awakenings...

The AMBIEN 2.5-hour half-life helps balance restful nights with refreshed awakenings



- Long enough to provide restful nights
- Short enough to provide refreshed awakenings

AMBIEN treats all 3 symptoms of insomnia[2,3]

- Trouble falling asleep
- Trouble staying asleep
- Waking too early

...on course toward better days

Patients wake up refreshed the next day so they're ready to perform

- Minimal drug effect on next-day functioning[1]

Low abuse potential at recommended doses[1,4]

AMBIEN is the #1 prescribed sleep agent in the US[5]

- More physicians prescribe AMBIEN than any other sleep agent[6]

November

S	M	T	W	T	F	S
		1	2	3	4	5
6	7	8	9	10	11	12
13	14	15	16	17	18	19
20	21	22	23	24	25	26
27	28	29	30			

December

S	M	T	W	T	F	S
				1	2	3
4	5	6	7	8	9	10
11	12	13	14	15	16	17
18	19	20	21	22	23	24
25	26	27	28	29	30	31

National Sleep Awareness Week is a registered trademark of the National Sleep Foundation.

Please see important safety information on reverse side and provided full prescribing information.

AMBIEN®
ZOLPIDEM TARTRATE

Restful nights, refreshed awakenings.

Ambien® CR

zolpidem tartrate

DESCRIPTION

CLINICAL PHARMACOLOGY

* Next-day residual effects were evaluated in 7 studies involving normal volunteers. In 3 studies in adults (including 1 study in a phase-advance model of transient insomnia) and 1 study in elderly subjects, a small but statistically significant decrease in performance was observed in the Digit Symbol Substitution Test (DSST) when compared with placebo. Studies in nonelderly patients with insomnia did not detect evidence of next-day residual effects using the DSST, the Multiple Sleep Latency Test (MSLT), and patients' ratings of alertness.[1]

AMBIEN is indicated for the short-term treatment of insomnia. In elderly or debilitated patients, or patients with hepatic dysfunction, treatment should be initiated with a 5-mg dose and patients closely monitored. Due to its rapid onset of action, patients should take AMBIEN right before going to bed and when ready for sleep. Patients should not take AMBIEN unless they are prepared to get a full night's sleep (7 to 8 hours) to avoid residual effects. Until they know how it will affect them physical or mental performance upon awakening, patients should not drive or operate hazardous machinery after taking AMBIEN or any other sleep medication. During short-term treatment with AMBIEN, the most commonly observed adverse effects in controlled clinical trials were drowsiness (2%), dizziness (1%), and diarrhea (1%). Because individuals with a history of addiction or substance abuse are at increased risk of habituation and dependence, they should be under careful surveillance when receiving AMBIEN or any other hypnotic. AMBIEN is classified as a Schedule IV controlled substance. Sedative-hypnotics have produced withdrawal signs and symptoms following abrupt discontinuation. Hypnotics should generally be limited to 7 to 10 days of use, and reevaluation of the patient is recommended if they are taken for more than 2 to 3 weeks. Prescriptions for AMBIEN should not exceed a 1 month supply.

References: 1. Data on file, sanofi-aventis. 2. Sanofi-Aventis. Ambien CR package insert. Bridgewater, NJ: sanofi-aventis U.S. LLC; 2009. 3. Walsh JK, Krystal AD, Amato DA, et al. Nocturnal sleep duration and daytime functioning in insomnia patients treated for 6 months with eszopiclone 3 mg, zolpidem 10 mg or placebo: evidence for dose-specific effects. Sleep Med. 2007;8(6):608-621. 4. Hajak G, Rodenbeck A, Adler L, et al. Nocturnal and next-day profile of subjective and objective measures of sleepiness, hypnotic efficacy and adverse events after acute administration of zolpidem 10 mg and 20 mg in healthy subjects. Arzneimittelforschung. 1994;44(11):1246-1255. 5. IMS Health. National Sales Perspectives™. Data for the 52 weeks ending December 26, 2008. 6. IMS Health. National Prescription Audit Plus™. Data for the 52 weeks ending December 26, 2008.

