UNITED STATES SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

SCHEDULE 14A

PROXY STATEMENT PURSUANT TO SECTION 14(a) OF THE SECURITIES EXCHANGE ACT OF 1934

Filed by the Registrant \square

Filed by a Party other than the Registrant o

Check the appropriate box:

o Preliminary Proxy Statement

o Confidential, for Use of the Commission Only (as permitted by Rule 14a-6(e)(2))

o Definitive Proxy Statement

o Definitive Additional Materials

☑ Soliciting Material Pursuant to § 240.14a-12

BIOGEN IDEC INC.

(Name of Registrant as Specified In Its Charter)

N.A.

(Name of Person(s) Filing Proxy Statement, if other than the Registrant)

Payment of Filing Fee (Check the appropriate box):

☑ No fee required.

o Fee computed on table below per Exchange Act Rules 14a-6(i)(1) and 0-11.

(1) Title of each class of securities to which transaction applies:

(2) Aggregate number of securities to which transaction applies:

(3) Per unit price or other underlying value of transaction computed pursuant to Exchange Act Rule 0-11 (set forth the amount on which the filing fee is calculated and state how it was determined):

(4) Proposed maximum aggregate value of transaction:

(5) Total fee paid:

o Fee paid previously with preliminary materials.

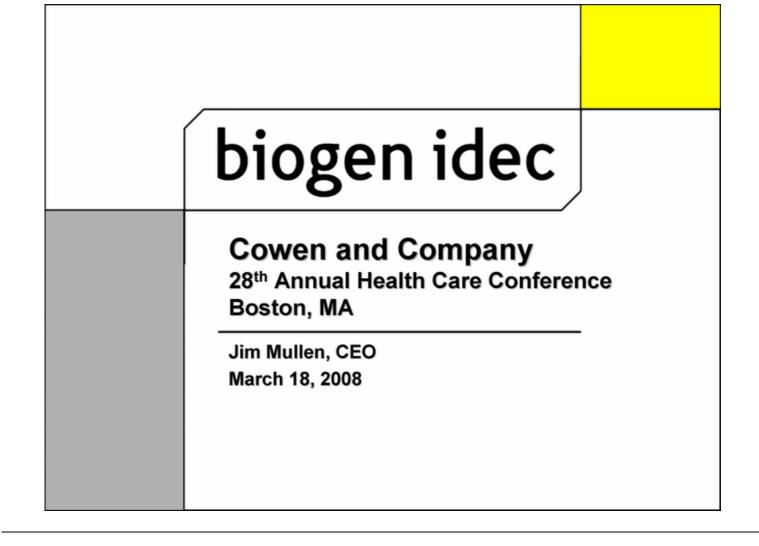
o Check box if any part of the fee is offset as provided by Exchange Act Rule 0-11(a)(2) and identify the filing for which the offsetting fee was paid previously. Identify the previous filing by registration statement number, or the Form or Schedule and the date of its filing.

(1) Amount Previously Paid:

(2) Form, Schedule or Registration Statement No.:

(3) Filing Party:

(4) Date Filed:



Forward Looking & Proxy Solicitation Statements

This presentation includes forward-looking statements about:

- our expected revenues, earnings, and cash flows
- the size and growth of the markets for our products,
- estimates of sales for our products,
 our expected filings with regulatory
- our expected filings with regulatory agencies,
 the anticipated development and timing of programs in our clinical pipeline
- our external business development initiatives
- the sales potential of TYSABRI®

Each forward-looking statement is subject to risks and uncertainties that could cause actual results to differ materially from those that we express or imply.

