BizInt Smart Charts

What's New with the BizInt Smart Charts Family

ICIC, Berlin 15 October 2012 John Willmore, VP Product Development

- Improvements on STN, Orbit, Patbase, etc.
- Added support for Thomson Reuters Cortellis
- Generate links to Thomson Innovation and Patentscope
- Control over publication number display
- Many new non-patent literature files (STN)



- Continued improvements to ClinicalTrials.gov
- Improved mapping of fields between Citeline TrialTrove Adis Clinical Trials Insight ClinicalTrials.gov
- Generate Common Trial ID



- Added support for: DGENE and PCTGEN on STN
- Create a standardized sequence ID (e.g. from PSL or Note in DGENE)
- Combine results from multiple queries



- Improved column selection
- Easier combine/update operations
- New Excel and Acrobat exports
- Move rows command
- New toolbar, application icons



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	Common Drug Name	Database	Status	Companies		
1	FP-1039	Adis R&D Insight	Phase I	Five Prime Therapeutics (Originator) Human Genome Sciences (Licensee)		
2	FP-1039	IMS R&D Focus	1. Multiple Records for the	vePrime (USA) Iman Genome Sciences (USA)		
3	FP-1039	Thomson Pharma	Same Entity	vePrime Therapeutics Inc Iman Genome Sciences Inc		
4	FP-1039	Citeline Pipeline	Phase II	Five Prime Therapeutics Human Genome Sciences		
5	lexatumumab	Thomson Pharma	Phase 2 Clinical	Cambridge Antibody Technology Group plc Human Genome Sciences Inc		
6	lexatumumab	Adis R&D Insight	Discontinued I	Cambridge Antibody Technology (Originator) Human Genome Sciences (Licensee)		
7	lexatumumab	Citeline Pipeline	No Development Reported	Human Genome Sciences AstraZeneca		
8	lexatumumab	IMS R&D Focus	Phase I	Human Genome Sciences (USA)		





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1	FP-1039	Adis R&D Insight	Phase I	Five Prime Therapeutics (Originator) Human Genome Sciences (Licensee)			
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5	lexatumumab	Thomson Pharma	Phase 2 Clinical	Cambridge Antibody Technology Group plc Human Genome Sciences Inc			
6	lexatumumab	Adis R&D Insight	Discontinued I	Cambridge A 2. Conflicting Itor) Human Gence Data for the			
7	lexatumumab	Citeline Pipeline	No Development Reported	Human Genc AstraZeneca Same Entity			
8	lexatumumab	IMS R&D Focus	Phase I	Human Genome Sciences (USA)			





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3	FP-1039	Thomson Pharma	Phase 2 Clinical	FivePrime Therapeutics Inc. Human 3. Terminology		
4	FP-1039	Citeline Pipeline	Phase II	Five Pr Human Variation		
5	lexatumumab	Thomson Pharma	Phase 2 Clinical	Cambridge Antibody Technology Group plc Human Genome Sciences Inc		
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7	lexatumumab	Citeline Pipeline	No Development Reported	Human Genome Sciences AstraZeneca		
8	lexatumumab	IMS R&D Focus	Phase I	Human Genome Sciences (USA)		





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	Common Drug Name	Database	Status	the
1	FP-1039	Adis R&D Insight	Phase I	Five vantage point
2	FP-1039	IMS R&D Focus	1. Multiple Records for the	e BizInt Smart Charts
3	FP-1039	Thomson Pharma	Same Entity	rePrime Therapeutics Inc. ^{Imal} 3. Terminology
4	FP-1039	Citeline Pipeline	Phase II	Five Pr Human Variation
5	[∎] BizIn	t Smart C	harts 🖆	Cambridge Antibody Technology Group plc Human Genome Sciences Inc
6	le	erence R		Cambridge A 2. Conflicting tor) Human Gence Data for the
7	IE		Reported	Human Gence Same Entity
8	lexatumumab	IMS R&D Focus	Phase I	Human Genome Sciences (USA)



- Customized version of VantagePoint designed for BizInt Smart Charts users.
- Provides tools for normalization, cleanup, analysis and visualization.
- Uses the XML Smart Data Exchange format.
- Sold separately (\$3000 for first user)
- Contact us to arrange a trial!



Clean-up company names...