Please see provided full prescribing information.

Visit our Web site at www.ambien.com

AMBIEN®
ZOLPIDEM TARTRATE

Restful nights, refreshed awakenings.

sanofi aventis

EVERSENSE REDUCES THE DAY-TO-DAY BURDEN OF DIABETES, HELPING PATIENTS MANAGE THEIR GLUCOSE LEVELS MORE EASILY AND MEET THEIR THERAPY GOALS.

- → **90-DAY SENSOR** provides freedom from frequent insertions
- → **4 SENSOR CHANGES/YEAR** vs. 26-52 for other CGM systems
- → **REDUCED # OF UNRELIABLE** day-1 readings
- → **MINIMIZED MAINTENANCE** and management of supplies
- → **ZERO IRRITATION** with gentle, silicone-based adhesives[4]

ONLY **4** X **PER YEAR**

versus

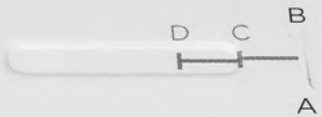
26-52 SENSOR CHANGES FOR **OTHER CGMs**

Left: The Eversense transmitter comes on and off easily, while the sensor stays put.

"Right now I've done about 50 insertions of Eversense. **PATIENTS LOVE IT. THEY COME BACK AND THEY SAY 'THIS IS THE BEST THING I'VE EVER SEEN.'** They never want to leave home without it."

Dr. Bruce Bode, *Eversense CGM certified*

CEMENT TECHNIQUE
SIMPLE STEPS[1]



[~5mm incision]

Step 1: Inject lidocaine; make ~5mm incision in upper arm (between A–B)

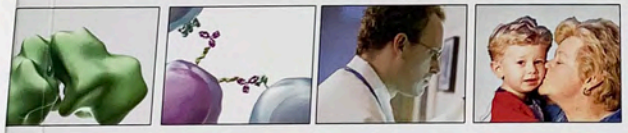
Step 2: Create pocket with custom incision tool

Step 3: Insert sensor with custom insertion tool (along C-D

eversense
Continuous Glucose Monitoring System
PULMONARY ARTERIAL HYPERTENSION
1 WHAT IS PULMONARY ARTERIAL HYPERTENSION
DIAGNOSED?
TREATED?
INTRODUCING THE EVERSENSE BRIDGE PATIENT ACCESS PROGRAM
EVERSENSE® BRIDGE —A NEW PATIENT ACCESS PROGRAM IS HERE!
Now you can try the Eversense® 90-day CGM system for just $99.*
EVERSENSE BRIDGE PROGRAM
PRIOR AUTHORIZATION & APPEALS SERVICES*
Aetna is now covering eversense
Continuous Glucose Monitoring System
FOR MORE INFORMATION, VISIT:
www.eversensediabetes.com
Today, Tomorrow and Beyond
EVERSENSE DELIVERS UNMATCHED WEAR TIME AND IMPROVED GLYCEMIC OUTCOMES.
Eversense Procedure Reference Guide
109
Sensor Insertion
Sensor Removal
Dial 1-844-SENSE4U for support before, during or after performing the procedure.

The Pipeline of Novartis Oncology

open



NOVARTIS
ONCOLOGY

Our vision

Through consistently delivering innovative therapies, which significantly reduce the burden of cancer, we will improve and extend the lives of cancer patients as we make therapies broadly available.

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All compounds are either investigational or studied in new indications. Efficacy and safety have not been established. There is no guarantee that they will become commercially available.