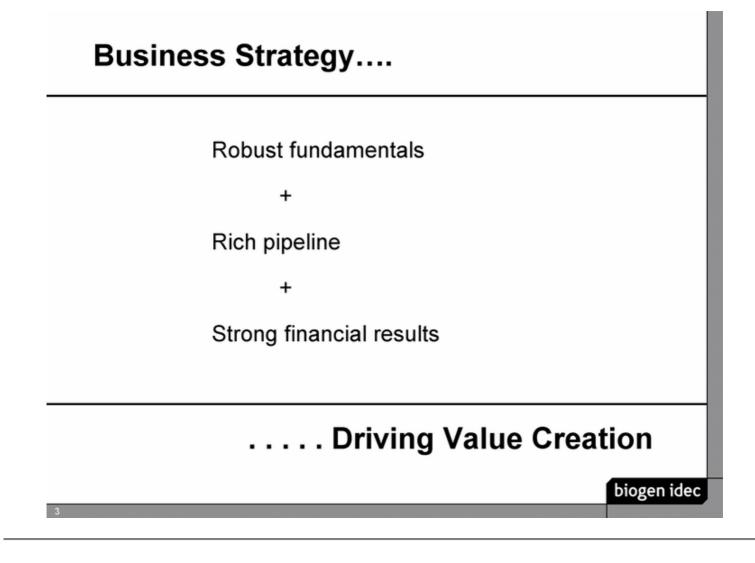
Important factors that could cause our actual results to differ include our continued dependence on our two principal products, AVONEX® and RITUXAN®, the uncertainty of success in commercializing other products including TYSABRI®, the occurrence of adverse safety events with our products, the consequences of the nomination of directors for election to our Board by an activist shareholder, the failure to execute our growth strategy successfully or to compete effectively in our markets, our dependence on collaborations over which we may not always have full control, possible adverse impact of government regulation and changes in the availability of reimbursement for our products, problems with our manufacturing processes and our reliance on third parties, fluctuations in our operating results, our ability to protect our intellectual property rights and the cost of doing so, the risks of doing business internationally and the other risks and uncertainties that are described in Item 1.A. Risk Factors in our annual report on Form 10-K, in our quarterly reports on Form 10-Q and in other periodic and current reports we file with the SEC.

These forward-looking statements speak only as of the date of this presentation, and we do not undertake any obligation to publicly update any forward-looking statements, whether as a result of new information, future events, or otherwise.

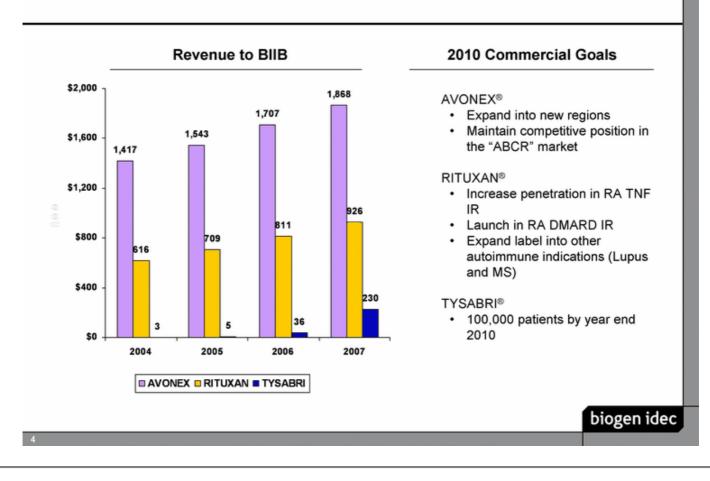
Biogen Idec and its directors, executive officers and other members of its management and employees may be deemed to be participants in the solicitation of proxies from the stockholders of Biogen Idec in connection with the Company's 2008 annual meeting of stockholders. Information concerning the interests of participants in the solicitation of proxies will be included in any proxy statement filed by Biogen Idec in connection with the Company's 2008 annual meeting of stockholders. In addition, Biogen Idec files annual, quarterly and special reports with the Securities and Exchange Commission (the "SEC"). The proxy statement filed in connection with the Company's 2008 annual meeting of stockholders. Biogen Idec files annual, quarterly and special reports with the Securities and Exchange Commission (the "SEC"). The proxy statement filed in connection with the Company's 2008 annual meeting of stockholders when it becomes available before making any voting or investment decision. The Company's proxy statement will also be available for free by writing to Biogen Idec Inc., 14 Cambridge Center, Cambridge, MA 02142. In addition, copies of the proxy materials may be requested from our proxy solicitor, Innisfree M&A Incorporated, by toll-free telephone at (877) 750-5886 or by e-mail at info@innisfreema.com.