					60	6S	es	
	# Records	# Instances	stances	# Records	# Instances	Companies:		
	¥.	#		1	219	219	Human Genome Sciences	
1	69	69	Hun	2	38	39	GlaxoSmithKline	
2	54	54	Hun	3	14	14	Medimmune	
3	49	49	Hun	4	9	9	AstraZeneca	
4	42	42	Hun	5	9	9	Aventis Behring LLC	
5	14	14	Gla	6	9	9	Transgene	
6	12	12	Med	7	8	8	Takeda	
7	10	10	Glap	8	7	7	Amgen	
8	10	10	Gla	9	7	7	Cambridge Antibody Technology	
9	9	9	Ave	10	7	7	Pfizer	
10	8	8	Ast	11	6	6	Novartis	
11	5	5	Am	12	5	5	Genentech	
12	5	5	Hun	13	5	5	Teva	
13	4	4	Can	in lage	Annou	5100	noogy Group pr	www.bizcharts.com

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Normalize phase terminology...

	# Records	# Instances	Status	-	sp	ces	÷
1 2 3 4	26 11 3 7	26 11 3 7	Discontinued Discontinued I Discontinued II Discontinued Preclinical		# Records	# Instances	Status (1)
5	1	1	Discontinued preregistra	1	79	79	No Development Reported
7	3	3	L. hed	2	52	52	Discontinued
3	2 62	2 62	Man No Dev. me porti	3	45	45	Preclinical
10 11	1	1	No devel reporte		13	13	Phase 2
12	3	3	Phase nic.	5	8	8	Phase 1
13 14	2	2	Phr Clinical	6	8	8	Phase 3
15	10	10	ase II Phase III	7	5	5	Launched
16 17	1	1	Pre-registration	8	4	4	Technology
18	40	40	Preclinical	9	3	3	Biological Testing
19 20	2	2	Preregistration Research	10	3	3	Preregistration
21	3	3	Suspended				
22 23	1	1	Suspended II Technology				

BizInt Smart Charts

Reference Rows[™]



- Information from related records is displayed in a single "Reference Row"...
- ...based on database rankings and rules that you define.
- A separate utility working with patent, drug pipeline, and clinical trial reports.
- Included in all BizInt Smart Charts licenses.



Now we can provide better tabular reports.

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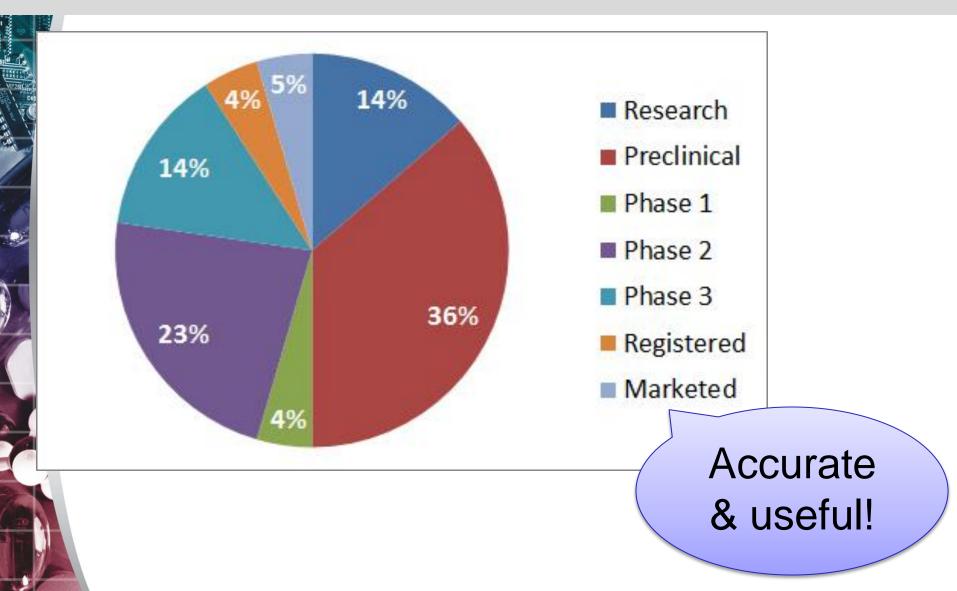
Human Genome Sciences - Pipeline (TP, RDI, RDF, CP - Aug 2012)