We'll bring our team of
STRATEGISTS and AFFILIATE MEMBERS
directly to you.
Work through real solutions.
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For dealers in
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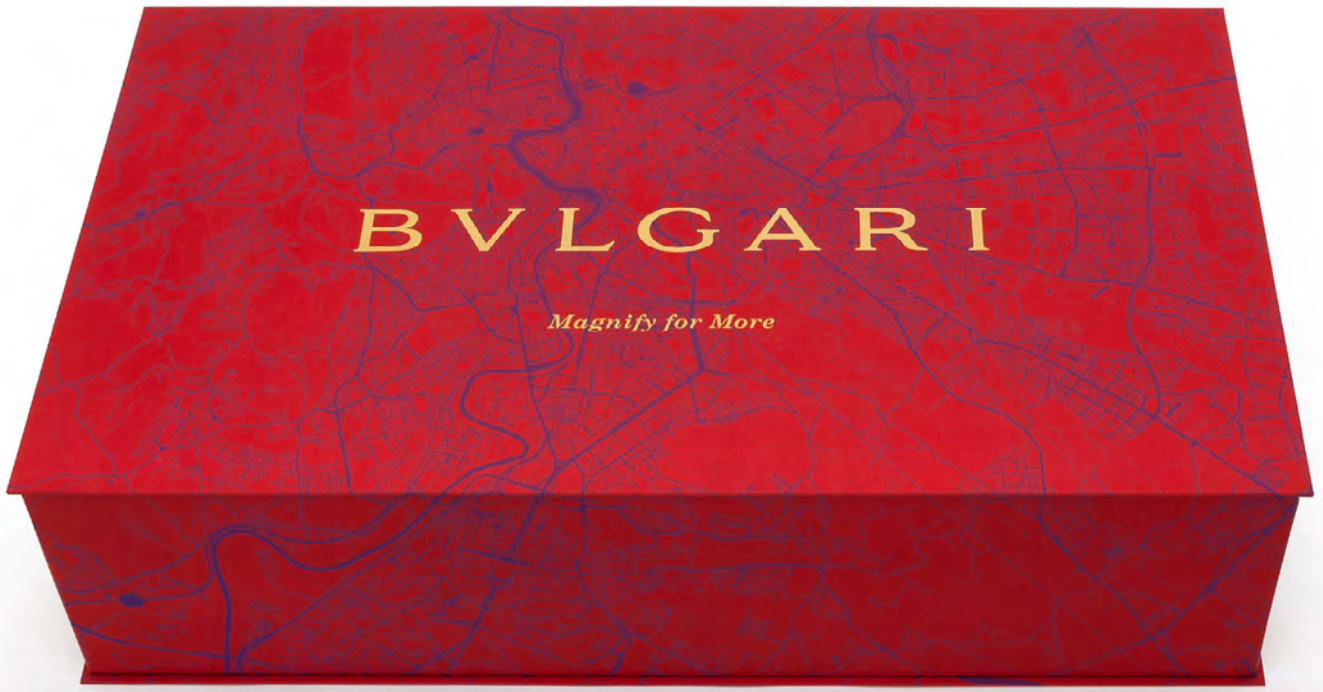
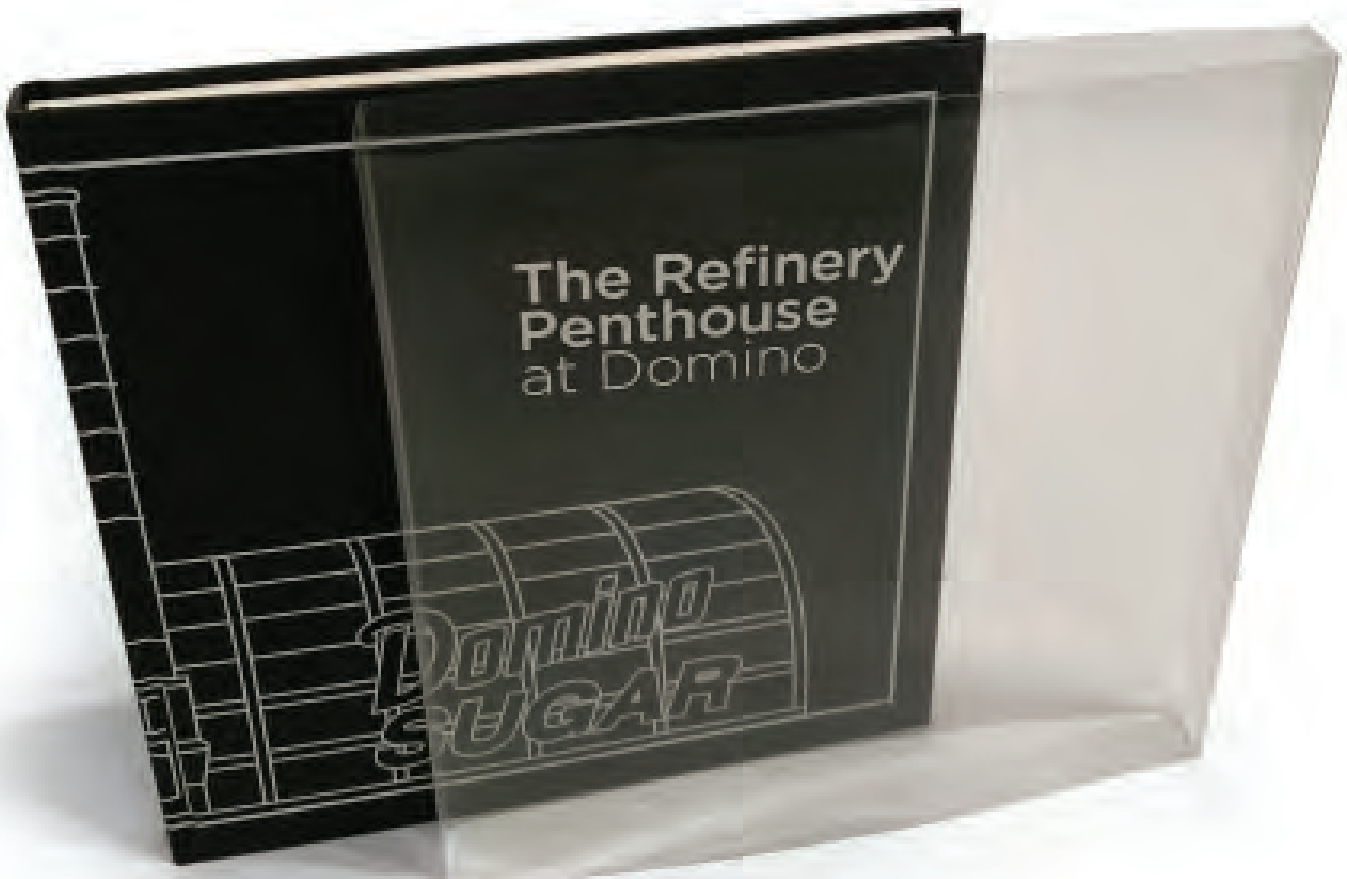
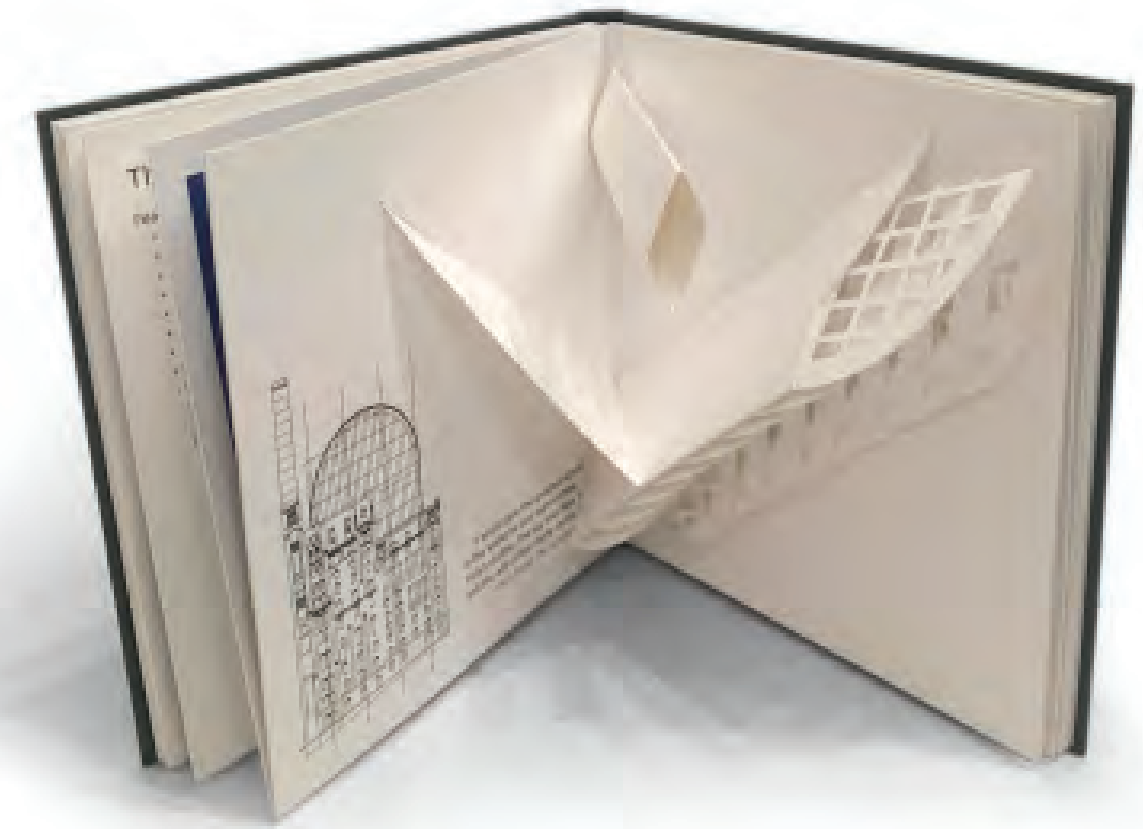

