# HIPAA Transactions The Next Generation

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Intelligently Linking Information Systems



## Today's Session

Objective: Provide information that allows impacted organizations to track and participate in future HIPAA transactions activities; thereby managing their futures

#### Topics:

- ✓ Materials Used in HIPAA Transactions
- ✓ Processes for Creating Materials
- ✓ Status, Predictions, and Key Issues
- ✓ Obtaining Further Information

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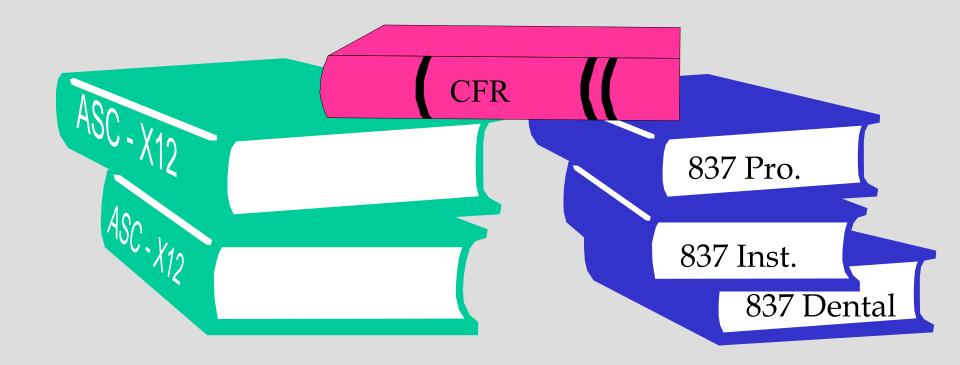
#### Caveats

For clarity and simplicity, today's discussion primarily illustrates the ANSI SDO processes of Accredited Standards Committee X12. Similar but differing processes also exist at other HIPAA SSO's.

The predictions contained in today's presentation are solely those of the author and do not represent the views, official or unofficial, of anybody else.



#### Materials





## HIPAA Transactions Specifications

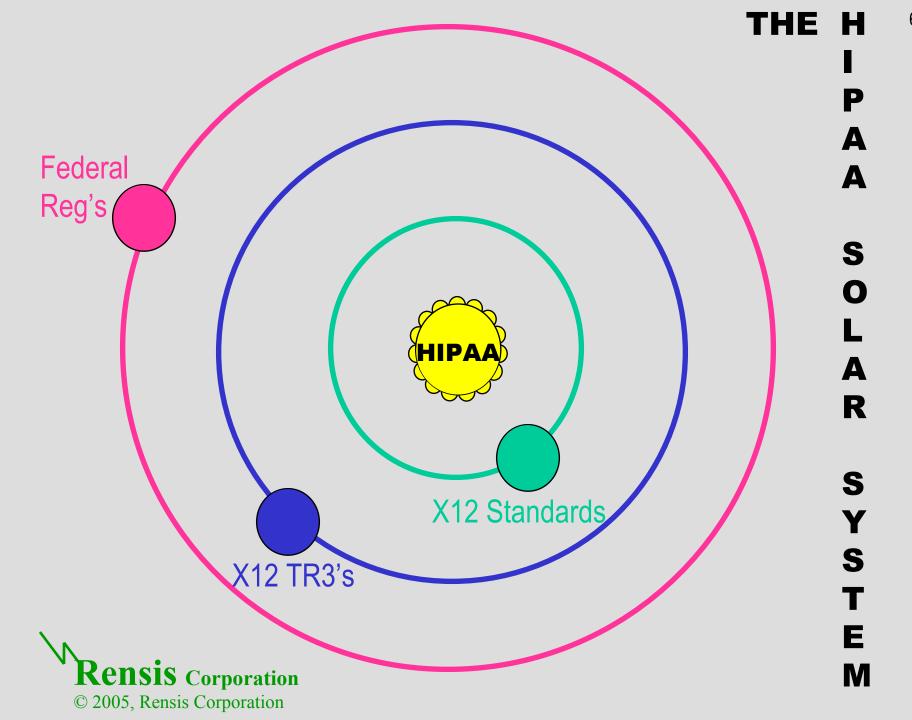
Mandatory Federal Regulations ["Rules"]
 which "adopt" and promulgate

Voluntarily published X12 (and equivalent)
 Type 3 Technical Reports (TR3's)
 a.k.a. "HIPAA Standards"

which define precise uses of

Voluntarily published X12 Standards





#### X12 Standards

Publication Cycle 3 times a year

Publisher Data Interchange

**Standards Association** 

Governing Materials Standing Doc. 2 (SD2)

Authoring Entities X12N Workgroups

Supporting Entities X12N / TG8 (Architecture)

X12J (Tech. Assessment)

Procedures Review Board

(PRB)



## X12 Type 3 Technical Reports

Publication Cycle

not yet fully stabilized, trying for every 2 years

Publisher

Washington Publishing Co.