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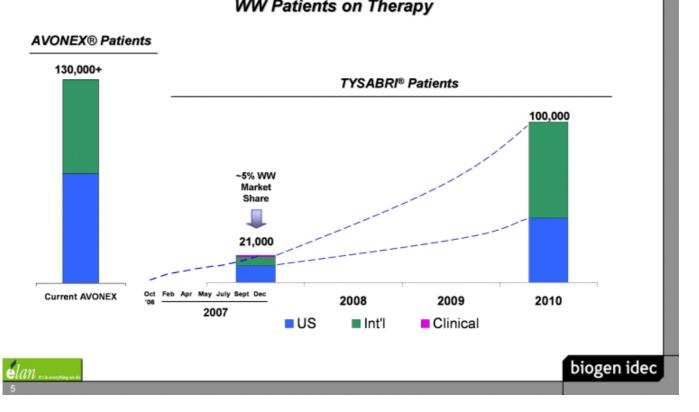
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Robust Fundamentals



100,000 TYSABRI® Patients by 2010



WW Patients on Therapy

Robust Pipeline Near Term Data Readouts

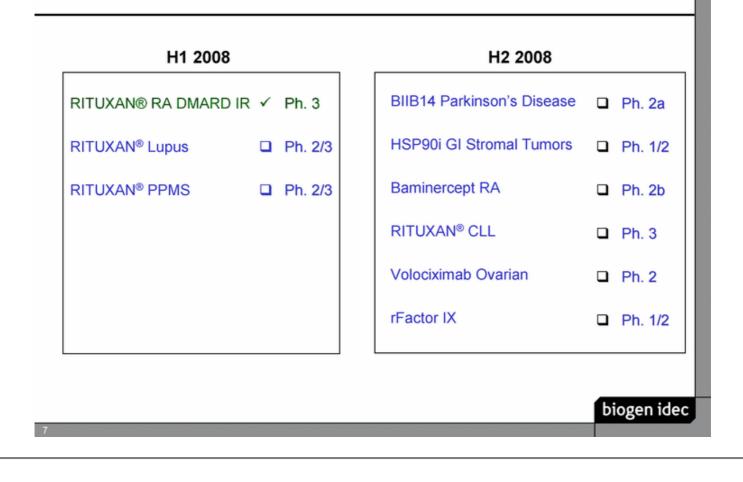
- Over the past 3 years, significantly strengthened pipeline
 - Internal programs
 - External programs
 - >10 molecules accessed via BD strategy
- 4 products in registrational trials... with a 5th by year end
- >4x increase in number of patients in clinical trials in 2008
- 15 programs in Ph. 2 or beyond

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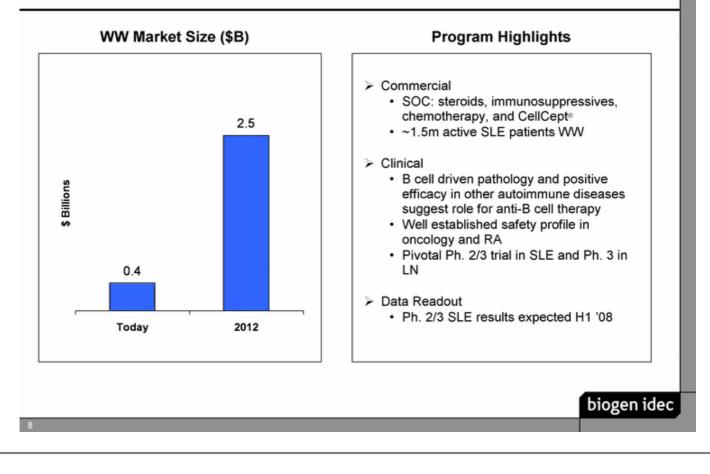
• 8 important data readouts throughout 2008