	Drug	Database	Status (Arabic)	Companies (Cleaned)	Update Date	Summary
1.	balugrastim	1.1TPharm link1.2IMS link1.3Pipeln link	Phase 3	Aventis Behring LLC CoGenesys Inc Human Genome Sciences Teva	2012-08-07	Teva Pharmaceutical Industries, following the acquisition of CoGenesys, under license from Human Genome Sciences (HGS, formerly Principia), and under license from Aventis Behring (now ZLB Behring), is developing balugrastim (CG-10639, Neugranin, Albugranin), a long-acting form of G-CSF created by fusion to albumin, for the potential treatment of chemotherapy-induced neutropenia [447883], [678434], [778202], [780827], [878888] [CONT.]
	1.1 TPharm				1.1 TPharm	1.1 TPharm
2.	Darapladib	2.1Adis link2.2Adis link2.3IMS link2.4Pipeln link	Phase 3	GlaxoSmithKline Human Genome Sciences	2012-08-23	Darapladib (SB-480848) is an oral once-daily lipoprotein- associated phospholipase A2 (Lp-PLA2) inhibitor isolated from Pseudomonas fluorescens, under development by GlaxoSmithKline (GSK) for the treatment of atherosclerosis and diabetic macular oedema (DMO) (Company pipeline, GSK, Feb 2001; USAN Web Page, 8 Nov 2005; Company presentation, GSK, 5 Feb 2009; Ann Rep, GSK, 2011, page 236, http://www. [CONT.]
	2.1 Adis				2.3 IM S	2.4 Pipeln
3.	FP-1039 3.1 TPharm	3.1 TPharm link 3.2 Adis link 3.3 IMS link 3.4 Pipeln link	_	Five Prime Therapeutics	2012-08-07	FivePrime Therapeutics and licensee Human Genome Sciences (HGS), a subsidiary of GlaxoSmithKline, are developing FP-1039 (HGS-1036), a soluble fusion protein comprising the extracellular domain of FGFR-1C linked to the Fc domain of IgG1 as a FGFR-1C antagonist that binds to several ligands that are key drivers of tumor growth, for the potential iv treatment of cancer such as breast, lung, ovarian and endometrial tumors [664645], [1108287], [1108288], [1162681], [1176971] [CONT.] 3.1 TPharm
-						

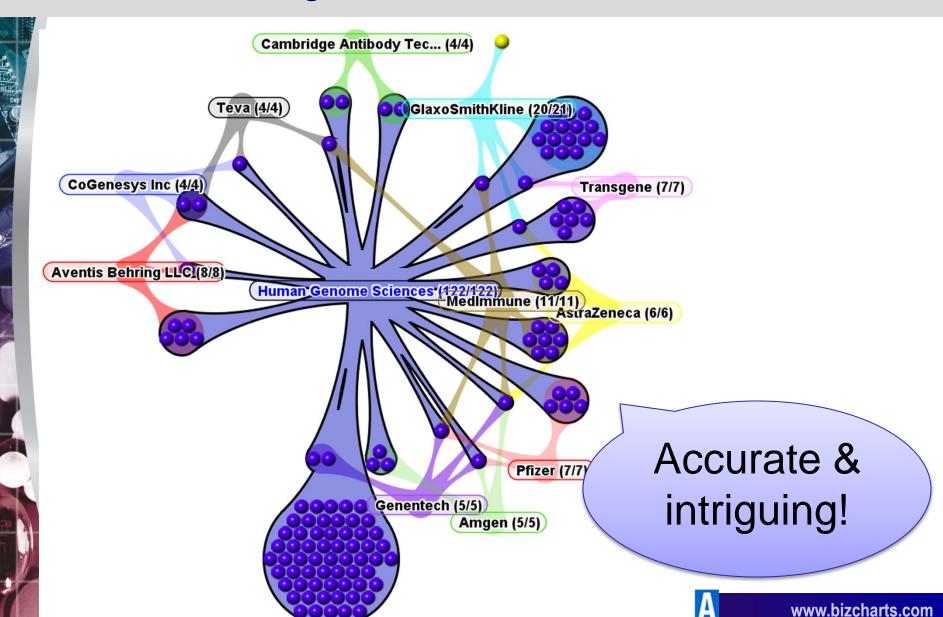
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Or graphs...



Or network diagrams.





A "Cookbook" of Visualizations Created with the

BizInt Smart Charts

Product Family

Pharma-Bio-Med

Lisbon, Portugal October 2012 BizInt Smart Charts
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