Governing Materials

IG Handbooks

Authoring Entities

X12N Workgroups

Supporting Entities

X12N / TG4 (IG Coord.) X12J (Tech. Assessment)



#### Federal Regulations

Publication Cycle as recommended

Publisher Government Printing Office

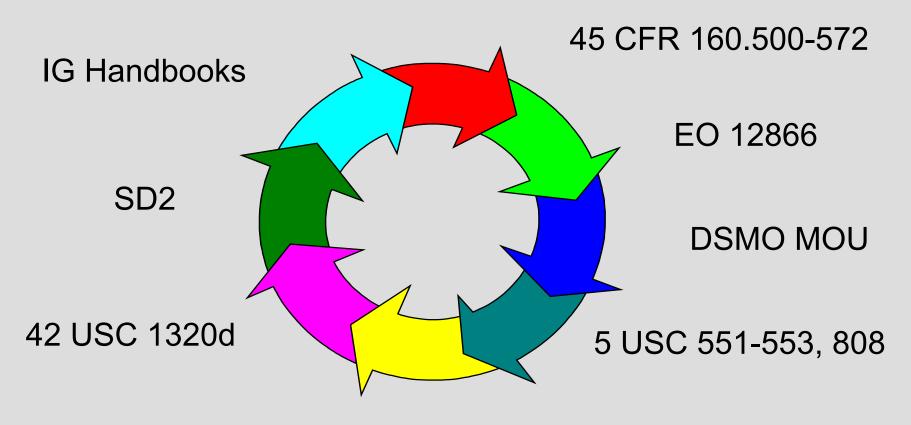
Governing Materials HIPAA Legislation
 Administrative Procedures
 Act

Authoring Entity CMS' Office of HIPAA Stds.

Supporting Entities DSMO Steering Committee
 NCVHS

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#### Processes



45 CFR 162.910

63 FR 88, pp 25274-25294



#### **Transactions Processes**

- Updating and creating new X12 standards; including internal code lists
- Creating and modifying Type 3 Technical Reports (TR3's); including internal code lists subsets
- Adopting TR3's for HIPAA



## **Updating Standards**

- X12 has two formal processes documented in Standing Document 2 (SD2)
  - Data Maintenance (DM)
    - For message structure, format, data element definitions, and internal code lists values
    - Can take many months or years
  - ° Code Maintenance Request (CMR)
    - For internal code lists values only
    - Expedited process to speed-up changes
    - Can still take 4 8 months



- X12N process summary
  - Work groups within authoring task groups, in conjunction with Washington Publishing Company, establish schedule [including change request cutoff dates for various sources] and then create new TR3's
  - ° Following internal approvals for technical accuracy and proper process from supporting task groups, work groups commence X12N public comment period for new TR3's

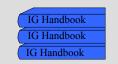
- X12N process summary
  - ° TR3's <u>public comment period</u> occurs
    - were 30 days; but being expanded to approximately 60 days
  - Work groups resolve any issues raised during public comment period and make any needed adjustments to TR3's
  - Work groups hold <u>public Informational Forums</u> during X12 Trimester Meetings to confirm resolved issues and TR3's adjustments

- X12N process summary
  - ° Work groups vote to move TR3's to task group for publication approval
  - ° Task groups [only TG2 Healthcare, at present] vote to move TR3's to subcommittee X12N – Insurance for publication approval
  - ° X12N approves TR3's for publication
  - ° Any other affected X12 subcommittees approve TR3's for publication [new for TR3's]



- X12N process summary
  - ° X12J Technical Assessment subcommittee approves TR3's for publication [new for TR3's]
  - ° Procedures Review Board approves TR3's for publication [new for TR3's]
  - Washington Publishing Company publishes





- X12N process timing
  - Overall IG and TR3 timing is, at present, variable
  - ° Focused effort is underway to manage volunteer resources and demands to establish two year TR3 publication cycle
    - Six approval slots every two years
    - Groups of TR3's HIPAA and non-HIPAA allocated to each slot on rotating schedule



#### Two cycle process – first iteration

- X12N proposes new version of published Type 3 Technical Reports (TR3's)
- Designated Standards Maintenance Organizations (**DSMO**) Steering Committee approves new version
- National Committee on Vital and Health Statistics (NCVHS) recommends new version

- Centers for Medicare and Medicaid Services (CMS) prepares Notice of Proposed Rule Making (NPRM) announcing new version
- Department of Health and Human Services (DHHS) clears NPRM
- Other affected federal agencies (e.g., Office of Management and Budget) approve NPRM

- NPRM is published in Federal Register
- Public comment period occurs
  - normally 60 days
- CMS, with any needed support from DSMO Steering Committee, X12N, et. al., analyzes comments received about NPRM