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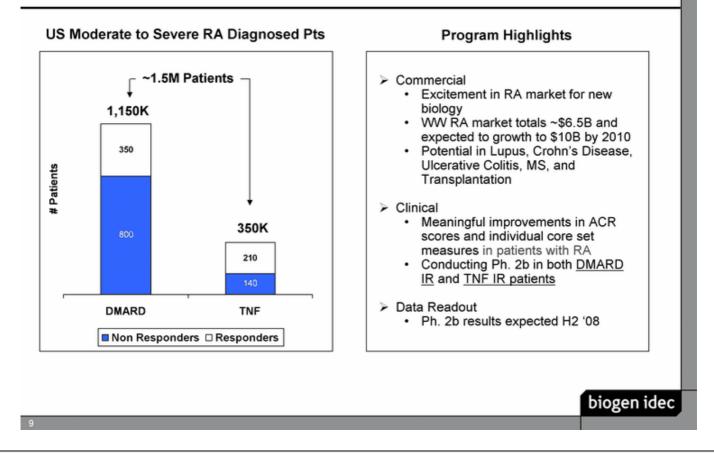
2008 Pipeline Data Readouts



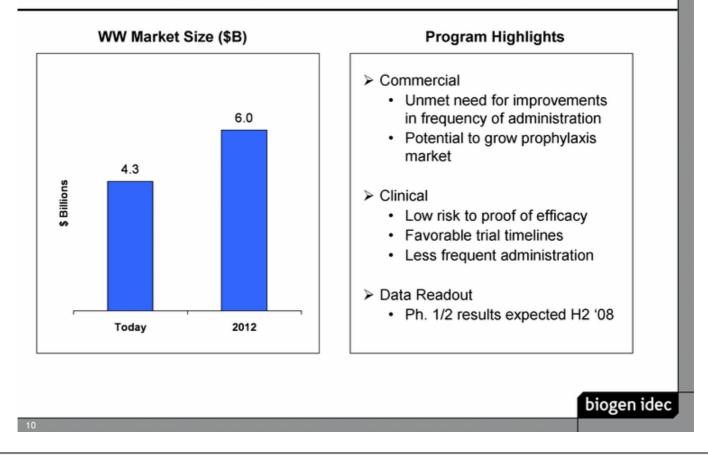
Select Programs with 2008 Readouts RITUXAN®: Ph. 2/3 Lupus