BVLGARI
Magnify for More

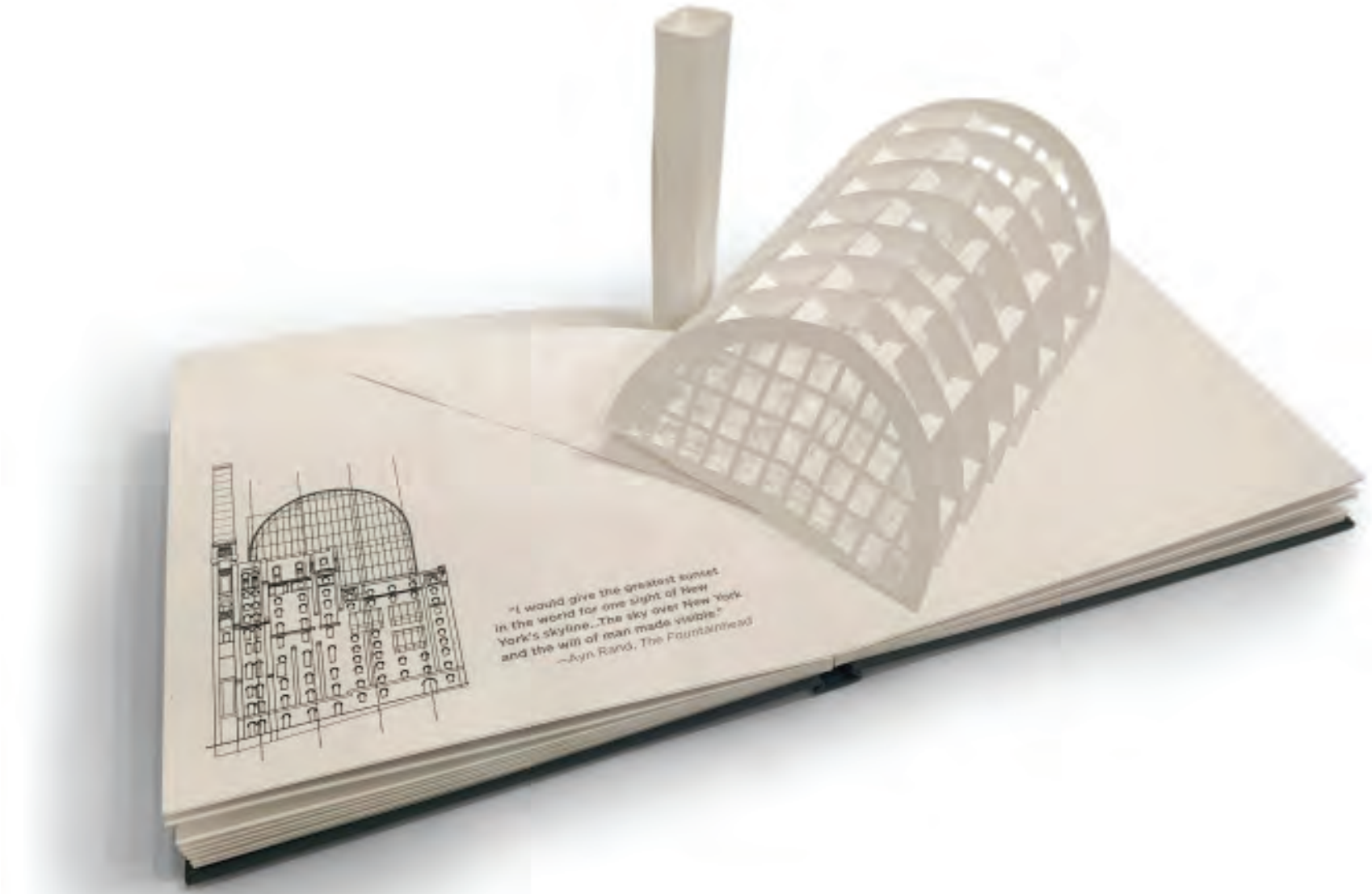
VLGARI
FIORI D'AMORE
DOLCE ESTASI
ROCK'N'ROME