#### Two cycle process – second iteration

- Based on received comments, if necessary, X12N incorporates changes into next published new version of TR3's
- DSMO Steering Committee approves new version
- NCVHS recommends new version
- CMS prepares Final Rule promulgating new version



- DHHS clears Final Rule
- Other affected federal agencies (e.g., OMB) approve Final Rule

- Final Rule is published in Federal Register
  - Specifies explicit Effective Date
     [Effective Date also known as Adoption Date]
  - ° Specifies explicit Compliance Date(s)
- For an existing HIPAA standard, any
   Effective Date for a modified standard must be at least 12 months following any previous Effective Date



Effective Date occurs no earlier than the end of mandatory Congressional Review period which is normally 60 days

#### **Compliance Date(s)**

- New Standards 24 months after Effective Date; small health plans get 36 months
- Modified Standards established within the Final Rule, but must be at least 180 days after Effective Date



## Status, Predictions, and Key Issues



as of 5 August 2005



## David A. Feinberg, C.D.P.

- Consultant and Teacher -- Healthcare Interfaces and EDI
- Author, "Understanding HIPAA Communications"
- Member, Accredited Standards Committee X12 and its Insurance Subcommittee (X12N)
- Member, Health Level Seven (HL7)
- Co-Chair, X12N HIPAA Implementation Work Group
- Member, HL7 Attachments Special Interest Group (ASIG) and X12N Patient Information Work Group (TG2/WG9)
- Member, HL7 Imaging Integration Special Interest Group (IISIG) and DICOM Image Integration Group (WG20)
- Member, concluded HL7 Master Person Index Mediation Special Interest Group (MPISIG)
- Commercial and Technology Arbitrator, American Arbitration Association





#### **Transactions Futures**

- Claims Attachments
- New Versions of Current Transactions
- Potential New Transactions



#### Claims Attachments

- Defined by HL7 Attachments Special Interest Group (ASIG) in "Specifications"
- Presently planned to incorporate XML within EDI; i.e.,
  - ° X12's 275 transaction ... contains
    - HL7's Clinical Document Architecture (CDA) ...
       made up of
      - Structured data elements,
      - Narrative, unstructured, text, and/or
      - Scanned, non-diagnostic, images [many formats]

#### Claims Attachments

- First Round
  - ° Ambulance
  - Emergency Department
  - Rehabilitative Services
  - Laboratory Results
  - Medications
  - ° Clinical Notes





#### Claims Attachments

#### Status

- ° X12 and HL7 materials ready ... again
- NPRM sent to OMB on 6/02/2005 for last major clearance
- ° NPRM targeted for publication on 9/23/2005
- Pilot project in final stages
- ° Comments on NPRM and lessons from pilot project plus any other proofs of concepts expected to be fed back to update X12 and HL7 materials for use in final rule

#### New Versions of Current Txns.

- X12 version 004010 + 004010A1
   Implementation Guides (IG's)
  - Remain current HIPAA standards
  - ° X12 web site for obtaining HIPAA IG interpretations opened to public on 11/08/2004

www.x12n.org/portal

#### New Versions of Current Txns.

- X12 version 004050 counterpart IG's approved for publication during 2003
  - Contain additional useful explanations that can be applied to current HIPAA standards
  - Not presently planned to be generally proposed as modified HIPAA standards, but ...

## v4050 835 IG Proposed for HIPAA

From the HIPAA DSMO change request system site, www.hipaa-dsmo.org/crs

Change Request ID: 1008 Submission Date: 8/30/2004

Request Type: Payment of a Health Care Claim

<u>Business Reason</u>: The later version of 835 Implementation Guide contains additional valuable information that will benefit the industry for those attempting to use the 835.

The Claim Payment workgroup and the Health Care Task Group of ASC X12 Insurance Sub Committee believes that this new guide version 4050 designated X124 should be considered as a candidate for the next HIPAA version of the 835.

Suggested Change: Recommend to NCVHS that the 4050 version of the 835 Implementation Guide (X124) be named as the HIPAA standard.

Copies of the **004050X124** document for the 835 "Health Care Claim Payment/Advice" transaction, may be obtained for a modest fee from <a href="https://www.wpc-edi.com/products/publications">www.wpc-edi.com/products/publications</a>



#### New Versions of Current Txns.

- Writing of X12 version 005010 counterpart TR3's now underway
  - Planned changes include
    - Additional useful explanations
    - Accumulated and timely new routine requests
    - National Provider Identifier (NPI) adaptations
  - ° Planned changes do not at this time include
    - ICD-10 modifications
  - New change requests now being considered only for subsequent versions (e.g., 005030)



#### New Versions of Current Txns.

- Writing of X12 version 005010 counterpart TR3's now underway
  - ° Public comment periods projected in phases during 2004 2006 [see schedule slides]
  - Presently targeted for publication in stages during 2005 - 2006
  - ° Tentatively planned to be proposed for reference in NPRM for modified HIPAA standards: 2006 - 2007 - 2008?!
  - ° Volunteers welcomed! Contact me if you'd like further information.