Select Programs with 2008 Readouts Baminercept: Ph. 2b RA

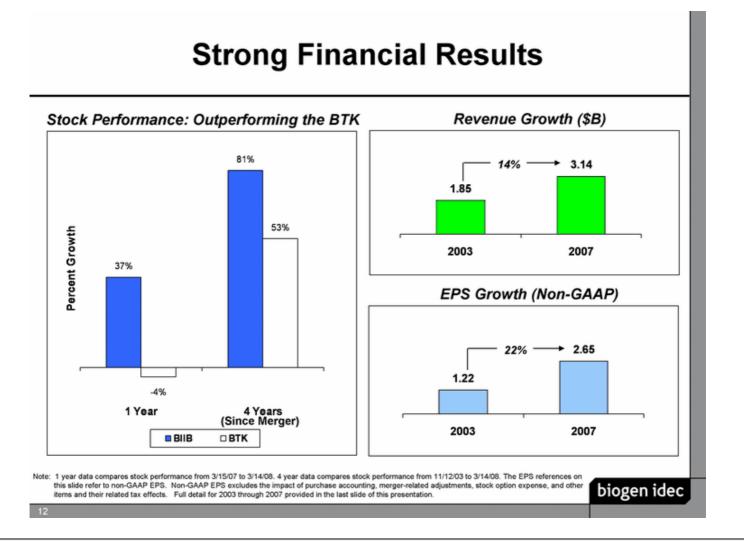


Select Programs with 2008 Readouts Long Acting rFactor IX: Ph.1/2 Hemophilia

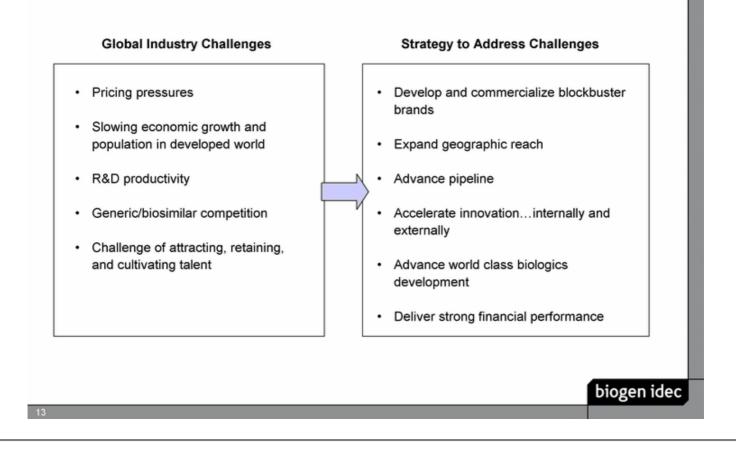


Late Stage Programs in Registrational Trials

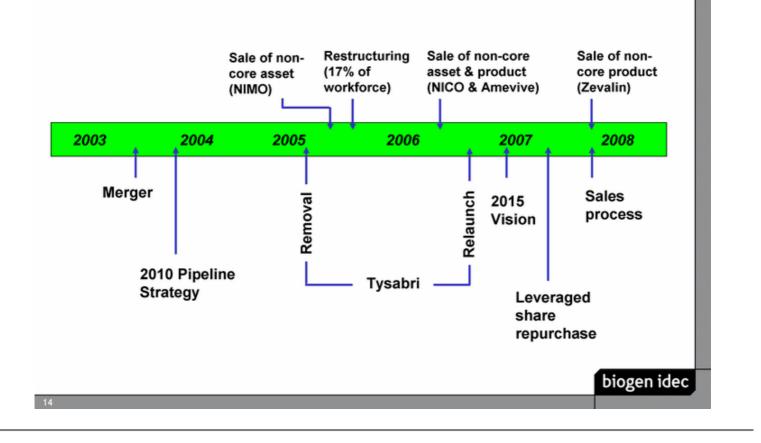
Program	Indication
• BG-12	MS (oral)
 Galiximab 	NHL
 Lixivaptan 	Hyponatremia/CHF
Lumiliximab	CLL
 Adentri (expected H2 '08) 	CHF
	biogen idec
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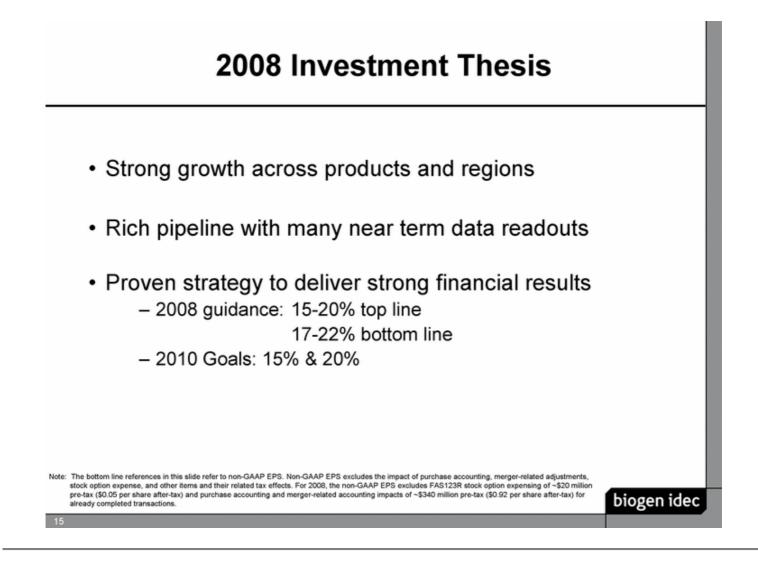


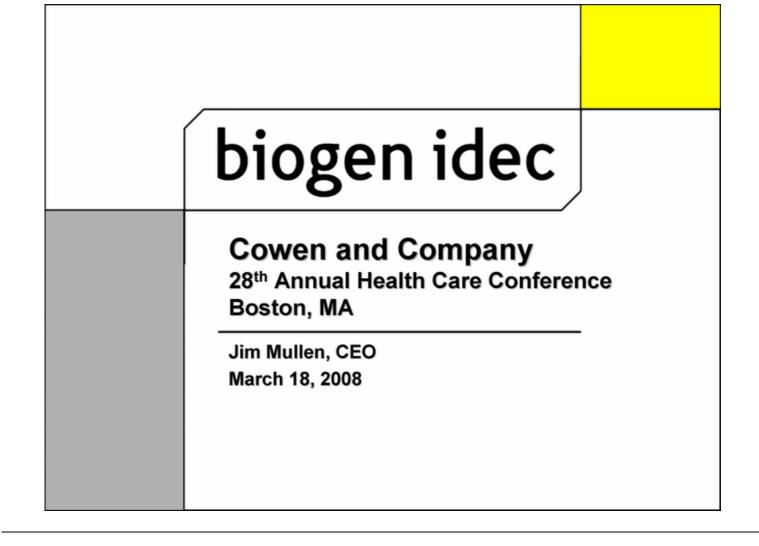
Pioneering New Solutions to Industry Challenges