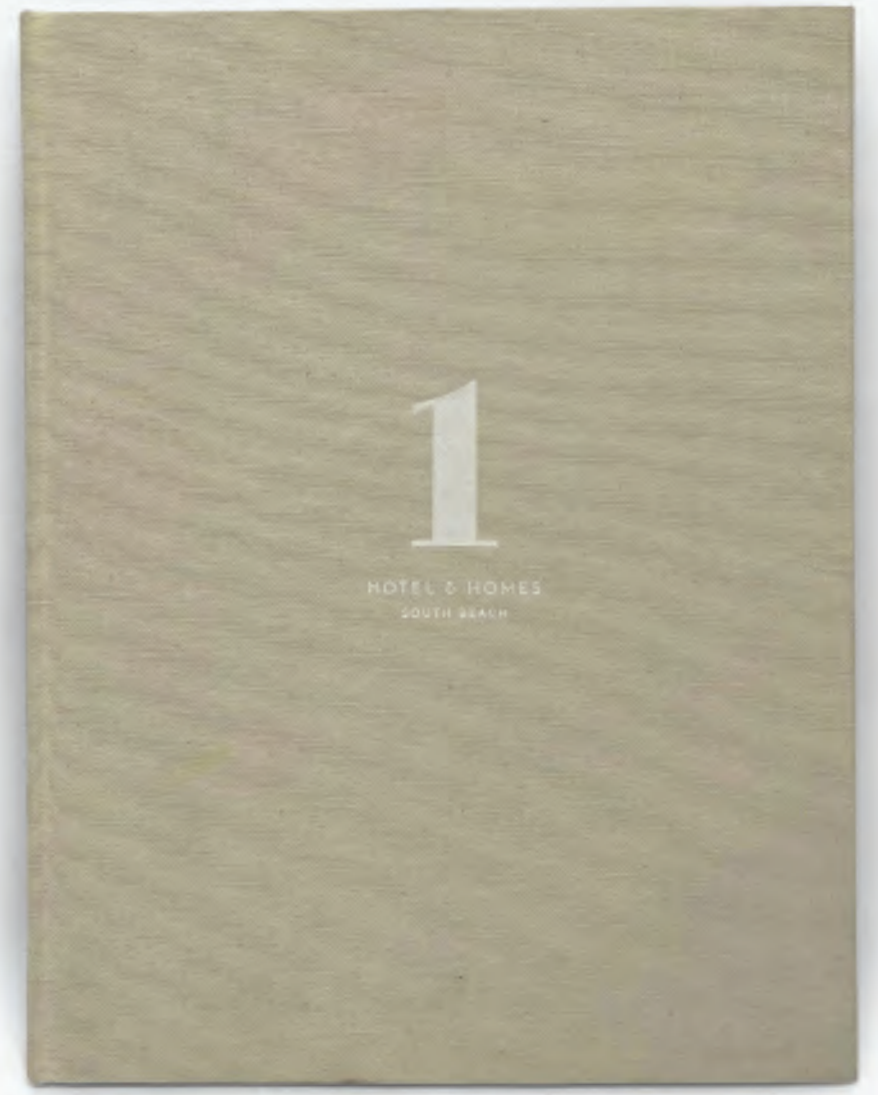
BVLGARI
We hope you enjoy these authentic gifts sourced directly from the Italian cities that have inspired each Allegra fragrance. Accompanying your virtual experience, you will also find a lifetime subscription to Rosetta Stone to brush up on your Italian and transport you to the bustling streets of Florence or Sicily. The rich history of Italian music plays an integral part of its culture and daily lifestyle, from acapella street performers in Rome to the famous Opera at St. Mark's Basilica in Venice. We encourage you to explore these musings through curated playlists to enhance and inspire each of the emotions explored in these fragrances.
As an Italian brand, we are thankful for the time you took to immerse yourself in our culture and to explore the Allegra Franchise virtual experience from the comfort of your own home. Grazie!
DUE FETTE DI TORRONE
RIVA SOLARE
Authentic limoncello in hand-painted ceramic from the Amalfi Coast
FIORI D'AMORE
Rossi 1931 Stationery from Florence
DOLCE ESTASI
Due Fette Di Torrone from Sicily
ROCK'N'ROME
Aperol Spritz Lemon Candles from Rome
FANTASIA VENETA
Primo Aperitivo Negroni from Venice