## New Versions of Current Txns. 005010 TR3 Public Comment Periods

Actual X12 TR3 Comment Period

HIPAA Counterpart Transactions

Other, beyond HIPAA, Healthcare Transactions

Oct. – Nov. 2004

824 Acknowledgements

Feb. – Mar. 2005 835 Remittance Advice

834 Enrollment

837 Professional Claim

837 Institutional Claim

837 Dental Claim

274 Provider Inquiry – 4050

837 Data Reporting

997 Functional Ack.



continued on next slide ...

Note: TR3's are presently known as IG's

### New Versions of Current Txns. 005010 TR3 Public Comment Periods

Projected X12

TR3 Comment Period

HIPAA Counterpart

**Transactions** 

Other, beyond HIPAA, Healthcare Transactions

June – Sept.

2005

820 Premium Payment

278 Auth. Req. Review & Resp.

278 Auth. Serv. Review Notice

278 Auth. Inquiry & Response

269 Benefit Verification

999 Implementation Ack.

824 Acknowledgement – 4010

274 Provider Inquiry – 4050

277 Claim Acknowledgement

(moving to later period)

Informational Forums targeted for 26-28 Sept. at X12 Trimester Meeting in Atlanta. <a href="https://www.x12.org/x12org/meetings/x12trimt/index.cfm">www.x12.org/x12org/meetings/x12trimt/index.cfm</a>



continued on next slide ...

Note: TR3's are presently known as IG's

### New Versions of Current Txns. 005010 TR3 Public Comment Periods

Projected X12

**TR3 Comment** 

Period

HIPAA Counterpart

**Transactions** 

Other, beyond HIPAA, **Healthcare Transactions** 

Oct. – Nov.

2005

276/277 Claim Status

270/271 Eligibility Inq. & Resp.

275 Auth. Attachment

Feb. – Mar.

2006

271 Roster

274 Provider Directory

274 Provider Credentialing

277 Req. for Additional Info.

275 Claim Attachment

June - July





 Should the participating organizations (X12, DSMO Steering Committee, NCVHS, OESS ... CMS ... DHHS) execute their portions of the HIPAA adoption process on a staggered schedule as groups of TR3's are published, or wait until a complete suite (e.g., 005010, 005030) is available?

- Pilot projects
  - ° How many, if any, are needed?
  - ° How comprehensive should they be?
  - ° How long should they run?
  - ° When should they be executed in relation to development and adoption process steps?
  - ° Who will participate? How will participation be arranged and funded?
  - ° Who will manage and/or consolidate results?



 Cost vs. benefit (i.e., return on investment) analyses



- ° When and how extensive should any be?
- ° Who should perform them?
- Should HIPAA adoption be done just because new transaction versions are simply necessary to comply with other federal regulations (e.g., NPI, e-prescribing)?
- ° Can there ever be a pay-back for moving to new versions of current transactions?

- At what point should the federal government commence its portions of the HIPAA adoption process? What triggers these activities?
- How do the individual DSMO, who are continually developing new materials (e.g., X12 TR3's), interact during the 2+ year federal adoption sequence; especially with comments received from NPRM's?

- Bigger picture, what will be the impacts of
  - ° e-prescribing transactions standards?
  - Office of the National Coordinator for Health Information Technology (ONCHIT) proposed contract for Standards Coordination and Harmonization?
  - ° American Health Information Community (AHIC) federal advisory committee?
  - National Healthcare Information Infrastructure (NHII)?
  - ° Consolidated Health Informatics (CHI) Initiative?



#### Potential New Transactions

[999, 997, 824, 277] Acknowledgement Coordination of Benefit Confirmation **Provider Information** [274] Eligibility / Enrollment Roster [271] **Authorization Attachment** 275 **Additional Claims Attachments** [HL7 CDA]

Any HIPAA adoption activities will only begin by prior industry acceptance, use, and request



#### Further Information

Rensis Corporation Seminar:

"HIPAA TCS – What's Next? Products, Processes, and Prognostications"

- Feinberg's Free Focused HIPAA Mailing List Send e-mail request to one or more of
  - Opening of the computer of
  - \* HIPAA-TCS-subscribe@yahoogroups.com
  - ShareHIPAA-subscribe@yahoogroups.com

#### Further Information

Other Online Presentations – mostly free:

www.complyassistant.com/presentations.html

 White Papers on managing never-ending HIPAA TCS, Privacy, and Security compliance

www.complyassistant.com (left border)



Comments?

Questions?

Smart Remarks?

Contact Dave Feinberg





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