Focused on Creating Value







GAAP to non-GAAP Reconciliation 2003 - 2007 Diluted EPS and Net Income

Condensed Consolidated Statements of Income – Operating Basis	FY 2003	FY 2004	FY 2005	FY 2006	FY 2007
GAAP diluted EPS	(4.92)	0.07	0.47	0.63	1.99
Adjustment to net income (see below)	6.14	1.38	1.10	1.62	0.75
Effect of FAS128 and ETIF 0306		(0.05)			
Non-GAAP diluted EPS	1.22	1.40	1.57	2.25	2.74
GAAP Net Income (\$M)	(875.1)	25.1	160.7	217.5	638.2
Revenue - Pre-merger Biogen product, royalty and corporate partner revenue	1,173.1	-			-
COGS - Fair value step up of inventory acquired from Biogen and Fumapharm	231.6	295.5	34.2	7.8	-
COGS - Pre-merger Biogen cost of sales	(179.2)	-		-	-
COGS - Royalties related to Corixa	1.8				
COGS – Amevive divesture			36.4		
R&D – Pre-merger Biogen net R&D	(301.1)				
R&D - Severance and restructuring		3.1	20.3	0.3	1.2
R&D - Sale of plant		-	1.9	-	-
SG&A – Pre-merger Biogen SG&A	(346.7)	-			
SG&A – Merger related and purchase accounting costs		-		0.1	
SG&A – Severance and restructuring	13.2	9.3	19.3	2.0	0.6
Amortization of intangible assets primarily related to Biogen merger	33.2	347.7	302.3	267.0	257.5
In-process R&D related to the Biogen Idec merger, acquisitions of Conforma,					
Syntonix, and Furnapharm, and consolidation of Cardiokine, Neurimmune and Escoublec	823.0	-		330.5	84.2
Loss/(gain) on settlement of license agreements with Fumedica and Fumapharm				(6.1)	
(Gain)/loss on sale of long lived assets			111.8	(16.5)	(0.4)
Other income, net: Pre-merger Biogen	32.9				
Other income, net: Consolidation of Cardiokine and Neurimmune and gain on sale of					(70.0)
long lived assets		•			(72.3)
Write down of investments		12.7			•
Charitable donations and legal settlements	30.7				
Income taxes - Effect of reconciling items	(205.8)	(195.4)	(145.2)	(70.3)	(65.5)
Stock option expense				44.5	35.6
Non-GAAP Net Income	431.7	498.0	541.7	776.8	879.1

Notes: The non-GAAP financial measures presented in this table are utilized by Biogen Idec management to gain an understanding of the comparative financial performance of the Company. Our non-GAAP financial measures are defined as reported, or GAAP, values excluding (1) purchase accounting and mergerrelated adjustments, (2) stock option expense and the cumulative effect of an accounting change relating to the initial adoption of SFAS No. 123R and (3) other financial measures to establish financial goals and to gain an understanding of the comparative financial performance of the Company from year to year and quarter to quarter. Accordingly, we believe investors' understanding of the Company's financial performance is enhanced as a result of our disclosing these non-GAAP financial measures. Non-GAAP net income and non-GAAP diluted EPS should not be viewed in isolation or as a substitute for reported, or GAAP, net income and filuted EPS.

The GAAP figures reflect:

* 2004 and beyond - the combined Biogen Idec

 2003 – a full year of IDEC Pharmaceuticals and 7 weeks of the former Biogen, Inc. (for the period 11/13/03 through 12/31/03)

Numbers may not foot due to rounding.

Source: Biogen Idec Annual Reports, 10-K filings and earnings press releases (FY 2003-2007).

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