BVLGARI
Bvlgari Parfums invites you to further immerse yourself in Italian culture with a complimentary lifetime subscription to Rosetta Stone. Enjoy exploring your Italian literary skills at your leisure, preparing you for any future Italian pursuits.
Reference the leaflet for details on how to redeem


The Refinery
Penthouse
at Domino
Domino
SUGAR




"I would give the greatest sunset
in the world for one sight of New
York's skyline...The sky over New York
and the will of man made visible."
—Ayn Rand, The Fountainhead


1
HOTEL & HOMES
SOUTH BEACH

MANHATTAN
HOUSE

MANHATTAN
HOUSE

RALPH
WALKER
TRIBECA
AT
100
BARCLAY

ALWAYS MODERN

With a nod to revered Finnish architect Alvar Aalto, Circa's elegant curve and organic modernist lines mimic its location on Frederick Douglass Circle. Featuring a tiered profile of terraced residences, New York-based architects FXFOWLE have created a striking 11-story presence that is designed to achieve LEED certification. A beautiful and efficient louver system on the façade frames views and shades southern-facing units. Sensitivity to environmental issues and a preoccupation with the new building's role in the larger community speak to the value of this meaningful new landmark.

THE
"ARCHITECTURE IS THE ART AND SCIENCE OF
CREATING THE FRAMEWORK OF OUR LIVES."
BJARKE INGELS

The indulgent spa areas truly appeal to the senses – sight, sound and touch. The water-themed spa features a 52-foot heated pool and indoor sun deck, a cedar-lined dry sauna, and the unique Aqua Spa, a lavish space with tiered steps of heated marble reminiscent of a Turkish hammam.
INVITINGLY LAVISH.
INCREDIBLY INDULGENT.

RALPH WALKER
100 BARCLAY
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WELCOME
TO THE LIVE YOUR BEST LIFE WEEKEND!
This box contains some of the items you'll need this weekend. There is a checklist in your Weekend Guide to ensure that you know what to bring with you each day. There are also a few gifts just for you!
The most important thing to remember: you must have your weekend pass on both Saturday and Sunday in order to gain entry to all the events.

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DIP
INTO some
GOOD FUN
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PERISHABLE
REFRIGERATE IMMEDIATELY
UP
UP

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TIPS
Pizza Hut

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REBEL T7i
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